

APPROVED DRUG PRODUCTS

WITH

**THERAPEUTIC
EQUIVALENCE
EVALUATIONS**

44th EDITION

**THE PRODUCTS IN THIS LIST HAVE BEEN APPROVED UNDER
SECTION 505 OF THE FEDERAL FOOD, DRUG, AND COSMETIC ACT.**

**U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
OFFICE OF MEDICAL PRODUCTS AND TOBACCO
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF GENERIC DRUGS
OFFICE OF GENERIC DRUG POLICY**

2024

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The products in this list have been approved under section 505 of the Federal Food, Drug, and Cosmetic Act. This volume is current through December 31, 2023.

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**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPROVED DRUG PRODUCTS
With
Therapeutic Equivalence Evaluations**

PREFACE TO FORTY FOURTH EDITION

The publication, *Approved Drug Products With Therapeutic Equivalence Evaluations* (the List, commonly known as the Orange Book), identifies drug products approved on the basis of safety and effectiveness by the Food and Drug Administration (FDA) under the Federal Food, Drug, and Cosmetic Act (the FD&C Act). The main criterion for the inclusion of any product is that the product is the subject of an application with an approval that has not been withdrawn for safety or effectiveness reasons. Inclusion of products in the Orange Book is independent of any current regulatory action being taken administratively or judicially against a drug product. In addition, the Orange Book contains therapeutic equivalence evaluations for approved multisource prescription drug products. These evaluations have been prepared to serve as public information and advice to state health agencies, prescribers, and pharmacists to promote public education in the area of drug product selection and to foster containment of health care costs. Therapeutic equivalence evaluations in this publication are not official FDA actions affecting the legal status of products under the FD&C Act.

Background of the Publication. To contain drug costs, virtually every state has adopted laws and/or regulations that encourage the substitution of drug products. These state laws generally require either that substitution be limited to drugs on a specific list (the positive formulary approach) or that it be permitted for all drugs except those prohibited by a particular list (the negative formulary approach). Because of the number of requests in the late 1970s for FDA assistance in preparing both positive and negative formularies, it became apparent that FDA could not serve the needs of each state on an individual basis. The Agency also recognized that providing a single list based on common criteria would be preferable to evaluating drug products on the basis of differing definitions and criteria in various state laws. As a result, on May 31, 1978, the Commissioner of the Food and Drug Administration sent a letter to officials of each state announcing FDA's intent to provide a list of all prescription drug products that are approved by FDA for safety and effectiveness, along with therapeutic equivalence determinations for multisource prescription products.

The Orange Book was distributed as a proposal in January 1979. It included only currently marketed prescription drug products approved by FDA through new drug applications (NDAs) and abbreviated new drug applications (ANDAs) under the provisions of Section 505 of the FD&C Act and FDA regulations at that time.

The therapeutic equivalence evaluations in the Orange Book reflect FDA's application of specific criteria to the multisource prescription drug products listed in the Orange Book and approved under Section 505 of the FD&C Act. These evaluations are presented in the form of code letters that indicate the basis for the evaluation made. An explanation of the codes appears in the *Introduction*.

A complete discussion of the background and basis of FDA's therapeutic equivalence evaluation policy was published in the *Federal Register* on January 12, 1979 (44 FR 2932). The final rule, which includes FDA's responses to the public comments on the proposal, was published in the *Federal Register* on October 31, 1980 (45 FR 72582). The first publication of the Orange Book in October 1980, concurrent with finalization of the rule, incorporated appropriate revisions. Each subsequent edition has included new approvals and made appropriate changes in data.

On September 24, 1984, the President signed into law the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments). The Hatch-Waxman Amendments amended the FD&C Act to establish, among other things, the 505(b)(2) and 505(j) approval pathways. The Hatch-Waxman Amendments require that FDA, among other things, make publicly available a list of approved drug products with monthly supplements. The Orange Book and its monthly Cumulative Supplements satisfy this requirement. The *Addendum* to this publication identifies drugs that have qualified under the FD&C Act for periods of exclusivity and provides patent information concerning the approved drug products in the Orange Book. The *Addendum* also provides additional information that may be helpful to those submitting an NDA under Section 505(b) of the FD&C Act or an ANDA under Section 505(j) of the FD&C Act to the Agency.

The Agency intends to use this publication to further its objective of obtaining input and comment on the publication itself and related Agency procedures. Therefore, if you have comments on how the publication can be improved, please send them to the Central Document Room, Attn: Director, Division of Orange Book Publication and Regulatory Assessment (DOBPR), Office of Generic Drug Policy, Center for Drug Evaluation and Research, Food and Drug Administration, 5901-B Ammendale Rd., Beltsville, MD 20705-1266. Comments received are publicly available to the extent allowable under the Freedom of Information Act and FDA regulations.

1.0 INTRODUCTION

1.1 Content and Exclusion

The Orange Book is composed of four parts: (1) approved prescription drug products with therapeutic equivalence evaluations; (2) approved over-the-counter (OTC) drug products for those drugs that may not be marketed without NDAs or ANDAs because they are not subject to 505G; (3) drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research; and (4) a cumulative list of approved products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing and we have not determined that they were withdrawn from sale for safety or effectiveness reasons, or have had their approvals withdrawn for other than safety or effectiveness reasons subsequent to being discontinued from marketing.¹ This publication also includes indices of prescription and OTC drug products by proprietary name (brand name or trade name) or, if no proprietary name exists, established name of the active ingredient and by applicant name, which have been abbreviated for this publication. Established names for active ingredients generally conform to compendial names or *United States Adopted Names* (USAN) as described in 21 CFR 299.4(e). A list of uniform terms is provided in Appendix C.

The *Addendum* contains patent and exclusivity information for the Prescription, OTC, Discontinued Drug Product Lists, and for the Drug Products with Approval under Section 505 of the FD&C Act Administered by the Center for Biologics Evaluation and Research. The publication may include additional information that the Agency deems appropriate to disseminate.

Prior to the 6th Edition, the publication had excluded OTC drug products and drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research. The Hatch-Waxman Amendments required the Agency to begin publishing an up-to-date list of all marketed drug products, OTC as well as prescription, that have been approved for safety and effectiveness and for which NDAs are required.

Under the FD&C Act, some drug products are given tentative approvals. The Agency will not include drug products with tentative approvals in the Orange Book because a drug product that is granted tentative approval is not an approved drug product. Tentative approval lists by month are available on FDA's website [Drugs@FDA](#). When the tentative approval becomes a final approval through a subsequent action letter to the applicant, the Agency will list the drug product and the date of approval in the appropriate approved drug product list. In addition, we note that Section 505(x) of the FD&C Act affects the date of approval for certain drug products subject to scheduling under the Controlled Substances Act. For these drug products subject to scheduling, the Agency will list the drug product on FDA's website [Drugs@FDA](#) upon approval under Section 505(c) of the FD&C Act, and will list the drug product in the Orange Book upon the date of approval as determined under Section 505(x).

¹ Generally, newly approved products are added to the Active Section of the Orange Book (i.e., the Prescription Drug Product List or the Over-the-Counter Drug Product List), depending on the dispensing requirements (prescription or OTC) or approval authority, unless the Division of Orange Book Publication and Regulatory Assessment is otherwise notified before publication. See Section 1.12.

The Orange Book identifies the application holder of a drug product and does not identify distributors or repackagers.

1.2 Therapeutic Equivalence-Related Terms

Pharmaceutical Equivalents. Pharmaceutical equivalents are drug products in identical dosage forms and route(s) of administration that contain identical amounts of the identical active drug ingredient, i.e., the same salt or ester of the same therapeutic moiety, or, in the case of modified-release dosage forms that require a reservoir or overage or such forms as prefilled syringes where the residual volume may vary, that deliver identical amounts of the active drug ingredient over the identical dosing period; do not necessarily contain the same inactive ingredients; and meet the identical compendial or other applicable standard of identity, strength, quality, and purity, including potency and, where applicable, content uniformity, disintegration times, and/or dissolution rates.² They may differ in characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, within certain limits, labeling.

Pharmaceutical Alternatives. Pharmaceutical alternatives are drug products that contain the identical therapeutic moiety, or its precursor, but not necessarily in the same amount or dosage form, or the same salt or ester (e.g., tetracycline hydrochloride, 250 mg capsules vs. tetracycline phosphate complex, 250 mg capsules; quinidine sulfate, 200 mg tablets vs. quinidine sulfate, 200 mg capsules).³ Each such drug product individually meets either the identical or its own respective compendial or other applicable standard of identity, strength, quality, and purity, including potency and, where applicable, content uniformity, disintegration times, and/or dissolution rates.⁴ Different dosage forms and strengths within a product line by a single manufacturer are pharmaceutical alternatives, as are extended-release products when compared with immediate-release or standard-release formulations of the same active ingredient.

Therapeutic Equivalents. Approved drug products are considered to be therapeutic equivalents if they are pharmaceutical equivalents for which bioequivalence has been demonstrated, and they can be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.⁵

FDA classifies as therapeutically equivalent those drug products that meet the following general criteria: (1) they are approved as safe and effective; (2) they are pharmaceutical equivalents in that they (a) contain identical amounts of the identical active drug ingredient in the identical dosage form and route of administration, and (b) meet compendial or other applicable standards of strength, quality, purity, and identity; (3) they are bioequivalent in that (a) they do not present a known or potential bioequivalence problem, and they meet an acceptable *in vitro* standard, or (b) if they do present such a known or potential problem, they are shown to meet an appropriate bioequivalence standard; (4) they are adequately labeled; and (5) they are manufactured in compliance with Current Good Manufacturing Practice regulations. The concept of therapeutic equivalence applies only to drug products containing the identical active ingredient(s) and does not encompass a comparison of different therapeutic agents used for the same

² 21 CFR 314.3(b).

³ See 21 CFR 314.3(b).

⁴ 21 CFR 314.3(b).

⁵ 21 CFR 314.3(b).

condition (e.g., meperidine hydrochloride vs. morphine sulfate for the treatment of pain). Any drug product in the Orange Book repackaged and/or distributed by other than the applicant is considered to be therapeutically equivalent to the applicant's drug product even if the applicant's drug product is single source or coded as non-equivalent (e.g., **BN**). Distributors or repackagers of an applicant's drug product are not identified in the Orange Book.

FDA considers drug products to be therapeutically equivalent if they meet the criteria outlined above, even though they may differ in certain other characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, certain aspects of labeling (e.g., the presence of specific pharmacokinetic information), and storage conditions. When such differences are important in the care of a particular patient, it may be appropriate for the prescribing physician to require that a specific product be dispensed as a medical necessity. With this limitation, however, FDA believes that products classified as therapeutically equivalent can be substituted with the full expectation that the substituted product can be expected to have the same clinical effect and safety profile as the prescribed product when administered to patients under the conditions specified in the labeling.

Strength. Strength refers to the amount of drug substance contained in, delivered, or deliverable from a drug product, which includes: (1)(a) the total quantity of drug substance in mass or units of activity in a dosage unit or container closure (e.g., weight/unit dose, weight/volume or weight/weight in a container closure, or units/volume or units/weight in a container closure); and/or, as applicable, (b) the concentration of the drug substance in mass or units of activity per unit volume or mass (e.g., weight/weight, weight/volume, or units/volume); or (2) such other criteria the Agency establishes for determining the amount of drug substance contained in, delivered, or deliverable from a drug product if the weights and measures described in clause (1)(a) do not apply (e.g., certain drug-device combination products for which the amount of drug substance is emitted per use or unit time).⁶ Note that if the criteria the Agency establishes for determining and expressing the amount of drug substance in a product evolves over time, the Agency generally does not intend to revise the expressions of strength for drug products already included in the Orange Book, but rather intends to apply the criteria prospectively to drug products added to the Orange Book.

Although the strength of drug products in the Orange Book is generally expressed in terms of the amount of drug substance (active ingredient) in the drug product, it is sometimes expressed in terms of the amount of the active moiety.⁷ For example, certain drug products included in the Orange Book include a designation of "EQ" next to their expression of strength. This "EQ" designation generally is used in connection with salt drug products to indicate that the strength of such drug product is being expressed in terms of the equivalent strength of the active moiety (e.g., "EQ 200 MG BASE"), rather than in terms of the strength of the active ingredient.

Bioavailability. Bioavailability is the rate and extent to which the active ingredient or active moiety is absorbed from a drug product and becomes available at the site of drug action. For drug products that are not intended to be absorbed into the bloodstream, bioavailability may be assessed

⁶ 21 CFR 314.3(b).

⁷ Active moiety is the molecule or ion, excluding those appended portions of the molecule that cause the drug to be an ester, salt (including a salt with hydrogen or coordination bonds), or other noncovalent bonds (such as a complex, chelate, or clathrate) of the molecule, responsible for the physiological or pharmacological action of the drug substance. 21 CFR 314.3(b).

by scientifically valid measurements intended to reflect the rate and extent to which the active ingredient or active moiety becomes available at the site of drug action.⁸

Bioequivalence. Bioequivalence is the absence of a significant difference in the rate and extent to which the active ingredient or active moiety in pharmaceutical equivalents or pharmaceutical alternatives becomes available at the site of drug action when administered at the same molar dose under similar conditions in an appropriately designed study.⁹ Section 505(j)(8)(B) of the FD&C Act describes certain conditions under which a test drug and reference listed drug (RLD)(see Section 1.4) shall be considered bioequivalent:

- (i) the rate and extent of absorption of the [test] drug do not show a significant difference from the rate and extent of absorption of the [reference] listed drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses; or
- (ii) the extent of absorption of the [test] drug does not show a significant difference from the extent of absorption of the [reference] listed drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses and the difference from the [reference] listed drug in the rate of absorption of the drug is intentional, is reflected in its proposed labeling, is not essential to the attainment of effective body drug concentrations on chronic use, and is considered medically insignificant for the drug.

Where these above methods are not applicable (e.g., for drug products that are not intended to be absorbed into the bloodstream), other scientifically valid *in vivo* or *in vitro* test methods to demonstrate bioequivalence may be appropriate.

For example, bioequivalence may sometimes be demonstrated using an *in vitro* bioequivalence standard, especially when such an *in vitro* test has been correlated with human *in vivo* bioavailability data. In other situations, bioequivalence may sometimes be demonstrated through comparative clinical trials or pharmacodynamic studies.¹⁰

1.3 Further Guidance on Bioequivalence

FDA's regulations and guidance documents provide additional information regarding bioequivalence and bioavailability, including methodologies and statistical criteria used to establish the bioequivalence of drug products.¹¹

1.4 Reference Listed Drug and Reference Standard

⁸ 21 CFR 314.3(b).

⁹ 21 CFR 314.3(b).

¹⁰ 21 CFR 320.24.

¹¹ We note that prior to the 36th edition of the Orange Book, the Preface to the Orange Book included a section entitled "Statistical Criteria for Bioequivalence." Please see FDA's regulations and guidance documents for additional information regarding bioequivalence and bioavailability. See generally 21 CFR part 320. See FDA Drugs guidance Web page at <https://www.fda.gov/drugs/guidance-compliance-regulatory-information/guidances-drugs> and FDA Drugs guidance (Product-Specific Guidances for Generic Drug Development) Web page at <https://www.fda.gov/drugs/guidances-drugs/product-specific-guidances-generic-drug-development>.

A reference listed drug is the listed drug¹² identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA.¹³ FDA's general practice is to designate as RLDs drug products that have been approved for safety and effectiveness under Section 505(c) of the FD&C Act. For an ANDA based on an approved suitability petition (a petitioned ANDA), the reference listed drug generally is the listed drug referenced in the approved suitability petition.¹⁴

A reference standard is the drug product selected by FDA that an applicant seeking approval of an ANDA must use in conducting an *in vivo* bioequivalence study required for approval.¹⁵ FDA generally selects a single reference standard that ANDA applicants must use in *in vivo* bioequivalence testing. Ordinarily, FDA will select the reference listed drug as the reference standard. However, in some instances, the reference listed drug and the reference standard may be different. For example, where the reference listed drug has been withdrawn from sale for reasons other than safety or effectiveness, FDA may select an ANDA that is therapeutically equivalent to this reference listed drug as the reference standard.

FDA identifies reference listed drugs in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists. Listed drugs identified as reference listed drugs represent drug products upon which an applicant can rely in seeking approval of an ANDA. FDA intends to update periodically the reference listed drugs identified in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists, as appropriate.

If FDA has not designated a reference listed drug for a drug product the applicant intends to duplicate, the potential applicant may submit a controlled correspondence to the Office of Generic Drugs to ask FDA to designate a reference listed drug for that drug product. Section 1.7, *Therapeutic Equivalence Evaluations Codes (products meeting necessary bioequivalence requirements)* explains the character coding system (e.g., **AB**, **AB1**, **AB2**, **AB3**...) for multisource prescription drug products listed under the same heading with two or more reference listed drugs.

FDA also identifies reference standards in the Prescription Drug Product and OTC Drug Product Lists. Listed drugs identified as reference standards represent FDA's best judgment at this time as to the appropriate comparator for purposes of conducting any *in vivo* bioequivalence studies required for approval.

A potential applicant should consult Agency guidance related to referencing approved drug products in ANDA submissions for information on submitting a request for selection of a reference standard. FDA may, on its own initiative, select a new reference standard when doing so will help to ensure that applications for generic drugs may be submitted and evaluated, e.g., in the event that the listed drug currently selected as the reference standard has been withdrawn from sale.

¹² A "listed drug" is a new drug product that has been approved under Section 505(c) of the FD&C Act for safety and effectiveness or under Section 505(j) of the FD&C Act, which has not been withdrawn or suspended under Section 505(e)(1) through (5) or Section 505(j)(6) of the FD&C Act, and which has not been withdrawn from sale for what FDA has determined are reasons of safety or effectiveness. Listed drug status is evidenced by the drug product's identification in the current edition of FDA's "Approved Drug Products With Therapeutic Equivalence Evaluations" (the list) as an approved drug. A drug product is deemed to be a listed drug on the date of approval for the NDA or ANDA for that drug product (21 CFR 314.3(b)).

¹³ 21 CFR 314.3(b).

¹⁴ 21 CFR 314.94(a)(3)(i).

¹⁵ 21 CFR 314.3(b).

If an applicant has a question related to the appropriate reference standard, it is recommended that an applicant planning to conduct an *in vivo* bioequivalence study submit a controlled correspondence to the Office of Generic Drugs.

1.5 General Policies and Legal Status

The Orange Book contains public information and advice. It does not mandate the drug products that are purchased, prescribed, dispensed, or substituted for one another, nor does it, conversely, mandate the products that should be avoided. To the extent that the Orange Book sets forth FDA's evaluations of the therapeutic equivalence of drug products that have been approved, it contains FDA's advice to the public, to practitioners, and to the states regarding drug product selection. These evaluations do not constitute determinations that any product is in violation of the FD&C Act or that any product is preferable to any other. Therapeutic equivalence evaluations are a scientific judgment based upon evidence, while generic substitution may involve social and economic policy administered by the states, e.g., reducing the cost of drugs to consumers. To the extent that the Orange Book identifies drug products approved under Section 505 of the FD&C Act, it sets forth information that the Agency is required to publish and that the public is entitled to under the Freedom of Information Act. Exclusion of a drug product from the Orange Book does not necessarily mean that the drug product is in violation of Section 505 of the FD&C Act, that such a product is not safe or effective, or that such a product is not therapeutically equivalent to other drug products. Rather, the exclusion may be based on the fact that FDA has not evaluated the safety, effectiveness, and quality of the drug product.

1.6 Practitioner/User Responsibilities

Professional care and judgment should be exercised in using the Orange Book. Evaluations of therapeutic equivalence for prescription drugs are based on scientific and medical evaluations by FDA. Products evaluated as therapeutically equivalent can be expected, in the judgment of FDA, to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling. However, these products may differ in other characteristics that are not required by statute or regulation to be the same, such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, in some instances, labeling. If products with such differences are substituted for each other, there is a potential for patient confusion, e.g., due to differences in color or shape of tablets, inability to provide a given dose using a partial tablet if the proper scoring configuration is not available, or decreased patient acceptance of certain products because of flavor. There may also be patient-specific allergic reactions in rare cases due to a coloring or a preservative ingredient.

FDA evaluation of therapeutic equivalence in no way relieves practitioners of their professional responsibilities in prescribing and dispensing such products with due care and with appropriate information to individual patients. In those circumstances where the characteristics of a specific product, other than its active ingredient, are important in the therapy of a particular patient, the practitioner's prescribing of that product may be appropriate. Pharmacists must also be familiar with the different characteristics of therapeutically equivalent products, e.g., expiration dates/times and labeling directions for storage of the different products

(particularly for reconstituted products), so they can properly advise patients when one product is substituted for another.

Multisource and single-source drug products. In the Orange Book, FDA has evaluated for therapeutic equivalence only multisource prescription drug products approved under Section 505 of the FD&C Act, which in most instances means those pharmaceutical equivalents available (i.e., not on the Discontinued Drug Product list) from more than one manufacturer. For such products, a therapeutic equivalence code generally is included and product information is highlighted in bold face and underlined. Those products with approved applications that are single source (i.e., there is only one approved product available for that active ingredient, dosage form, route of administration, and strength) are also included in the Orange Book, but no therapeutic equivalence code is included with such products. Any drug product in the Orange Book repackaged and/or distributed by the applicant or some other person authorized by the applicant (e.g., an authorized generic) is considered to be therapeutically equivalent to the applicant's drug product even if the applicant's drug product is single source or coded as non-equivalent (e.g., **BN**). Distributors or repackagers of an applicant's drug product are not identified in the Orange Book. The details of therapeutic equivalence codes and the policies underlying them are discussed in Section 1.7, *Therapeutic Equivalence Evaluations Codes*.

Products in the Orange Book are identified by the names of the holders of approved applications (applicants) who may not necessarily be the manufacturer of the product. There are numerous entities other than the applicant that may be involved in the development, manufacturing, and/or marketing of a product. Products listed in the Orange Book are identified by the applicant's name (firm name on the Form FDA 356h in the application). Where the applicant's name does not appear on the label, a person wishing to relate a specific product to the applicant name in the Orange Book may refer to FDA's NDC Directory¹⁶ and match its search terms to information on the label, such as the NDC Code if available.

Every product in the Orange Book is subject at all times to regulatory action. From time to time, approved products may be found in violation of one or more provisions of the FD&C Act. In such circumstances, the Agency may commence appropriate enforcement action to correct the violation, if necessary, by securing removal of the product from the market by voluntary recall, seizure, or other enforcement actions. Such regulatory actions are, however, independent of the inclusion of a product in the Orange Book. The main criterion for inclusion of a product is that it has an NDA or ANDA that has been approved and that has not been withdrawn for safety or effectiveness reasons. FDA believes that retention of a violative product in the Orange Book will not have any significant adverse health consequences, because other legal mechanisms are available to the Agency to prevent the product's actual marketing. FDA may, however, change a product's therapeutic equivalence rating if the circumstances giving rise to the violation change or otherwise call into question the Agency's assessment of whether a product meets the criteria for therapeutic equivalence.

1.7 Therapeutic Equivalence Evaluations Codes

Generally, prescription drug products that the Agency considers multisource have been assigned a therapeutic equivalence code. The coding system for therapeutic equivalence evaluations is designed to allow users to determine quickly whether the Agency has evaluated a particular approved

¹⁶ <https://www.fda.gov/drugs/drug-approvals-and-databases/national-drug-code-directory>.

prescription drug product (e.g., a particular strength of an approved drug that is not on the Discontinued Drug Product list) as therapeutically equivalent to other pharmaceutically equivalent prescription drug products (first letter) and to provide additional information on the basis of FDA's evaluations (second letter). With some exceptions (e.g., therapeutic equivalence evaluations for certain 505(b)(2) applications), the therapeutic equivalence evaluation date is the same as the approval date.

The two basic categories into which multisource drugs have been placed are indicated by the first letter of the relevant therapeutic equivalence code as follows:

A Drug products that FDA considers to be therapeutically equivalent to other pharmaceutically equivalent products, i.e., drug products for which:

- (1) there are no known or suspected bioequivalence problems. These are designated **AA, AN, AO, AP, or AT**, depending on the dosage form; or
- (2) actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. These are designated **AB**.

B Drug products that FDA at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products, i.e.,

drug products for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence. Often the problem is with specific dosage forms rather than with the active ingredients. These are designated **BC, BD, BE, BN, BP, BR, BS, BT, BX, or B***.

Individual drug products have been evaluated as therapeutically equivalent to the reference product in accordance with the definitions and policies outlined below:

"A" CODES

Drug products that are considered to be therapeutically equivalent to other pharmaceutically equivalent products.

"A" products are those for which there are no known or suspected bioequivalence problems or for which actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. Drug products designated with an "A" code fall under one of two main policies:

- (1) for those active ingredients or dosage forms for which no *in vivo* bioequivalence issue is known or suspected, the information necessary to show bioequivalence between pharmaceutically equivalent products is either presumed and considered self-evident (based on other information in the application for some dosage forms (e.g., solutions)), or satisfied by a showing that an acceptable *in vitro* approach is met. A therapeutically equivalent rating is assigned such products so long as they are manufactured in accordance with Current Good Manufacturing Practice regulations and meet the other requirements of their approved applications (these are designated **AA, AN, AO, AP, or AT**, depending on the dosage form, as described below); or

- (2) for those Drug Efficacy Study Implementation (DESI) drug products containing active ingredients or dosage forms that have been identified by FDA as having actual or potential bioequivalence problems, and for post-1962 drug products presenting a potential bioequivalence problem, an evaluation of therapeutic equivalence is assigned to pharmaceutical equivalents only if the approved application contains adequate scientific evidence establishing through *in vivo* and/or *in vitro* studies the bioequivalence of the product to a selected reference product (these products are designated as **AB**).

There are some general principles that may affect the substitution of pharmaceutically equivalent products in specific cases. Prescribers and dispensers of drugs should be alert to these principles so as to deal appropriately with situations that require professional judgment and discretion.

There may be labeling differences among pharmaceutically equivalent products that require attention on the part of the health professional (e.g., pharmaceutically equivalent powders to be reconstituted for administration as oral or injectable liquids may vary with respect to their expiration time or storage conditions after reconstitution). FDA's determination that such products are therapeutically equivalent is applicable only when each product is reconstituted, stored, and used under the conditions specified in its labeling.

The Agency may use notes in this publication to point out special situations, such as potential differences between two drug products that have been evaluated as bioequivalent and otherwise therapeutically equivalent, when they should be brought to the attention of health professionals. These notes are contained in Section 1.8, *Description of Certain Special Situations*. For example, in certain instances, there may be variations among therapeutically equivalent products in their use or in conditions of administration. When such variations may, in the Agency's opinion, affect prescribing or substitution decisions by health professionals, a note may be added to Section 1.8.

For example, occasionally a situation may arise in which changes in a listed drug product after its approval (for example, a change in dosing interval) may have an impact on the substitutability of already approved generic versions of that product that were rated by the Agency as therapeutically equivalent to the listed product. When such changes in the listed drug product are considered by the Agency to have a significant impact on therapeutic equivalence, the Agency will change the therapeutic equivalence ratings for other versions of the drug product unless the manufacturers of those other versions of the product provide additional information to assure equivalence under the changed conditions. Pending receipt of the additional data, the Agency may add a note to Section 1.8, or, in rare cases, may even change the therapeutic equivalence rating.

In some cases (e.g., Isolyte® S w/ Dextrose 5% in Plastic Container and Plasma-Lyte® 148 and Dextrose 5% in Plastic Container), closely related products are listed as containing the same active ingredients, but in somewhat different amounts. In determining which of these products are pharmaceutically equivalent, generally the Agency has considered products to be pharmaceutically equivalent with labeled strengths of an ingredient that do not vary by more than 1%.

Different salts, esters or other noncovalent derivatives (such as a complex, chelate, or clathrate) of the same active moiety are regarded as different active ingredients. For the purpose of this publication, products containing such different active ingredients are considered pharmaceutical

alternatives and, thus, not therapeutically equivalent. Anhydrous and hydrated entities, as well as different polymorphs, are considered to be the same active ingredient and are expected to meet the same standards for identity to be considered pharmaceutical equivalents and therapeutic equivalents.

The codes in this book are not intended to preclude health care professionals from converting pharmaceutically different concentrations into pharmaceutical equivalents using accepted professional practice.

Where package size variations have therapeutic implications, products so packaged have not been considered pharmaceutically equivalent. For example, some oral contraceptives are supplied in 21-tablet and 28-tablet packets; the 28-tablet packets contain 7 placebo or iron tablets. These two packaging configurations are not regarded as pharmaceutically equivalent; thus, they are not designated as therapeutically equivalent.

Preservatives and other inactive ingredients may differ among some therapeutically equivalent drug products. These differences do not affect FDA's evaluation of therapeutic equivalence except in cases where these components may influence bioequivalence or routes of administration.

The specific sub-codes for those drugs evaluated as therapeutically equivalent and the policies underlying these sub-codes follow:

AA Products in conventional dosage forms not presenting bioequivalence problems

Multisource drug products coded as **AA** contain active ingredients and are in dosage forms that are not regarded as presenting either actual or potential bioequivalence problems or drug quality or standards issues. However, all oral dosage forms must, nonetheless, meet an appropriate *in vitro* bioequivalence standard that is acceptable to the Agency in order to be approved.

AB, AB1, AB2, AB3... Products meeting necessary bioequivalence requirements

Multisource drug products listed under the same heading (i.e., identical active ingredient(s), dosage form, and route(s) of administration) and having the same strength (see Section 1.2, *Therapeutic Equivalence-Related Terms, Strength*) generally will be coded **AB** if data and information are submitted demonstrating bioequivalence.

In certain instances, a number is added to the end of the **AB** code to make a three-character code (i.e., **AB1, AB2, AB3, etc.**). Three-character codes generally are assigned only in situations when more than one reference listed drug of the same strength has been designated under the same heading. If a study is submitted that demonstrates bioequivalence to a reference listed drug product, the generic product will be given the same three-character code as the reference listed drug it was compared against. For example, Adalat® CC and Procardia XL®, extended-release tablets, are listed under the active ingredient nifedipine. These drug products, listed under the same heading, are not bioequivalent to each other. Adalat® CC and Procardia XL® have been assigned ratings of **AB1** and **AB2**, respectively. Generic drug products deemed by FDA to be bioequivalent to Adalat® CC and Procardia XL® have been approved. As a result, the generic drug products bioequivalent to Adalat® CC have been assigned a rating of **AB1** and those bioequivalent to Procardia XL® have been assigned a rating of **AB2**. (The assignment of an **AB1** or **AB2** rating to a specific product does not imply product preference.) Even though drug products of distributors and/or repackagers are not included in the Orange Book, they are considered therapeutically equivalent to the applicant's drug product if the

applicant's drug product is rated either with an **AB** or three-character code or is single source in the Orange Book. Drugs coded as **AB** under a heading are considered therapeutically equivalent only to other drugs coded as **AB** under that heading. Drugs coded with a three-character code under a heading are considered therapeutically equivalent only to other drugs coded with the same three-character code under that heading.

AN Solutions and powders for aerosolization

Uncertainty regarding the therapeutic equivalence of aerosolized products arises primarily because of differences in the drug delivery system. Solutions and powders intended for aerosolization that are marketed for use in general-use delivery systems are considered to be pharmaceutically and therapeutically equivalent and are coded **AN**. Those products that are compatible only with a specific delivery system or those products that are packaged in and with a specific delivery system are coded **BN**, unless they have met an appropriate bioequivalence standard and are otherwise determined to be therapeutically equivalent. Solutions or suspensions in a specific delivery system will be coded **AN** if the bioequivalence standard is based upon *in vitro* methodology. If bioequivalence needs to be demonstrated by *in vivo* methodology, then the drug products will be coded **AB**.

AO Injectable oil solutions

The absorption of drugs in injectable (parenteral) oil solutions may vary substantially with the type of oil employed as a vehicle and the concentration of the active ingredient. Injectable oil solutions are therefore considered to be pharmaceutically and therapeutically equivalent only when the active ingredient, its concentration, and the type of oil used as a vehicle are all identical.

AP Injectable aqueous solutions and, in certain instances, intravenous non-aqueous solutions

It should be noted that even though injectable (parenteral) products under a specific listing may be evaluated as therapeutically equivalent, there may be important differences among the products in the general category, Injectable; Injection. For example, historically some injectable products that are rated therapeutically equivalent are labeled for different routes of administration. In addition, some products evaluated as therapeutically equivalent may have different preservatives or no preservatives at all. Injectable products available as dry powders for reconstitution, concentrated sterile solutions for dilution, or sterile solutions ready for injection are pharmaceutical alternative drug products. They are not rated as therapeutically equivalent (AP) to each other even if these pharmaceutical alternative drug products are designed to produce the same concentration prior to injection and are similarly labeled. Consistent with accepted professional practice, it is the responsibility of the prescriber, dispenser, or individual administering the product to be familiar with a product's labeling to assure that it is given only by the route(s) of administration stated in the labeling.

Certain commonly used large volume intravenous products in glass containers are not included in the Orange Book (e.g., dextrose injection 5%, dextrose injection 10%, sodium chloride injection 0.9%) since these products are on the market without FDA approval and FDA has not published

conditions for marketing such parenteral products under approved NDAs. When packaged in plastic containers, however, FDA regulations require approved applications prior to marketing. Approval then depends on, among other things, the extent of the available safety data involving the specific plastic component of the product. All large volume parenteral products are manufactured under similar standards, regardless of whether they are packaged in glass or plastic. Thus, FDA has no reason to believe that the packaging container of large volume parenteral drug products that are pharmaceutically equivalent would have any effect on their therapeutic equivalence.

Consistent with the definition of strength included in Section 1.2, *Therapeutic Equivalence-Related Terms*, the strength of parenteral drug products generally is identified by both the total drug content and the concentration of drug substance in a container approved by FDA.¹⁷ In the past, the strength of liquid parenteral drug products in the Orange Book has not been fully displayed. Rather, the strength of liquid parenteral drug products in the Orange Book has been displayed in terms of concentration, expressed as x mg/mL. Generally, the amount of dry powder or lyophilized powder in a container is identified as the strength, expressed as x mg/vial.

However, FDA subsequently realized that the format of the Orange Book with respect to parenteral solutions should be changed to reflect that each strength of a drug is considered to be a separate listed drug. The Orange Book generally displays the strength of new approvals of parenteral solutions. Previously (i.e., prior to 2003), we would have displayed only the concentration of an approved parenteral solution, e.g., 1 mg/mL. For example, if this application had a 125 mL and 250 mL container approved, we would now display two product strengths, listing both total drug content and concentration of drug substance in the relevant approved container, e.g., 125 mg/125 mL (1 mg/mL) and 250 mg/250 mL (1 mg/mL).

AT Topical products

There are a variety of topical dosage forms available for dermatologic, ophthalmic, otic, rectal, and vaginal administration, including creams, gels, lotions, oils, ointments, pastes, solutions, sprays, suppositories, and inserts. Even though different topical dosage forms may contain the same active ingredient and potency, these dosage forms are not considered pharmaceutically equivalent. Therefore, they are not considered therapeutically equivalent. All solutions and DESI drug products containing the same active ingredient in the same topical dosage form for which a waiver of *in vivo* bioequivalence has been granted, or the application contains adequate scientific evidence establishing through an *in vitro* approach the bioequivalence of the product to a selected reference product, and for which chemistry and manufacturing processes are adequate to demonstrate bioequivalence, are considered therapeutically equivalent and coded **AT**. Pharmaceutically equivalent topical products that raise questions of bioequivalence and for which a waiver of *in vivo* bioequivalence has not been granted, including all post-1962 non-solution topical drug products, are coded **AB** when supported by adequate *in vivo* bioequivalence data, and **BT** in the absence of such data.

"B" CODES

¹⁷ The strengths of certain parenteral drug products, including contrast agents, may be expressed as a percentage.

Drug products that FDA, at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products.

"B" products, for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence, often have a problem with specific dosage forms rather than with the active ingredients. Drug products designated with a "B" code fall under one of three main policies:

- (1) the drug products contain active ingredients or are manufactured in dosage forms that have been identified by the Agency as having documented bioequivalence problems or a significant potential for such problems and for which no adequate studies demonstrating bioequivalence have been submitted to FDA; or
- (2) the quality standards are inadequate or FDA has an insufficient basis to determine therapeutic equivalence; or
- (3) the drug products are under regulatory review.

The specific coding definitions and policies for the "B" sub-codes are as follows:

B* Drug products requiring further FDA investigation and review to determine therapeutic equivalence

The code **B*** is assigned to products previously assigned an **A** or **B** code when FDA receives new information that raises a significant question regarding therapeutic equivalence that can be resolved only through further Agency investigation and/or review of data and information submitted by the applicant. The **B*** code signifies that the Agency will take no position regarding the therapeutic equivalence of the product until the Agency completes its investigation and review.

BC Extended-release dosage forms (capsules, injectables and tablets)

Extended-release tablets are formulated in such a manner as to make the contained drug substance available over an extended period of time following ingestion.

Although bioavailability studies have been conducted on these dosage forms, they may be subject to bioavailability differences, primarily because applicants developing extended-release products for the same active ingredient rarely employ the same formulation approach. FDA, therefore, does not consider different extended-release dosage forms containing the same active ingredient in equal strength to be therapeutically equivalent unless equivalence between individual products in both rate and extent has been specifically demonstrated through appropriate bioequivalence studies. Extended-release products for which such bioequivalence data have not been submitted are coded **BC**, while those for which such data are available have been coded **AB**.

BD Active ingredients and dosage forms with documented bioequivalence problems

The **BD** code denotes products containing active ingredients with known bioequivalence problems and for which adequate studies have not been

submitted to FDA demonstrating bioequivalence. Where studies showing bioequivalence have been submitted, the product has been coded **AB**.

BE Delayed-release oral dosage forms

Where the drug may be destroyed or inactivated by the gastric juice or where it may irritate the gastric mucosa, the use of "enteric" coatings is indicated. Such coatings are intended to delay the release of the medication until the tablet has passed through the stomach. Drug products in delayed-release dosage forms containing the same active ingredients are subject to significant differences in absorption. Unless otherwise specifically noted, the Agency considers different delayed-release products containing the same active ingredients as presenting a potential bioequivalence problem and codes these products **BE** in the absence of *in vivo* studies showing bioequivalence. If adequate *in vivo* studies have demonstrated the bioequivalence of specific delayed-release products, such products are coded **AB**.

BN Products in aerosol-nebulizer drug delivery systems

This code applies to drug solutions or powders that are marketed only as a component of, or as compatible with, a specific drug delivery system. There may, for example, be significant differences in the dose of drug and particle size delivered by different products of this type. Therefore, the Agency does not consider different metered aerosol dosage forms containing the same active ingredient(s) in equal strengths to be therapeutically equivalent unless the drug products meet an appropriate bioequivalence standard; such products are coded **AB**.

BP Active ingredients and dosage forms with potential bioequivalence problems

FDA's bioequivalence regulations (21 CFR 320.33) contain criteria and procedures for determining whether a specific active ingredient in a specific dosage form has a potential for causing a bioequivalence problem. It is FDA's policy to consider an ingredient meeting these criteria as having a potential bioequivalence problem even in the absence of positive data demonstrating inequivalence. Pharmaceutically equivalent products containing these ingredients in oral dosage forms are coded **BP** until adequate bioequivalence data are submitted, after which such products are coded **AB**. Injectable suspensions containing an active ingredient suspended in an aqueous or oleaginous vehicle have also been coded **BP**. Injectable suspensions are subject to bioequivalence problems because differences in particle size, polymorphic structure of the suspended active ingredient, or the suspension formulation can significantly affect the rate of release and absorption. FDA does not consider pharmaceutical equivalents of these products bioequivalent without adequate evidence of bioequivalence; such products would be coded **AB**.

BR Suppositories or enemas that deliver drugs for systemic absorption

The absorption of active ingredients from suppositories or enemas that are intended to have a systemic effect (as distinct from suppositories administered for local effect) can vary significantly from product to product. Therefore, FDA considers pharmaceutically equivalent systemic suppositories or enemas bioequivalent only if *in vivo* evidence of bioequivalence is available. In those cases where *in vivo* evidence is

available, the products are coded **AB**. If such evidence is not available, the products are coded **BR**.

BS Products having drug standard deficiencies

If the drug standards for an active ingredient in a particular dosage form are found by FDA to be deficient so as to prevent an FDA evaluation of either pharmaceutical or therapeutic equivalence, all drug products containing that active ingredient in that dosage form are coded **BS**. For example, if the standards permit a wide variation in pharmacologically active components of the active ingredient such that pharmaceutical equivalence is in question, all products containing that active ingredient in that dosage form are coded **BS**.

BT Topical products with bioequivalence issues

This code applies mainly to post-1962 dermatologic, ophthalmic, otic, rectal, and vaginal products for topical administration, including creams, gels, lotions, oils, ointments, pastes, solutions, sprays, suppositories, and inserts not intended for systemic drug absorption. Topical products evaluated as having acceptable clinical performance, but that are not bioequivalent to other pharmaceutically equivalent products or that lack sufficient evidence of bioequivalence, will be coded **BT**.

BX Drug products for which the data are insufficient to determine therapeutic equivalence

The code **BX** is assigned to specific drug products for which the data that have been reviewed by the Agency are insufficient to determine therapeutic equivalence under the policies stated in this document. In these situations, the drug products are presumed to be therapeutically inequivalent until the Agency has determined that there is adequate information to make a full evaluation of therapeutic equivalence.

1.8 Description of Certain Special Situations

Certain drugs listed in the Orange Book present special situations that merit further discussion. The following are descriptions of certain examples of those special situations:

Amino Acid and Protein Hydrolysate Injections. These products differ in the amount and kinds of amino acids they contain and, therefore, are not considered pharmaceutical equivalents. For this reason, these products are not considered therapeutically equivalent. At the same time, the Agency believes that it is appropriate to point out that where nitrogen balance is the sole therapeutic objective and individual amino acid content is not a consideration, pharmaceutical alternatives with the same total amount of nitrogen content may be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.

Gaviscon®. Gaviscon® is an OTC product that has been marketed since September 1970. The active ingredients in this product, aluminum hydroxide and magnesium trisilicate, were reviewed by the Agency's OTC Antacid Panel and were considered to be safe and effective ingredients (Category I) by that Panel. However, the tablet failed to pass the antacid test that is required of all antacid products. The Agency, therefore, placed the tablet in Category III for lack of effectiveness. A full NDA with clinical studies was

submitted by Marion Laboratories, Inc., and approved by FDA on December 9, 1983. Gaviscon®'s activity in treating reflux acidity is made possible by the physical-chemical properties of the inactive ingredients, sodium bicarbonate and alginic acid. Therefore, *all ANDAs that cite Gaviscon® tablets as the reference listed drug must contain the inactive ingredients sodium bicarbonate and alginic acid.* A full NDA will be required to support the effectiveness of the drug product if different inactive ingredients are to be substituted for sodium bicarbonate or alginic acid or if different proportions of these ingredients are to be used.

Levothyroxine Sodium.¹⁸ Because there are multiple reference listed drugs for levothyroxine sodium tablets and some reference listed drugs' sponsors have conducted studies to establish their drugs' therapeutic equivalence to other reference listed drugs, FDA has determined that its usual practice of assigning two- or three-character therapeutic equivalence codes may be potentially confusing and inadequate for these drug products. Looking at the Orange Book listing alone for a product identified as a reference listed drug or reference standard, it may be difficult to determine to which therapeutic equivalence code the reference listed drugs and/or reference standard designation corresponds. For example, Unithroid 0.3 mg strength has been assigned the therapeutic equivalence codes AB1, AB2, and AB3 and it is identified as the reference listed drug and reference standard, but it is unclear that the reference listed drug and reference standard designations are associated with the AB1 therapeutic equivalence code.

Accordingly, FDA provides the following chart, which identifies (1) a reference listed drug for each therapeutic equivalence code in the Orange Book and (2) the reference standard products in the Active Section of the Orange Book.¹⁹

- Therapeutic equivalence has been established between products that have the same AB+number therapeutic equivalence code (i.e., AB1, AB2, AB3 or AB4).
- More than one therapeutic equivalence code may apply to some products. One common therapeutic equivalence code indicates therapeutic equivalence between products. For example, Unithroid has been assigned therapeutic equivalence codes AB1, AB2, and AB3, and therefore, Unithroid tablets are considered therapeutically equivalent to other levothyroxine sodium products of the same strength with these therapeutic equivalence codes.

TE Code	Proprietary Name	Applicant	Strength	Appl No	RLD	RS
AB1	UNITHROID	STEVENS J	0.3 MG	N021210	RLD	RS
AB2	SYNTHROID	ABBVIE	0.3 MG	N021402	RLD	RS

¹⁸ In previous editions of the Orange Book, FDA provided a chart outlining therapeutic equivalence codes for all 0.025 mg levothyroxine sodium drug products in the Active Section of the Orange Book. FDA has decided, for ease of review, to revise the chart to identify the NDAs for the reference listed drugs for each therapeutic equivalence code (i.e., AB1, AB2, AB3, and AB4), and their corresponding reference standards, which are identified in 0.2 and 0.3 mg strengths.

¹⁹ The chart is current as of the date of publication of the annual edition. See the most current monthly cumulative supplement for updates to this information available at <https://www.fda.gov/media/72973/download>. Please consult the Active Section for information on other strengths.

AB3	LEVOXYL	KING PHARMS	0.2 MG	N021301	RLD	RS
AB4	THYRO-TABS	ALVOGEN INC	0.3 MG	N021116	RLD	-
AB4	LEVOTHYROXINE SODIUM ²⁰	MYLAN	0.3 MG	A076187	-	RS

Patent Certification(s) and Reference Standard for ANDAs Duplicating a Drug Product Approved in a Petitioned ANDA. To submit an ANDA for a generic drug that is not the same as its reference listed drug because it has one different active ingredient in a fixed-combination drug product, or has a different route of administration, dosage form, or strength than that of the reference listed drug, an applicant first must obtain permission from FDA through what is known as a suitability petition pursuant to Section 505(j)(2)(C) of the FD&C Act. A petitioned ANDA relies on the reference listed drug described in the suitability petition. An ANDA seeking approval of a drug that is the same as a drug product approved in a petitioned ANDA should use as its reference listed drug, the reference listed drug that served as the basis for the approved suitability petition, and use the drug product approved in the petitioned ANDA as its reference standard for conducting an *in vivo* bioequivalence study required for approval. However, the reference listed drug for any such ANDA is generally the listed drug referenced in the approved suitability petition. The ANDA must include appropriate patent certification(s) and an exclusivity statement with respect to the reference listed drug that served as the basis for the approved suitability petition.²¹ (This concept also generally applies to an ANDA applicant that utilizes a reference standard that is not a reference listed drug, as such an application must include appropriate patent certification(s) and an exclusivity statement with respect to the reference listed drug.)

Waived exclusivity. If an NDA submitted under Section 505(b) of the FD&C Act qualifies for exclusivity under the FD&C Act, the exclusivity is generally listed in the Patent and Exclusivity Section of the Orange Book. If a drug product has qualified for this exclusivity, FDA will not accept for review and/or will not approve, as applicable, other applications blocked by the relevant exclusivity. If the listed drug is also protected by one or more patents, the approval date for an ANDA or 505(b)(2) application that relies on the listed drug will be determined based on an analysis of the applicant's patent certification(s) or statement(s) for each relevant patent and the effect of relevant exclusivity listed in the Orange Book. However, the

²⁰ Alvogen, Inc.'s tablets (NDA 021116) (previously known as Levotheroid) previously was listed in the Discontinued Drug Product List section of the Orange Book. It is the RLD for therapeutic equivalents identified with the AB4 code. During this time, Mylan's levothyroxine product (ANDA 076187) was selected as the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. It remains the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. If an ANDA that uses Mylan's levothyroxine product as its reference standard is approved, the ANDA will receive an AB4 rating. The ANDA applicant also may obtain an AB rating for its product to the other reference listed drugs (i.e., Unithroid, Synthroid, and Levoxyl) by submitting supplements that demonstrate that the generic product is bioequivalent to these other reference listed drugs and satisfies all other therapeutic equivalence criteria with respect to these reference listed drugs. See Letter from Janet Woodcock, M.D., Director, Center for Drug Evaluation and Research, FDA to Teri Nataline, Principal Consultant, Lachman Consultant Services, Inc., Docket No. FDA-2015-P-0403 (May 27, 2016).

²¹ If after approval of a suitability petition and before approval of an ANDA submitted pursuant to the approved petition, a drug product is approved in an NDA for the change described in the petition, the suitability petition and the listed drug identified in the petition can no longer be the basis of submission for such ANDA. Under these circumstances, an applicant seeking approval for a drug product with the change approved in the suitability petition must submit a new ANDA that identifies the drug product approved under such NDA as the RLD and comply with applicable regulatory requirements. See 21 CFR 314.93(f)(2).

holder of the NDA may waive its exclusivity as to any or all applications that might otherwise be blocked by such exclusivity. If an NDA holder waives its exclusivity, qualified applications may be accepted for review and/or approved, as applicable. An NDA for which the holder has waived its exclusivity as to all applications will be coded with a "W" in the Patent and Exclusivity Section of the Orange Book. The applicant whose product might otherwise be blocked by this exclusivity should indicate in the exclusivity statement in its application that the holder of the listed drug has waived its exclusivity.

1.9 Therapeutic Equivalence Code Change for a Category of Multisource Drug Products

The Agency will use the following procedures when, in response to a petition or on its own initiative, it is considering a change in the therapeutic equivalence code for approved multisource drug products. Such changes will generally occur when the Agency becomes aware of new scientific information affecting the therapeutic equivalence of an entire category of multisource drug products in the Orange Book (e.g., information concerning the active ingredient or the dosage form), rather than information concerning a single drug product within the category. These procedures will be used when a change in therapeutic equivalence code is under consideration for all drug products found in the Prescription Drug Product List under a specific active ingredient and dosage form. The change may be from the code signifying that the drug does not present a bioequivalence problem (e.g., **AA**) to a code signifying an actual or potential bioequivalence problem (e.g., **BP**), or vice versa. This procedure does not apply to a change of a particular product code (e.g., a change from **BP** to **AB** or from **AB** to **BX**).

Before making a change in a therapeutic equivalence code for an entire category of multisource drug products as described above, the Agency will announce in the *Introduction* to the Cumulative Supplement that it is considering the change and will invite comments. Comments, along with scientific data, may be sent to the Director, Office of Bioequivalence, Office of Generic Drugs, Center for Drug Evaluation and Research, Food and Drug Administration, Central Document Room, 5901-B Ammendale Rd., Beltsville, MD 20705-1266.

The comment period will generally be 60 days in length, and the closing date for comments will be listed in the description of the proposed change for each drug entity.

The most useful type of scientific data submitted to support comments is generally an *in vivo* bioavailability/bioequivalence study conducted on batches of the subject drug products. Comments including scientific data from an *in vivo* bioavailability/bioequivalence study should present a full description of the analytical procedures and equipment used, a validation of the analytical methodology, including the standard curve, a description of the method of calculating results, and a description of the pharmacokinetic and statistical models used in analyzing the data. Anecdotal or testimonial information is the least useful to the Agency, and submission of comments based on such information is discouraged. However, when there is supporting published or unpublished scientific literature, copies should be submitted with comments.

1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product

The procedure described in Section 1.9 does not apply to a change in a single drug product code. For example, a change in a single drug product's

code from **BP** to **AB** as a result of the submission of an acceptable bioequivalence study ordinarily will not be the subject of notice and comment in the Cumulative Supplement. Likewise, a change in a single drug product's code from **AB** to **BX** (e.g., as a result of new information raising a significant question as to bioequivalence) does not require notice and comment. The Agency's responsibility to provide the public with the Agency's most current information related to therapeutic equivalence may require a change in a drug product's code prior to any formal notice and opportunity for the applicant to be heard. The publication in the *Federal Register* of a proposal to withdraw approval of a drug product will ordinarily result in a change in a product's code from **AB** to **BX** if this action has not already been taken.

We recognize that certain drug products approved in 505(b)(2) applications may not have therapeutic equivalence codes, and that FDA may undertake therapeutic equivalence evaluations with respect to such drug products. A person seeking to have a therapeutic equivalence rating for a drug product approved in a 505(b)(2) application may petition the Agency through the citizen petition procedure (see 21 CFR 10.25(a) and 21 CFR 10.30).²²

1.11 Discontinued Section

Those drug products in the discontinued section of the Orange Book (Discontinued Drug Product List) for which a determination has been made that the products were not withdrawn for safety or effectiveness reasons have been annotated with a footnote following the product strength: "***Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons***". The determinations listed in the Orange Book are only reflective of determinations made since 1995 and published in the Federal Register. The identification of these drug products in the Discontinued Drug Product List should avoid the submission of multiple citizen petitions requesting a determination for the same drug product.

Generally, approved products are added to the Discontinued Drug Product List when the applicant notifies the Division of Orange Book Publication and Regulatory Assessment (DOBPR) of the products' not-marketed status. Products may also be added to the Discontinued Drug Product List if annual reports or other submissions to the Agency indicate the product is not being marketed or as a result of other Agency administrative actions.²³ Changes to the Orange Book are not affected by the drug registration and listing requirements of Section 510 of the FD&C Act.

1.12 Changes to the Orange Book

Every effort is made to ensure the Annual Edition is current and accurate. Applicants are requested to inform DOBPR of any changes or corrections, including any change in ownership or a product's marketing status that would result in the product being moved to the Discontinued Drug Product List. FDA notes that under Section 506I(a) of the FD&C Act, application holders must notify the Agency in writing 180 days prior to withdrawing a drug product from sale, or if 180 days is not practicable, not later than the date of withdrawal from sale. Furthermore, Section 506I(b) of

²² Section 3222 of the Food and Drug Omnibus Reform Act of 2022 (enacted December 29, 2022) amended the FD&C Act by adding a new provision to Section 505(j)(7)(A). Section 505(j)(7)(A)(v)(I) sets forth certain conditions under which FDA considers therapeutic equivalence evaluation requests in an application for an eligible drug submitted or approved pursuant to Section 505(b)(2) of the FD&C Act.

²³ See, e.g., Section 506I(d) of the FD&C Act.

the FD&C Act requires that application holders notify the Agency in writing within 180 days of approval of a drug product if such drug product will not be available for sale within 180 days of approval. A request to include a newly approved product in the Discontinued Drug Product List, rather than parts 1 or 2 of the Orange Book (as discussed in Section 1.1), must be submitted to DOBPPRA by the end of the month in which the product is approved to ensure that the product is not included in the Active Section of the next published Orange Book update. The Prescription Drug Product List and the Over-the-Counter (OTC) Drug Product List are collectively referred to as the "Active Section."

In addition, DOBPPRA generally will act on requests to change a proprietary name for a listed drug only after approval of a supplement for the relevant change in proprietary name. To the extent that conventions for describing product identification information (i.e., active ingredients, dosage forms, routes of administration, product names, applicants, strengths) evolve over time, the Agency generally does not intend to revise such information for drug products already listed in the Orange Book, but rather intends to apply the change prospectively to drug products as they are added to the Orange Book.

You can contact DOBPPRA by email at orangebook@fda.hhs.gov.

1.13 Availability of the Edition

The Annual Edition and current monthly Cumulative Supplement are available in a Portable Document Format (PDF) at the [Orange Book](#) home page by clicking on Publications. An annual subscription of the PDF format may be obtained from the U.S. Government Publishing Office, <https://www.gpo.gov/>.

2.0 HOW TO USE THE DRUG PRODUCT LISTS

2.1 Key Sections for Using the Drug Product Lists

This publication contains illustrations, along with Drug Product Lists, indices, and lists of abbreviations and terms which facilitate their use.

Illustrations. The annotated *Drug Product Illustration* (see Section 2.2) and the *Therapeutic Equivalence Evaluations Illustration* (see Section 2.3) are offered to provide further clarification. These depict the format found in the Prescription Drug Product List (the only list in which therapeutic equivalence evaluation codes are displayed).

Drug Product Lists. The Prescription and OTC Drug Product Lists, arranged alphabetically by active ingredient(s), contain product identification information (active ingredients, dosage forms, routes of administration, product names, applicants, strengths) for single and multiple ingredient drug products. Also shown are the application number and drug product number (FDA internal computer data use only) and approval dates for those drug products approved on or after January 1, 1982. The application number preceded by "N" is a New Drug Application (NDA or commonly the innovator). The application number preceded by an "A" is an Abbreviated New Drug Application (ANDA or commonly the generic).

The Discontinued Drug Product List, arranged alphabetically by active ingredient(s), contains product identification information (active ingredients, dosage forms, routes of administration, product names, applicants, strengths).

If a prescription drug product is available from more than one source (multisource), a therapeutic equivalence code will appear in front of the applicant's name. If a product is therapeutically equivalent to one or more products or to an appropriate reference, it will be designated with a code beginning with "A" and the entry will be underlined and printed in bold font for emphasis.

Active ingredient headings for multiple ingredient (combination) drug products are arranged alphabetically. For purposes of this publication, this alphabetical sort takes precedence over United States Pharmacopeia official monograph order (i.e., Reserpine, Hydralazine Hydrochloride, Hydrochlorothiazide). For example, product information labeled as Reserpine, Hydrochlorothiazide and Hydralazine Hydrochloride appears under the active ingredient heading *Hydralazine Hydrochloride; Hydrochlorothiazide; Reserpine*. A cross-reference to the product information (for prescription and OTC products) appears for each additional active ingredient in the product. For combination drug products, the ingredient strengths are separated by semicolons and appear in the same relative sequence as the ingredients in the heading. Available strengths of the dosage form from an applicant appear on separate lines.

To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if necessary. Then, find the ingredient in the applicable Drug Product List. Proceed to the dosage form and route of administration and

compare products within that ingredient heading only. Therapeutic equivalence or inequivalence for prescription products is determined on the basis of the therapeutic equivalence codes provided within that specific dosage form and route heading. The OTC Drug Product List, Discontinued Drug Product List, and Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research List have their data arranged similarly.

The Discontinued Drug Product List contains approved products that have never been marketed, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons, are for military use, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing. All products having a "@" in the December Cumulative Supplement of the previous Edition List have been added to the Discontinued Drug Product List appearing in this Edition. In addition, approved drug products that are not in the commercial distribution channel e.g., approved drug products in applications for export only are also listed in the Discontinued Drug Product List.

Product Name Index (*Prescription and OTC Drug Product Lists*). This is an index of drug products by trade name or established name of the active ingredient, if no trade name exists. The second term of each entry indicates the active ingredient name under which product information can be found in the appropriate Drug Product List. For those drug products with multiple active ingredients, only the first active ingredient (in alphabetical order) will appear. OTC products are so designated.

Product Name Index Listed by Applicant (*Prescription and OTC Drug Product Lists*). This is an index that cross-references applicants to drug products. The bolded and underlined entry represents the applicant name abbreviation used in this publication. Each complete applicant name that is represented by the abbreviated name is marked with an asterisk (*). Listed under each complete applicant name is the first alphabetically arranged ingredient under which product information can be found in the appropriate Drug Product List.

OTC products are so designated. To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if appropriate.

Uniform Terms. To improve readability, uniform terms are used to designate dosage forms, routes of administration, and abbreviations used to express strengths. These terms are listed in Appendix C. In some cases, the terms used may differ from those used in product labels and other labeling.

2.2 DRUG PRODUCT ILLUSTRATION

SINGLE INGREDIENT

ACTIVE INGREDIENT	→	<u>MEPERIDINE HYDROCHLORIDE</u>					
DOSAGE FORM; ROUTE OF ADMINISTRATION	→	INJECTABLE; INJECTION					
TRADE OR GENERIC NAMES	→	<u>HEXANON</u>					
REFERENCE LISTED DRUG* (+)	→	<u>AP</u> +!	PAGE PHARMA	<u>25MG/ML</u>	<u>N013111</u>	<u>001</u>	AUG 22, 1983
REFERENCE STANDARD * (!)	→	<u>AP</u> +!		<u>50MG/ML</u>	<u>N013111</u>	<u>002</u>	AUG 22, 1983
	→	<u>AP</u> +!		<u>75MG/ML</u>	<u>N013111</u>	<u>003</u>	AUG 22, 1983
	→	<u>AP</u> +!		<u>100MG/ML</u>	<u>N013111</u>	<u>004</u>	JAN 04, 1989
	→	<u>MEPERIDINE HCL</u>					
THERAPEUTIC EQUIVALENCE (TE)	→	<u>AP</u>	DAVIS PHARM	<u>25MG/ML</u>	<u>A064890</u>	001	FEB 29, 1987
CODE FOR MULTISOURCE PRODUCT	→	<u>AP</u>		<u>50MG/ML</u>	<u>A064890</u>	002	FEB 29, 1987
	→	<u>AP</u>		<u>75MG/ML</u>	<u>A064890</u>	003	FEB 29, 1987
	→	<u>AP</u>		<u>100MG/ML</u>	<u>A064890</u>	004	MAR 08, 1992
SINGLE SOURCE PRODUCT (NO TE CODE)	→		! TIMOKIM LLC	10MG/ML	A099225	001	DEC 12, 1995
	→	<u>AP</u>	JOHNSON MED	<u>25MG/ML</u>	<u>A099226</u>	<u>001</u>	NOV 27, 1993
	→		! KENDRA PHARM	150MG/ML	A079444	001	OCT 31, 1999
APPLICANT	→						
AVAILABLE STRENGTH(S) OF A PRODUCT	→						
APPLICATION NUMBER	→						
PRODUCT NUMBER IS FOR FDA INTERNAL COMPUTER DATA USE ONLY	→						
APPROVAL DATE	→						

*NOTE: REFERENCE LISTED DRUG AND REFERENCE STANDARD ARE DISCUSSED IN THE PREFACE SECTION 1.4

MULTIPLE INGREDIENTS WITH PRODUCT INFORMATION

ALPHABETICALLY SORTED BY		
ACTIVE INGREDIENT	→	<u>HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE</u>
PRODUCT INFORMATION	→	TABLET; ORAL HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL REINWALD LABS 25MG; 15MG; 0.1MG A069808 001 JAN 18, 1982

THIS EXAMPLE IS FOR PURPOSE OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

2.3 THERAPEUTIC EQUIVALENCE EVALUATIONS ILLUSTRATION

DRUG PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") UNDER AN INGREDIENT AND DOSAGE FORM HEADING ARE CONSIDERED THERAPEUTICALLY EQUIVALENT ONLY TO OTHER PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") AND **NOT** TO THOSE CODED **BP** (OR ANY CODE BEGINNING WITH "B") AND ANY PRODUCTS NOT LISTED. DRUG PRODUCTS CODED **BP** (OR ANY CODE BEGINNING WITH A "B") ARE **NOT** CONSIDERED THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCT. FOR A COMPLETE EXPLANATION OF THE **TE** CODES REFER TO SECTION 1.7 OF THE *INTRODUCTION*.

SULFASALAZINE

TABLET; ORAL

FAZINE

AB PARKLAND **500MG** **A042999** **001**

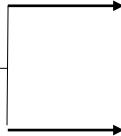
SULAZINE

AB URSA **500MG** **A042222** **001**

SULFASALAZINE

BP BROWN 500MG A041297 001

PRODUCTS CONSIDERED THERAPEUTICALLY EQUIVALENT TO EACH OTHER



PRODUCTS CONSIDERED **NOT** THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCTS LISTED

SULFASALAZINE

TABLET; ORAL

FAZINE

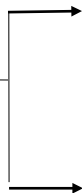
AB PARKLAND **500MG** **A042999** **001**

SULFASALAZINE

BP BROWN 500MG A041297 001

SOUTH 500MG A067627 001

PRODUCTS CONSIDERED **NOT** THERAPEUTICALLY EQUIVALENT TO EACH OTHER



NOTE: BOLD FONT AND UNDERLINING DENOTES MULTISOURCE PRODUCTS WHICH ARE CONSIDERED THERAPEUTICALLY EQUIVALENT.

THIS EXAMPLE IS FOR PURPOSES OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

PRESCRIPTION DRUG PRODUCT LIST

ABACAVIR SULFATE

SOLUTION;ORAL

ABACAVIR SULFATE

AA	AUROBINDO PHARMA LTD	EQ 20MG BASE/ML	A077950 001	Mar 14, 2018
AA	HETERO LABS LTD III	EQ 20MG BASE/ML	A201107 001	Sep 26, 2016

ZIAGEN

AA	+! VIIV HLTHCARE	EQ 20MG BASE/ML	N020978 001	Dec 17, 1998
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TABLET;ORAL

ABACAVIR SULFATE

AB	AUROBINDO PHARMA LTD	EQ 300MG BASE	A077844 001	Dec 17, 2012
AB	CIPLA	EQ 300MG BASE	A078119 001	Nov 21, 2017
AB	! HETERO LABS LTD III	EQ 300MG BASE	A091560 001	Sep 13, 2013
AB	MYLAN PHARMS INC	EQ 300MG BASE	A091294 001	Jun 18, 2012
AB	STRIDES PHARMA	EQ 300MG BASE	A091050 001	Oct 28, 2016

ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

TRIUMEQ

+!	VIIV HLTHCARE	EQ 600MG BASE;EQ 50MG BASE;300MG	N205551 001	Aug 22, 2014
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TABLET, FOR SUSPENSION;ORAL

TRIUMEQ PD

+!	VIIV HLTHCARE	EQ 60MG BASE;EQ 5MG BASE;30MG	N215413 001	Mar 30, 2022
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ABACAVIR SULFATE; LAMIVUDINE

TABLET;ORAL

ABACAVIR SULFATE AND LAMIVUDINE

AB	AUROBINDO PHARMA LTD	EQ 600MG BASE;300MG	A090159 001	Nov 15, 2018
AB	CIPLA	EQ 600MG BASE;300MG	A091144 001	Mar 28, 2017
AB	LAURUS	EQ 600MG BASE;300MG	A216332 001	Jul 25, 2022
AB	LUPIN LTD	EQ 600MG BASE;300MG	A204990 001	Mar 28, 2017

EPZICOM

AB	+! VIIV HLTHCARE	EQ 600MG BASE;300MG	N021652 001	Aug 02, 2004
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TABLET, FOR SUSPENSION;ORAL

ABACAVIR SULFATE AND LAMIVUDINE

+!	MYLAN LABS LTD	EQ 60MG BASE;30MG	N204311 001	Dec 22, 2023
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ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE

!	LUPIN LTD	EQ 300MG BASE;150MG;300MG	A202912 001	Dec 05, 2013
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ABALOPARATIDE

SOLUTION;SUBCUTANEOUS

TYMLOS

+!	RADIUS	3.12MG/1.56ML (2MG/ML)	N208743 001	Apr 28, 2017
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ABEMACICLIB

TABLET;ORAL

VERZENIO

+	ELI LILLY AND CO	50MG	N208716 001	Sep 28, 2017
+		100MG	N208716 002	Sep 28, 2017
+		150MG	N208716 003	Sep 28, 2017
+!		200MG	N208716 004	Sep 28, 2017

ABIRATERONE ACETATE

TABLET;ORAL

ABIRATERONE ACETATE

AB	AMNEAL PHARMS	250MG	A208327 001	Jan 07, 2019
AB		500MG	A208327 002	Dec 23, 2020
AB	APOTEX	250MG	A208453 001	Oct 31, 2018
AB	DR REDDYS	250MG	A208416 001	May 18, 2020
AB		500MG	A208416 002	Sep 01, 2023
AB	FLORIDA	500MG	A215086 001	Mar 23, 2023
AB	GLENMARK PHARMS	250MG	A209227 001	Oct 16, 2019
AB		500MG	A209227 002	May 19, 2022
AB	HIKMA	250MG	A208339 001	Oct 31, 2018
AB	MSN	250MG	A210686 001	Jul 10, 2019
AB	MYLAN	250MG	A208446 001	Oct 31, 2018
AB		500MG	A208446 002	Dec 14, 2020
AB	NOVUGEN	250MG	A215947 001	Jan 05, 2022
AB		500MG	A215947 002	Jan 05, 2022
AB	QILU	250MG	A212462 001	Sep 27, 2019
AB		500MG	A212462 002	Jun 25, 2021

PRESCRIPTION DRUG PRODUCT LIST

ABIRATERONE ACETATE

TABLET; ORAL

ABIRATERONE ACETATE

<u>AB</u>	RISING	<u>250MG</u>	<u>A208371</u>	<u>001</u>	Feb 25, 2019
<u>AB</u>	TEVA PHARMS USA	<u>250MG</u>	<u>A208432</u>	<u>001</u>	Oct 31, 2018
<u>AB</u>		<u>500MG</u>	<u>A210726</u>	<u>001</u>	Jan 26, 2023
<u>AB</u>	WOCKHARDT BIO AG	<u>250MG</u>	<u>A208380</u>	<u>001</u>	Feb 27, 2019
<u>ZYTIGA</u>					
<u>AB</u>	+ JANSSEN BIOTECH	<u>250MG</u>	<u>N202379</u>	<u>001</u>	Apr 28, 2011
<u>AB</u>	+!	<u>500MG</u>	<u>N202379</u>	<u>002</u>	Apr 14, 2017
YONSA					
	+!	SUN PHARM	125MG	N210308	001 May 22, 2018

ABIRATERONE ACETATE; NIRAPARIB TOSYLATE

TABLET; ORAL

AKEEGA

	+ JANSSEN BIOTECH	500MG;EQ 50MG BASE	N216793	001	Aug 11, 2023
	+!	500MG;EQ 100MG BASE	N216793	002	Aug 11, 2023

ABROCITINIB

TABLET; ORAL

CIBINQO

	+ PFIZER	50MG	N213871	001	Jan 14, 2022
	+	100MG	N213871	002	Jan 14, 2022
	+!	200MG	N213871	003	Jan 14, 2022

ACALABRUTINIB

CAPSULE; ORAL

CALQUENCE

	+! ASTRAZENECA	100MG	N210259	001	Oct 31, 2017
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ACALABRUTINIB MALEATE

TABLET; ORAL

CALQUENCE

	+! ASTRAZENECA	EQ 100MG BASE	N216387	001	Aug 03, 2022
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ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL

ACAMPROSATE CALCIUM

<u>AB</u>	! GLENMARK GENERICS	<u>333MG</u>	<u>A202229</u>	<u>001</u>	Jul 16, 2013
<u>AB</u>	MYLAN	<u>333MG</u>	<u>A200142</u>	<u>001</u>	Mar 11, 2014
<u>AB</u>	ZYDUS PHARMS	<u>333MG</u>	<u>A205995</u>	<u>001</u>	May 26, 2017

ACARBOSE

TABLET; ORAL

ACARBOSE

<u>AB</u>	AVET LIFESCIENCES	<u>25MG</u>	<u>A202271</u>	<u>001</u>	Feb 07, 2012
<u>AB</u>		<u>50MG</u>	<u>A202271</u>	<u>002</u>	Feb 07, 2012
<u>AB</u>		<u>100MG</u>	<u>A202271</u>	<u>003</u>	Feb 07, 2012
<u>AB</u>	HIKMA	<u>25MG</u>	<u>A078470</u>	<u>001</u>	May 07, 2008
<u>AB</u>		<u>50MG</u>	<u>A078470</u>	<u>002</u>	May 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A078470</u>	<u>003</u>	May 07, 2008
<u>AB</u>	IMPAX LABS	<u>25MG</u>	<u>A078441</u>	<u>001</u>	May 14, 2009
<u>AB</u>		<u>50MG</u>	<u>A078441</u>	<u>002</u>	May 14, 2009
<u>AB</u>		<u>100MG</u>	<u>A078441</u>	<u>003</u>	May 14, 2009
<u>AB</u>	! STRIDES PHARMA	<u>25MG</u>	<u>A090912</u>	<u>001</u>	Jul 27, 2011
<u>AB</u>		<u>50MG</u>	<u>A090912</u>	<u>002</u>	Jul 27, 2011
<u>AB</u>		<u>100MG</u>	<u>A090912</u>	<u>003</u>	Jul 27, 2011
<u>AB</u>	VIRTUS PHARM	<u>25MG</u>	<u>A091343</u>	<u>001</u>	Oct 17, 2013
<u>AB</u>		<u>50MG</u>	<u>A091343</u>	<u>002</u>	Oct 17, 2013
<u>AB</u>		<u>100MG</u>	<u>A091343</u>	<u>003</u>	Oct 17, 2013
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A077532</u>	<u>001</u>	May 07, 2008
<u>AB</u>		<u>50MG</u>	<u>A077532</u>	<u>002</u>	May 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A077532</u>	<u>003</u>	May 07, 2008

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

<u>AB</u>	! AMNEAL PHARM	<u>EQ 200MG BASE</u>	<u>A075047</u>	<u>001</u>	Dec 30, 1999
<u>AB</u>	!	<u>EQ 400MG BASE</u>	<u>A075047</u>	<u>002</u>	Dec 30, 1999
<u>AB</u>	ANI PHARMS	<u>EQ 200MG BASE</u>	<u>A074007</u>	<u>001</u>	Oct 18, 1995
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A074007</u>	<u>002</u>	Oct 18, 1995

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN

SOLUTION; INTRAVENOUS

ACETAMINOPHEN

<u>AP</u>	ASPIRO	<u>1GM/100ML (10MG/ML)</u>	<u>A216617 001</u>	Jul 27, 2022
<u>AP</u>	BAXTER HLTHCARE CORP	<u>1GM/100ML (10MG/ML)</u>	<u>A214331 001</u>	Sep 17, 2021
<u>AP</u>	EUGIA PHARMA	<u>1GM/100ML (10MG/ML)</u>	<u>A210969 001</u>	Oct 21, 2020
<u>AP</u>	HIKMA	<u>1GM/100ML (10MG/ML)</u>	<u>A202605 001</u>	Jun 13, 2016
<u>AP</u>	INFORLIFE	<u>1GM/100ML (10MG/ML)</u>	<u>A215403 001</u>	May 03, 2023
<u>AP</u>	MYLAN	<u>1GM/100ML (10MG/ML)</u>	<u>A213255 001</u>	Aug 07, 2020
<u>AP</u>	! SANDOZ	<u>1GM/100ML (10MG/ML)</u>	<u>A204052 001</u>	Mar 22, 2016
<u>AP</u>	WOCKHARDT BIO AG	<u>1GM/100ML (10MG/ML)</u>	<u>A205746 001</u>	Apr 18, 2023
+!	B BRAUN MEDICAL INC	500MG/50ML (10MG/ML)	N204957 001	Feb 18, 2021
+!		1GM/100ML (10MG/ML)	N204957 002	Feb 18, 2021
	FRESENIUS KABI USA	1GM/100ML (10MG/ML)	N204767 001	Oct 28, 2015
+!	HIKMA	1GM/100ML (10MG/ML)	N206968 001	Jun 03, 2022

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BUTALBITAL AND ACETAMINOPHEN

<u>AA</u>	! DR REDDYS LABS SA	<u>300MG; 50MG</u>	<u>A207313 001</u>	Dec 27, 2017
<u>AA</u>	GRANULES	<u>300MG; 50MG</u>	<u>A213115 001</u>	Nov 22, 2019

TABLET; ORAL

ALLZITAL

<u>AA</u>	! LARKEN LABS INC	<u>325MG; 25MG</u>	<u>A203484 001</u>	Dec 04, 2015
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BUTALBITAL AND ACETAMINOPHEN

<u>AA</u>	LARKEN LABS INC	<u>325MG; 50MG</u>	<u>A203484 002</u>	Dec 04, 2015
<u>AA</u>	! LGM PHARMA	<u>300MG; 50MG</u>	<u>A090956 001</u>	Aug 23, 2011
<u>AA</u>	MIKART	<u>300MG; 50MG</u>	<u>A207386 001</u>	Nov 15, 2016
<u>AA</u>	NE RX PHARMA	<u>300MG; 50MG</u>	<u>A214955 001</u>	Aug 16, 2022
<u>AA</u>	SENORES PHARMS	<u>300MG; 50MG</u>	<u>A214088 001</u>	Apr 07, 2022
<u>AA</u>		<u>325MG; 25MG</u>	<u>A214088 002</u>	Apr 07, 2022
<u>AA</u>		<u>325MG; 50MG</u>	<u>A214088 003</u>	Apr 07, 2022

BUTAPAP

<u>AA</u>	! MIKART	<u>325MG; 50MG</u>	<u>A089987 001</u>	Oct 26, 1992
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ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AA</u>	AUROLIFE PHARMA LLC	<u>325MG; 50MG; 40MG</u>	<u>A204733 001</u>	Sep 26, 2018
<u>AA</u>	DR REDDYS LABS SA	<u>300MG; 50MG; 40MG</u>	<u>A210817 001</u>	Dec 17, 2019
<u>AA</u>	! GRANULES	<u>325MG; 50MG; 40MG</u>	<u>A089007 001</u>	Mar 17, 1986
<u>AA</u>	GRANULES	<u>300MG; 50MG; 40MG</u>	<u>A213321 001</u>	Apr 08, 2020
<u>AA</u>	LANNETT CO INC	<u>300MG; 50MG; 40MG</u>	<u>A212082 001</u>	Dec 17, 2019
<u>AA</u>		<u>325MG; 50MG; 40MG</u>	<u>A212083 001</u>	Dec 17, 2019
<u>AA</u>	! LGM PHARMA	<u>300MG; 50MG; 40MG</u>	<u>A040885 001</u>	Nov 16, 2009
<u>AA</u>	NOVAST LABS	<u>300MG; 50MG; 40MG</u>	<u>A215047 001</u>	Nov 17, 2021
<u>AA</u>	NUVO PHARMS INC	<u>300MG; 50MG; 40MG</u>	<u>A207118 001</u>	Oct 28, 2016
<u>AA</u>	RISE PHARMA	<u>300MG; 50MG; 40MG</u>	<u>A214087 001</u>	Aug 13, 2021
<u>AA</u>		<u>325MG; 50MG; 40MG</u>	<u>A214087 002</u>	Aug 13, 2021
<u>AA</u>	TARO	<u>300MG; 50MG; 40MG</u>	<u>A213046 001</u>	Jul 01, 2020
<u>AA</u>	XSPIRE PHARMA	<u>300MG; 50MG; 40MG</u>	<u>A206615 001</u>	Aug 04, 2017

SOLUTION; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

!	MIKART	325MG/15ML; 50MG/15ML; 40MG/15ML	A040387 001	Jan 31, 2003
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TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AA</u>	ABHAI LLC	<u>325MG; 50MG; 40MG</u>	<u>A211106 001</u>	Sep 26, 2018
<u>AA</u>	+ ACTAVIS LABS UT INC	<u>325MG; 50MG; 40MG</u>	<u>A088616 001</u>	Nov 09, 1984
<u>AA</u>	GRANULES	<u>325MG; 50MG; 40MG</u>	<u>A040864 001</u>	Dec 01, 2008
<u>AA</u>	LANNETT CO INC	<u>325MG; 50MG; 40MG</u>	<u>A200243 001</u>	Sep 13, 2012
<u>AA</u>	LGM PHARMA	<u>325MG; 50MG; 40MG</u>	<u>A209587 001</u>	Oct 31, 2018
<u>AA</u>	MIKART	<u>325MG; 50MG; 40MG</u>	<u>A089175 001</u>	Jan 21, 1987
<u>AA</u>	! STRIDES PHARMA	<u>325MG; 50MG; 40MG</u>	<u>A040511 001</u>	Aug 27, 2003
<u>AA</u>		<u>325MG; 50MG; 40MG</u>	<u>A203647 001</u>	Sep 21, 2020

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

<u>AB</u>	HIKMA	<u>300MG; 50MG; 40MG; 30MG</u>	<u>A215138 002</u>	Jan 26, 2022
<u>AB</u>		<u>325MG; 50MG; 40MG; 30MG</u>	<u>A215138 001</u>	Jan 26, 2022
<u>AB</u>	LGM PHARMA	<u>300MG; 50MG; 40MG; 30MG</u>	<u>A076560 002</u>	Jul 19, 2012
<u>AB</u>		<u>325MG; 50MG; 40MG; 30MG</u>	<u>A076560 001</u>	Jun 10, 2004
<u>AB</u>	NOSTRUM LABS INC	<u>325MG; 50MG; 40MG; 30MG</u>	<u>A075929 001</u>	Apr 22, 2002

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

AB	STRIDES PHARMA	325MG;50MG;40MG;30MG	A204649 001	Jul 08, 2020
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FIORICET W/ CODEINE

AB	+! ACTAVIS LABS UT INC	325MG;50MG;40MG;30MG	N020232 001	Jul 30, 1992
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ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

TREZIX

	XSPIRE PHARMA	320.5MG;30MG;16MG	A204785 001	Nov 26, 2014
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ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA	AKORN	120MG/5ML;12MG/5ML	A040119 001	Apr 26, 1996
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AA	CHARTWELL	120MG/5ML;12MG/5ML	A089450 001	Oct 27, 1992
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AA	! GENUS LIFESCIENCES	120MG/5ML;12MG/5ML	A087508 001	
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TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA	AMNEAL PHARMS NY	300MG;30MG	A040779 001	May 29, 2008
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AA	AUROLIFE PHARMA LLC	300MG;15MG	A202800 001	Apr 15, 2013
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AA		300MG;30MG	A202800 002	Apr 15, 2013
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AA		300MG;60MG	A202800 003	Apr 15, 2013
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AA	ELITE LABS INC	300MG;15MG	A212418 001	Sep 10, 2019
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AA		300MG;30MG	A212418 002	Sep 10, 2019
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AA		300MG;60MG	A212418 003	Sep 10, 2019
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AA	! SPECGX LLC	300MG;15MG	A040419 001	May 31, 2001
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AA	! SPECGX LLC	300MG;30MG	A040419 002	May 31, 2001
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AA		300MG;60MG	A040419 003	May 31, 2001
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AA	SUN PHARM INDS LTD	300MG;30MG	A085868 001	
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AA		300MG;60MG	A087083 001	
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AA	WES PHARMA INC	300MG;15MG	A211610 001	Jun 27, 2019
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AA		300MG;30MG	A211610 002	Jun 27, 2019
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AA		300MG;60MG	A211610 003	Jun 27, 2019
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ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA	! CHARTWELL	325MG/15ML;7.5MG/15ML	A040482 001	Sep 25, 2003
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AA	GENUS	325MG/15ML;7.5MG/15ML	A040894 001	Jul 19, 2011
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AA	PHARM ASSOC	325MG/15ML;7.5MG/15ML	A040838 001	May 10, 2013
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AA	WES PHARMA INC	325MG/15ML;7.5MG/15ML	A211023 001	Mar 08, 2019
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!	MIKART	300MG/15ML;10MG/15ML	A040881 001	Feb 25, 2010
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!	PHARM ASSOC	325MG/15ML;10MG/15ML	A040834 001	Apr 18, 2008
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TABLET;ORAL

ANEXSIA 5/325

AA	! SPECGX LLC	325MG;5MG	A040409 001	Oct 20, 2000
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ANEXSIA 7.5/325

AA	! SPECGX LLC	325MG;7.5MG	A040405 001	Sep 08, 2000
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HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA	ABHAI LLC	300MG;5MG	A209036 001	Jun 21, 2017
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AA		300MG;7.5MG	A209036 002	Jun 21, 2017
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AA		300MG;10MG	A209036 003	Jun 21, 2017
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AA		325MG;5MG	A209037 001	Jun 21, 2017
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AA		325MG;7.5MG	A209037 002	Jun 21, 2017
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AA		325MG;10MG	A209037 003	Jun 21, 2017
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AA	AMNEAL PHARMS	300MG;10MG	A207137 001	Nov 29, 2016
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AA	AMNEAL PHARMS NY	300MG;5MG	A206869 001	Jun 23, 2017
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AA		325MG;5MG	A040736 001	Aug 25, 2006
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AA		325MG;7.5MG	A040746 002	May 10, 2016
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AA		325MG;10MG	A040746 001	Aug 25, 2006
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AA	ASCENT PHARMS INC	325MG;5MG	A211487 002	Nov 07, 2018
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AA		325MG;7.5MG	A211487 003	Nov 07, 2018
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AA		325MG;10MG	A211487 004	Nov 07, 2018
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AA	AUROLIFE PHARMA LLC	300MG;5MG	A207709 001	Sep 13, 2018
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AA		300MG;7.5MG	A207709 002	Sep 13, 2018
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AA		300MG;10MG	A207709 003	Sep 13, 2018
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AA		325MG;5MG	A201013 001	Apr 11, 2012
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AA		325MG;7.5MG	A201013 002	Apr 11, 2012
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AA		325MG;10MG	A201013 003	Apr 11, 2012
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AA	! CHARTWELL	300MG;5MG	A040658 001	Jan 19, 2006
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AA	! CHARTWELL	300MG;7.5MG	A040658 002	Mar 24, 2006
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AA	! CHARTWELL	300MG;10MG	A040658 003	Jun 23, 2004
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PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A040432 001</u>	Jan 22, 2003
<u>AA</u>	EPIC PHARMA LLC	<u>325MG; 5MG</u>	<u>A203863 001</u>	Mar 30, 2018
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A203863 002</u>	Mar 30, 2018
<u>AA</u>		<u>325MG; 10MG</u>	<u>A203863 003</u>	Mar 30, 2018
<u>AA</u>	PRINSTON INC	<u>325MG; 5MG</u>	<u>A214928 001</u>	Dec 30, 2021
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A214928 002</u>	Dec 30, 2021
<u>AA</u>		<u>325MG; 10MG</u>	<u>A214928 003</u>	Dec 30, 2021
<u>AA</u>	RHODES PHARMS	<u>300MG; 5MG</u>	<u>A207808 001</u>	Mar 30, 2018
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A207808 002</u>	Mar 30, 2018
<u>AA</u>		<u>300MG; 10MG</u>	<u>A207808 003</u>	Mar 30, 2018
<u>AA</u>		<u>325MG; 5MG</u>	<u>A202991 001</u>	Apr 12, 2016
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A202991 002</u>	Apr 12, 2016
<u>AA</u>		<u>325MG; 10MG</u>	<u>A202991 003</u>	Apr 12, 2016
<u>AA</u>	SPECGX LLC	<u>300MG; 5MG</u>	<u>A206718 001</u>	Mar 31, 2017
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A206718 002</u>	Mar 31, 2017
<u>AA</u>		<u>300MG; 10MG</u>	<u>A206718 003</u>	Mar 31, 2017
<u>AA</u>	!	<u>325MG; 10MG</u>	<u>A040400 001</u>	Jul 26, 2000
<u>AA</u>	STRIDES PHARMA	<u>325MG; 5MG</u>	<u>A040655 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG; 5MG</u>	<u>A202935 002</u>	Jun 15, 2016
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A040656 001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A202935 003</u>	Jun 15, 2016
<u>AA</u>		<u>325MG; 10MG</u>	<u>A040355 001</u>	May 31, 2000
<u>AA</u>		<u>325MG; 10MG</u>	<u>A202935 004</u>	Jun 15, 2016
<u>AA</u>	TRIS PHARMA INC	<u>300MG; 5MG</u>	<u>A202214 004</u>	Mar 15, 2016
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A202214 005</u>	Mar 15, 2016
<u>AA</u>		<u>300MG; 10MG</u>	<u>A202214 006</u>	Mar 15, 2016
<u>AA</u>		<u>325MG; 5MG</u>	<u>A202214 001</u>	Mar 27, 2013
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A202214 002</u>	Mar 27, 2013
<u>AA</u>		<u>325MG; 10MG</u>	<u>A202214 003</u>	Mar 27, 2013
<u>AA</u>	WES PHARMA INC	<u>300MG; 5MG</u>	<u>A207509 001</u>	Oct 29, 2018
<u>AA</u>		<u>300MG; 7.5MG</u>	<u>A207509 002</u>	Oct 29, 2018
<u>AA</u>		<u>300MG; 10MG</u>	<u>A207509 003</u>	Oct 29, 2018
<u>AA</u>		<u>325MG; 5MG</u>	<u>A210211 001</u>	Oct 30, 2017
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A210211 002</u>	Oct 30, 2017
<u>AA</u>		<u>325MG; 10MG</u>	<u>A210211 003</u>	Oct 30, 2017
<u>AA</u>	!	ASCENT PHARMS INC 325MG; 2.5MG	A211487 001	Nov 07, 2018

ACETAMINOPHEN; IBUPROFEN SODIUM

SOLUTION; INTRAVENOUS

COMBOGESIC IV

+	!	HIKMA	1GM/100ML (10MG/ML); EQ 300MG BASE/100ML (EQ 3MG BASE/ML)	N215320 001	Oct 17, 2023
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ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>	!	ABHAI LLC	<u>325MG/5ML; 5MG/5ML</u>	<u>A211499 001</u>	Dec 31, 2018
<u>AA</u>		NOSTRUM LABS INC	<u>325MG/5ML; 5MG/5ML</u>	<u>A201448 001</u>	Aug 26, 2021
		MIKART	300MG/5ML; 10MG/5ML	A202142 001	Nov 27, 2018

TABLET; ORAL

OXYCET

<u>AA</u>		SPECGX LLC	<u>325MG; 5MG</u>	<u>A087463 001</u>	Dec 07, 1983
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OXYCODONE AND ACETAMINOPHEN

<u>AA</u>		ABHAI LLC	<u>325MG; 2.5MG</u>	<u>A210644 001</u>	Feb 09, 2018
<u>AA</u>			<u>325MG; 5MG</u>	<u>A210644 002</u>	Feb 09, 2018
<u>AA</u>			<u>325MG; 7.5MG</u>	<u>A210644 003</u>	Feb 09, 2018
<u>AA</u>			<u>325MG; 10MG</u>	<u>A210644 004</u>	Feb 09, 2018
<u>AA</u>	ACTAVIS ELIZABETH		<u>325MG; 2.5MG</u>	<u>A201447 001</u>	Apr 12, 2013
<u>AA</u>			<u>325MG; 5MG</u>	<u>A201447 002</u>	Apr 12, 2013
<u>AA</u>			<u>325MG; 7.5MG</u>	<u>A201447 003</u>	Apr 12, 2013
<u>AA</u>			<u>325MG; 10MG</u>	<u>A201447 004</u>	Apr 12, 2013
<u>AA</u>	ALVOGEN		<u>325MG; 5MG</u>	<u>A202677 003</u>	Mar 08, 2016
<u>AA</u>			<u>325MG; 7.5MG</u>	<u>A202677 001</u>	Jul 26, 2012
<u>AA</u>			<u>325MG; 10MG</u>	<u>A202677 002</u>	Jul 26, 2012
<u>AA</u>	AMNEAL PHARMS		<u>325MG; 5MG</u>	<u>A040777 001</u>	Nov 27, 2007
<u>AA</u>	AMNEAL PHARMS NY		<u>325MG; 7.5MG</u>	<u>A040778 002</u>	Jun 27, 2014
<u>AA</u>			<u>325MG; 10MG</u>	<u>A040778 001</u>	Nov 27, 2007
<u>AA</u>	ASCENT PHARMS INC		<u>325MG; 2.5MG</u>	<u>A207419 001</u>	Mar 22, 2017
<u>AA</u>			<u>325MG; 5MG</u>	<u>A207419 002</u>	Mar 22, 2017
<u>AA</u>			<u>325MG; 7.5MG</u>	<u>A207419 003</u>	Mar 22, 2017
<u>AA</u>			<u>325MG; 10MG</u>	<u>A207419 004</u>	Mar 22, 2017

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>	AUROLIFE PHARMA LLC	<u>325MG; 2.5MG</u>	<u>A201972 001</u>	Jul 15, 2013
<u>AA</u>		<u>325MG; 5MG</u>	<u>A201972 002</u>	Jul 15, 2013
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A201972 003</u>	Jul 15, 2013
<u>AA</u>		<u>325MG; 10MG</u>	<u>A201972 004</u>	Jul 15, 2013
<u>AA</u>	CHARTWELL	<u>325MG; 5MG</u>	<u>A207834 001</u>	Aug 15, 2019
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A207834 002</u>	Aug 15, 2019
<u>AA</u>		<u>325MG; 10MG</u>	<u>A207834 003</u>	Aug 15, 2019
<u>AA</u>	EPIC PHARMA LLC	<u>325MG; 5MG</u>	<u>A203864 001</u>	Jul 02, 2018
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A203864 002</u>	Jul 02, 2018
<u>AA</u>		<u>325MG; 10MG</u>	<u>A203864 003</u>	Jul 02, 2018
<u>AA</u>	NOVEL LABS INC	<u>325MG; 2.5MG</u>	<u>A204407 001</u>	Feb 24, 2017
<u>AA</u>		<u>325MG; 5MG</u>	<u>A204407 002</u>	Feb 24, 2017
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A204407 003</u>	Feb 24, 2017
<u>AA</u>		<u>325MG; 10MG</u>	<u>A204407 004</u>	Feb 24, 2017
<u>AA</u>	RHODES PHARMS	<u>325MG; 5MG</u>	<u>A201278 001</u>	Aug 28, 2014
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A201278 002</u>	Aug 28, 2014
<u>AA</u>		<u>325MG; 10MG</u>	<u>A201278 003</u>	Aug 28, 2014
<u>AA</u>	SPECGX LLC	<u>325MG; 7.5MG</u>	<u>A040545 001</u>	Jun 30, 2004
<u>AA</u>		<u>325MG; 10MG</u>	<u>A040545 002</u>	Jun 30, 2004
<u>AA</u>	WES PHARMA INC	<u>325MG; 5MG</u>	<u>A207510 001</u>	Mar 21, 2018
<u>AA</u>		<u>325MG; 7.5MG</u>	<u>A207510 002</u>	Mar 21, 2018
<u>AA</u>		<u>325MG; 10MG</u>	<u>A207510 003</u>	Mar 21, 2018

PERCOCET

<u>AA</u>	! VINTAGE PHARMS LLC	<u>325MG; 2.5MG</u>	<u>A040330 001</u>	Jun 25, 1999
<u>AA</u>	!	<u>325MG; 5MG</u>	<u>A040330 002</u>	Jun 25, 1999
<u>AA</u>	!	<u>325MG; 7.5MG</u>	<u>A040330 003</u>	Nov 23, 2001
<u>AA</u>	!	<u>325MG; 10MG</u>	<u>A040330 004</u>	Nov 23, 2001

OXYCODONE AND ACETAMINOPHEN

!	MIKART	300MG; 2.5MG	A040608 001	Dec 30, 2005
!		300MG; 5MG	A040608 002	Dec 30, 2005
!		300MG; 7.5MG	A040608 003	Dec 30, 2005
!		300MG; 10MG	A040608 004	Dec 30, 2005

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

<u>AB</u>	ALKEM LABS LTD	<u>325MG; 37.5MG</u>	<u>A202076 001</u>	Mar 30, 2012
<u>AB</u>	! AMNEAL PHARMS	<u>325MG; 37.5MG</u>	<u>A090485 001</u>	Dec 09, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>325MG; 37.5MG</u>	<u>A207152 001</u>	Mar 22, 2017
<u>AB</u>	MICRO LABS LTD INDIA	<u>325MG; 37.5MG</u>	<u>A201952 001</u>	Dec 14, 2012
<u>AB</u>	RISING	<u>325MG; 37.5MG</u>	<u>A077858 001</u>	Sep 26, 2008
<u>AB</u>	SUN PHARM INDS INC	<u>325MG; 37.5MG</u>	<u>A077184 001</u>	Dec 16, 2005
<u>AB</u>	ZYDUS PHARMS USA INC	<u>325MG; 37.5MG</u>	<u>A090460 001</u>	Sep 06, 2012

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

<u>AB</u>	ACCORD HLTHCARE	<u>500MG</u>	<u>A207659 001</u>	Oct 18, 2018
<u>AB</u>	ALEMBIC	<u>500MG</u>	<u>A210423 001</u>	Feb 19, 2019
<u>AB</u>	! CADILA	<u>500MG</u>	<u>A205301 001</u>	Jan 16, 2019
<u>AB</u>	HERITAGE PHARMA	<u>500MG</u>	<u>A040904 001</u>	Dec 10, 2008
<u>AB</u>	INDICUS PHARMA	<u>500MG</u>	<u>A090779 001</u>	Jul 14, 2011
<u>AB</u>	MICRO LABS LTD INDIA	<u>500MG</u>	<u>A207401 001</u>	Oct 01, 2020
<u>AB</u>	NOSTRUM LABS INC	<u>500MG</u>	<u>A204691 001</u>	Mar 29, 2016
<u>AB</u>	NOVAST LABS	<u>500MG</u>	<u>A203434 001</u>	Sep 30, 2016

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>125MG</u>	<u>A211151 001</u>	Sep 11, 2023
<u>AB</u>		<u>250MG</u>	<u>A211151 002</u>	Sep 11, 2023
<u>AB</u>	APPCO	<u>125MG</u>	<u>A211372 001</u>	Feb 22, 2021
<u>AB</u>		<u>250MG</u>	<u>A211372 002</u>	Feb 22, 2021
<u>AB</u>	CHARTWELL MOLECULAR	<u>250MG</u>	<u>A084840 001</u>	
<u>AB</u>	EYWA PHARMA	<u>125MG</u>	<u>A211556 001</u>	Oct 18, 2019
<u>AB</u>		<u>250MG</u>	<u>A211556 002</u>	Oct 18, 2019
<u>AB</u>	HERITAGE PHARMA	<u>125MG</u>	<u>A205530 001</u>	Oct 27, 2016
<u>AB</u>		<u>250MG</u>	<u>A205530 002</u>	Oct 27, 2016
<u>AB</u>	MANKIND PHARMA	<u>125MG</u>	<u>A214282 001</u>	Oct 07, 2020
<u>AB</u>		<u>250MG</u>	<u>A214282 002</u>	Oct 07, 2020

PRESCRIPTION DRUG PRODUCT LIST

ACETAZOLAMIDE

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>	NE RX PHARMA	<u>125MG</u>	<u>A217197 001</u>	May 09, 2023
<u>AB</u>		<u>250MG</u>	<u>A217197 002</u>	May 09, 2023
<u>AB</u>	NOVITIUM PHARMA	<u>125MG</u>	<u>A210588 001</u>	Oct 17, 2019
<u>AB</u>		<u>250MG</u>	<u>A210588 002</u>	Oct 17, 2019
<u>AB</u>	RUBICON	<u>125MG</u>	<u>A215101 001</u>	Aug 19, 2021
<u>AB</u>		<u>250MG</u>	<u>A215101 002</u>	Aug 19, 2021
<u>AB</u>	STRIDES PHARMA	<u>125MG</u>	<u>A209734 001</u>	Nov 20, 2017
<u>AB</u>		<u>250MG</u>	<u>A209734 002</u>	Nov 20, 2017
<u>AB</u>	TARO	<u>125MG</u>	<u>A040195 001</u>	May 28, 1997
<u>AB</u>	!	<u>250MG</u>	<u>A040195 002</u>	May 28, 1997
<u>AB</u>	ZYDUS	<u>125MG</u>	<u>A211069 001</u>	Apr 04, 2023
<u>AB</u>		<u>250MG</u>	<u>A211069 002</u>	Apr 04, 2023

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

<u>AP</u>	AVET LIFESCIENCES	<u>EQ 500MG BASE/VIAL</u>	<u>A202693 001</u>	Dec 19, 2014
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A040089 001</u>	Feb 28, 1995
<u>AP</u>	MYLAN ASI	<u>EQ 500MG BASE/VIAL</u>	<u>A200880 001</u>	May 09, 2012
<u>AP</u>	! XGEN PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A040784 001</u>	Dec 10, 2008
<u>AP</u>	ZYDUS PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A206533 001</u>	Apr 15, 2019

ACETIC ACID, GLACIAL

SOLUTION; IRRIGATION, URETHRAL

ACETIC ACID 0.25% IN PLASTIC CONTAINER

<u>AT</u>	B BRAUN	<u>250MG/100ML</u>	<u>N018161 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>250MG/100ML</u>	<u>N018523 001</u>	Feb 19, 1982
<u>AT</u>	+! ICU MEDICAL INC	<u>250MG/100ML</u>	<u>N017656 001</u>	

SOLUTION/DROPS; OTIC

ACETIC ACID

<u>AT</u>	RISING	<u>2%</u>	<u>A207280 001</u>	Mar 09, 2018
<u>AT</u>	SAPTALIS PHARMS	<u>2%</u>	<u>A040607 001</u>	Feb 24, 2005
<u>AT</u>	TARO	<u>2%</u>	<u>A088638 001</u>	Sep 06, 1984

VOSOL

<u>AT</u>	! HIKMA	<u>2%</u>	<u>N012179 001</u>	
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ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

HYDROCORTISONE AND ACETIC ACID

<u>AT</u>	COSETTE	<u>2%;1%</u>	<u>A040609 001</u>	Feb 06, 2006
<u>AT</u>	+! TARO	<u>2%;1%</u>	<u>A088759 001</u>	Mar 04, 1985
<u>AT</u>	+! SAPTALIS PHARMS	<u>2%;1%</u>	<u>N012770 001</u>	

ACETOHYDROXAMIC ACID

TABLET; ORAL

LITHOSTAT

+!	MISSION PHARMA	250MG	N018749 001	May 31, 1983
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ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

MIOCHOL-E

+!	BAUSCH AND LOMB	20MG/VIAL	N020213 001	Sep 22, 1993
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ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETADOTE

<u>AP</u>	+! CUMBERLAND PHARMS	<u>6GM/30ML (200MG/ML)</u>	<u>N021539 001</u>	Jan 23, 2004
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ACETYLCYSTEINE

<u>AP</u>	ASPEN	<u>6GM/30ML (200MG/ML)</u>	<u>A213693 001</u>	Feb 03, 2022
<u>AP</u>	EUGIA PHARMA	<u>6GM/30ML (200MG/ML)</u>	<u>A207358 001</u>	Feb 29, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>6GM/30ML (200MG/ML)</u>	<u>A200644 001</u>	Nov 07, 2012
<u>AP</u>	INDOCO	<u>6GM/30ML (200MG/ML)</u>	<u>A215620 001</u>	Feb 23, 2022
<u>AP</u>	RISING	<u>6GM/30ML (200MG/ML)</u>	<u>A203173 001</u>	Mar 24, 2015
<u>AP</u>	SAGENT PHARMS INC	<u>6GM/30ML (200MG/ML)</u>	<u>A091684 001</u>	Oct 31, 2017
<u>AP</u>	STERISCIENCE	<u>6GM/30ML (200MG/ML)</u>	<u>A217182 001</u>	Apr 19, 2023
<u>AP</u>	ZYDUS PHARMS	<u>6GM/30ML (200MG/ML)</u>	<u>A208166 001</u>	Jul 20, 2018

SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

<u>AN</u>	ALVOGEN	<u>10%</u>	<u>A204674 001</u>	Feb 11, 2014
<u>AN</u>		<u>20%</u>	<u>A203853 001</u>	Jun 21, 2012
<u>AN</u>	! AM REGENT	<u>10%</u>	<u>A072489 001</u>	Jul 28, 1995

PRESCRIPTION DRUG PRODUCT LIST

ACETYLCYSTEINE

SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

<u>AN</u>	!		<u>20%</u>	<u>A072547</u>	<u>001</u>	Jul 28, 1995
<u>AN</u>		EXELA PHARMA	<u>20%</u>	<u>A205643</u>	<u>001</u>	Nov 01, 2023
<u>AN</u>		HOSPIRA	<u>10%</u>	<u>A073664</u>	<u>001</u>	Aug 30, 1994
<u>AN</u>			<u>20%</u>	<u>A074037</u>	<u>001</u>	Aug 30, 1994

ACITRETIN

CAPSULE; ORAL

ACITRETIN

<u>AB</u>		BARR LABS INC	<u>10MG</u>	<u>A091455</u>	<u>001</u>	Apr 04, 2013
<u>AB</u>			<u>25MG</u>	<u>A091455</u>	<u>002</u>	Apr 04, 2013
<u>AB</u>		IMPAX LABS INC	<u>10MG</u>	<u>A202552</u>	<u>001</u>	Dec 23, 2015
<u>AB</u>			<u>17.5MG</u>	<u>A202552</u>	<u>002</u>	Dec 23, 2015
<u>AB</u>			<u>22.5MG</u>	<u>A202552</u>	<u>003</u>	Dec 23, 2015
<u>AB</u>			<u>25MG</u>	<u>A202552</u>	<u>004</u>	Dec 23, 2015
<u>AB</u>		MYLAN	<u>10MG</u>	<u>A202148</u>	<u>001</u>	Sep 10, 2015
<u>AB</u>			<u>25MG</u>	<u>A202148</u>	<u>002</u>	Sep 10, 2015
<u>AB</u>		SIGMAPHARM LABS LLC	<u>10MG</u>	<u>A204633</u>	<u>001</u>	May 22, 2015
<u>AB</u>			<u>17.5MG</u>	<u>A204633</u>	<u>002</u>	May 22, 2015
<u>AB</u>			<u>22.5MG</u>	<u>A204633</u>	<u>003</u>	May 22, 2015
<u>AB</u>	!		<u>25MG</u>	<u>A204633</u>	<u>004</u>	May 22, 2015
<u>AB</u>		TEVA PHARMS USA	<u>17.5MG</u>	<u>A202897</u>	<u>001</u>	Apr 04, 2013
<u>AB</u>			<u>22.5MG</u>	<u>A202897</u>	<u>002</u>	Apr 04, 2013

ACLIDINIUM BROMIDE

POWDER, METERED; INHALATION

TUDORZA PRESSAIR

+! COVIS

0.4MG/INH

N202450 001 Jul 23, 2012

ACLIDINIUM BROMIDE; FORMOTEROL FUMARATE

POWDER, METERED; INHALATION

DUAKLIR PRESSAIR

+! COVIS

0.4MG/INH; 0.012MG/INH

N210595 001 Mar 29, 2019

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

<u>AB</u>	!	APOTEX	<u>200MG</u>	<u>A075677</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>		CADILA	<u>200MG</u>	<u>A204313</u>	<u>001</u>	Mar 25, 2016
<u>AB</u>		CADILA PHARMS LTD	<u>200MG</u>	<u>A201445</u>	<u>001</u>	Mar 06, 2014
<u>AB</u>		CARLSBAD TECHNOLOGY	<u>200MG</u>	<u>A206261</u>	<u>001</u>	Aug 16, 2017
<u>AB</u>		HERITAGE PHARMS	<u>200MG</u>	<u>A074889</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>		INDOCO	<u>200MG</u>	<u>A075090</u>	<u>001</u>	Jan 26, 1999
<u>AB</u>		TEVA	<u>200MG</u>	<u>A074578</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>		YILING	<u>200MG</u>	<u>A212173</u>	<u>001</u>	Sep 14, 2020

CREAM; TOPICAL

ACYCLOVIR

<u>AB</u>		ALEMBIC	<u>5%</u>	<u>A212361</u>	<u>001</u>	Nov 21, 2023
<u>AB</u>		AMNEAL	<u>5%</u>	<u>A208766</u>	<u>001</u>	Nov 09, 2020
<u>AB</u>	!	PADAGIS ISRAEL	<u>5%</u>	<u>A208702</u>	<u>001</u>	Feb 04, 2019
<u>AB</u>		ZYDUS LIFESCIENCES	<u>5%</u>	<u>A206770</u>	<u>001</u>	Feb 28, 2023

ZOVIRAX

<u>AB</u>	+	BAUSCH	<u>5%</u>	<u>N021478</u>	<u>001</u>	Dec 30, 2002
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OINTMENT; TOPICAL

ACYCLOVIR

<u>AB</u>		ALEMBIC	<u>5%</u>	<u>A209000</u>	<u>001</u>	Apr 06, 2018
<u>AB</u>		AMNEAL PHARMS	<u>5%</u>	<u>A204605</u>	<u>001</u>	Jun 18, 2014
<u>AB</u>		APOTEX	<u>5%</u>	<u>A210774</u>	<u>001</u>	Sep 06, 2019
<u>AB</u>		CADILA	<u>5%</u>	<u>A205974</u>	<u>001</u>	Mar 15, 2019
<u>AB</u>		CHARTWELL RX	<u>5%</u>	<u>A212495</u>	<u>001</u>	Apr 07, 2020
<u>AB</u>		CIPLA	<u>5%</u>	<u>A211794</u>	<u>001</u>	Jan 18, 2019
<u>AB</u>		COSETTE	<u>5%</u>	<u>A205591</u>	<u>001</u>	Nov 13, 2017
<u>AB</u>		FOUGERA PHARMS INC	<u>5%</u>	<u>A206633</u>	<u>001</u>	May 11, 2016
<u>AB</u>		GLENMARK PHARMS SA	<u>5%</u>	<u>A205510</u>	<u>001</u>	Jul 31, 2017
<u>AB</u>		MACLEODS PHARMS LTD	<u>5%</u>	<u>A212444</u>	<u>001</u>	May 19, 2021
<u>AB</u>		MYLAN PHARMS INC	<u>5%</u>	<u>A202459</u>	<u>001</u>	Apr 03, 2013
<u>AB</u>		TARO	<u>5%</u>	<u>A205469</u>	<u>001</u>	Dec 21, 2016
<u>AB</u>		TORRENT	<u>5%</u>	<u>A209971</u>	<u>001</u>	Jan 11, 2019
<u>AB</u>		XIROMED	<u>5%</u>	<u>A201501</u>	<u>001</u>	Jan 29, 2020

ZOVIRAX

<u>AB</u>	+	BAUSCH	<u>5%</u>	<u>N018604</u>	<u>001</u>	Mar 29, 1982
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PRESCRIPTION DRUG PRODUCT LIST

ACYCLOVIR

SUSPENSION; ORAL

ACYCLOVIR

AB	ACTAVIS MID ATLANTIC	200MG/5ML	A074738 001	Apr 28, 1997
AB	AUROBINDO PHARMA	200MG/5ML	A216331 001	Mar 31, 2022
AB	CHARTWELL RX	200MG/5ML	A212718 001	Apr 23, 2020
AB	HETERO LABS LTD III	200MG/5ML	A215669 001	Jul 01, 2022
AB	HIKMA	200MG/5ML	A077026 001	Jun 07, 2005
AB	MSN	200MG/5ML	A217393 001	Mar 09, 2023
AB	! NOVITIUM PHARMA	200MG/5ML	A212252 001	Jul 10, 2020
AB	RUBICON	200MG/5ML	A215724 001	Aug 18, 2022

TABLET; BUCCAL

SITAVIG

+	! LNHC	50MG	N203791 001	Apr 12, 2013
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TABLET; ORAL

ACYCLOVIR

AB	APOTEX INC	400MG	A077309 001	Sep 29, 2005
AB		800MG	A077309 002	Sep 29, 2005
AB	CADILA PHARMS LTD	400MG	A202168 001	Nov 15, 2013
AB		800MG	A202168 002	Nov 15, 2013
AB	CARLSBAD	400MG	A075382 001	Apr 30, 1999
AB		800MG	A075382 002	Apr 30, 1999
AB	HERITAGE PHARMS	400MG	A074891 001	Oct 31, 1997
AB		800MG	A074891 002	Oct 31, 1997
AB	HETERO LABS LTD V	400MG	A203834 001	Oct 29, 2013
AB	!	800MG	A203834 002	Oct 29, 2013
AB	SQUARE PHARMS	400MG	A209366 001	Oct 07, 2019
AB		800MG	A209366 002	Oct 07, 2019
AB	STRIDES PHARMA	400MG	A074946 001	Nov 19, 1997
AB		800MG	A074946 002	Nov 19, 1997
AB	TEVA	400MG	A074556 002	Apr 22, 1997
AB		800MG	A074556 003	Apr 22, 1997
AB	YILING	400MG	A210401 001	Mar 07, 2018
AB		800MG	A210401 002	Mar 07, 2018
AB	ZYDUS PHARMS	400MG	A204314 001	Aug 19, 2014
AB		800MG	A204314 002	Aug 19, 2014

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

AP	EUGIA PHARMA	EQ 50MG BASE/ML	A203701 001	Oct 11, 2013
AP	! FRESENIUS KABI USA	EQ 50MG BASE/ML	A074930 001	May 13, 1998
AP	HAINAN POLY	EQ 50MG BASE/ML	A218111 001	Jan 08, 2024
AP	ZYDUS PHARMS	EQ 50MG BASE/ML	A206535 001	Aug 31, 2018

ACYCLOVIR; HYDROCORTISONE

CREAM; TOPICAL

XERESE

+	! BAUSCH	5%;1%	N022436 001	Jul 31, 2009
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ADAGRASIB

TABLET; ORAL

KRAZATI

+	! MIRATI THERAPS	200MG	N216340 001	Dec 12, 2022
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ADAPALENE

CREAM; TOPICAL

ADAPALENE

AB	FOUGERA PHARMS	0.1%	A090824 001	Jun 30, 2010
AB	! GALDERMA LABS LP	0.1%	N020748 001	May 26, 2000

GEL; TOPICAL

ADAPALENE

AB	ACTAVIS MID ATLANTIC	0.3%	A201000 001	Oct 27, 2014
AB	ALEMBIC	0.3%	A213508 001	Jun 18, 2020
AB	ENCUBE ETHICALS	0.3%	A200298 001	Jun 14, 2012
AB	TARO	0.3%	A208322 001	Jun 23, 2016

DIFFERIN

AB	! GALDERMA LABS LP	0.3%	N021753 001	Jun 19, 2007
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LOTION; TOPICAL

DIFFERIN

+	! GALDERMA LABS LP	0.1%	N022502 001	Mar 17, 2010
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PRESCRIPTION DRUG PRODUCT LIST

ADAPALENE

SOLUTION; TOPICAL

ADAPALENE

<u>AB</u>	CALL INC	<u>0.1%</u>	<u>A203981</u>	<u>001</u>	Sep 23, 2016
<u>AB</u>		<u>0.1%</u>	<u>A204593</u>	<u>001</u>	Jan 05, 2016

ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

<u>AB</u>	ACTAVIS LABS UT INC	<u>0.3%;2.5%</u>	<u>A209641</u>	<u>001</u>	Jun 22, 2022
<u>AB</u>	ALEMBIC	<u>0.3%;2.5%</u>	<u>A214185</u>	<u>001</u>	Aug 04, 2022
<u>AB</u>	ENCUBE	<u>0.1%;2.5%</u>	<u>A206164</u>	<u>001</u>	May 23, 2018
<u>AB</u>	GLENMARK PHARMS LTD	<u>0.1%;2.5%</u>	<u>A208108</u>	<u>001</u>	Nov 08, 2019
<u>AB</u>	PADAGIS ISRAEL	<u>0.1%;2.5%</u>	<u>A205033</u>	<u>001</u>	Jan 23, 2018
<u>AB</u>		<u>0.3%;2.5%</u>	<u>A212464</u>	<u>001</u>	May 31, 2022
<u>AB</u>	TARO	<u>0.1%;2.5%</u>	<u>A206959</u>	<u>001</u>	Jan 24, 2018
<u>AB</u>		<u>0.3%;2.5%</u>	<u>A209148</u>	<u>001</u>	Oct 17, 2018
<u>AB</u>	ZYDUS PHARMS	<u>0.3%;2.5%</u>	<u>A214553</u>	<u>001</u>	Jun 03, 2022

EPIDUO

<u>AB</u>	! GALDERMA LABS LP	<u>0.1%;2.5%</u>	<u>N022320</u>	<u>001</u>	Dec 08, 2008
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EPIDUO FORTE

<u>AB</u>	! GALDERMA LABS	<u>0.3%;2.5%</u>	<u>N207917</u>	<u>001</u>	Jul 15, 2015
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ADAPALENE; BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

CABTREO

!	BAUSCH	0.15%;3.1%;1.2%	N216632	001	Oct 20, 2023
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ADEFOVIR DIPIVOXIL

TABLET; ORAL

ADEFOVIR DIPIVOXIL

<u>AB</u>	APOTEX	<u>10MG</u>	<u>A205459</u>	<u>001</u>	Jul 06, 2018
<u>AB</u>	SIGMAPHARM LABS LLC	<u>10MG</u>	<u>A202051</u>	<u>001</u>	Aug 29, 2013

HEPSERA

<u>AB</u>	! GILEAD	<u>10MG</u>	<u>N021449</u>	<u>001</u>	Sep 20, 2002
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ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

<u>AP</u>	FRESENIUS KABI USA	<u>3MG/ML</u>	<u>A077133</u>	<u>001</u>	Apr 27, 2005
<u>AP</u>		<u>3MG/ML</u>	<u>A205568</u>	<u>001</u>	Apr 16, 2018
<u>AP</u>	GLAND PHARMA LTD	<u>3MG/ML</u>	<u>A077283</u>	<u>001</u>	Jun 14, 2007
<u>AP</u>		<u>3MG/ML</u>	<u>A206778</u>	<u>001</u>	Feb 16, 2018
<u>AP</u>	HIKMA	<u>3MG/ML</u>	<u>A076404</u>	<u>001</u>	Jun 16, 2004
<u>AP</u>		<u>3MG/ML</u>	<u>A076500</u>	<u>001</u>	Jun 16, 2004
<u>AP</u>	MYLAN LABS LTD	<u>3MG/ML</u>	<u>A078686</u>	<u>001</u>	May 13, 2009
<u>AP</u>	! RISING	<u>3MG/ML</u>	<u>A078076</u>	<u>001</u>	Oct 31, 2008

SOLUTION; INTRAVENOUS

ADENOSINE

<u>AP</u>	AVET LIFESCIENCES	<u>60MG/20ML (3MG/ML)</u>	<u>A202313</u>	<u>001</u>	Sep 15, 2014
<u>AP</u>		<u>90MG/30ML (3MG/ML)</u>	<u>A202313</u>	<u>002</u>	Sep 15, 2014
<u>AP</u>	EUGIA PHARMA	<u>60MG/20ML (3MG/ML)</u>	<u>A205331</u>	<u>001</u>	Nov 02, 2017
<u>AP</u>		<u>90MG/30ML (3MG/ML)</u>	<u>A205331</u>	<u>002</u>	Nov 02, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>60MG/20ML (3MG/ML)</u>	<u>A077897</u>	<u>001</u>	Nov 27, 2017
<u>AP</u>		<u>90MG/30ML (3MG/ML)</u>	<u>A077897</u>	<u>002</u>	Nov 27, 2017
<u>AP</u>	HOSPIRA	<u>60MG/20ML (3MG/ML)</u>	<u>A203883</u>	<u>001</u>	Mar 24, 2014
<u>AP</u>		<u>90MG/30ML (3MG/ML)</u>	<u>A203883</u>	<u>002</u>	Mar 24, 2014
<u>AP</u>	! MEITHEAL	<u>60MG/20ML (3MG/ML)</u>	<u>A077425</u>	<u>001</u>	Aug 29, 2013
<u>AP</u>	!	<u>90MG/30ML (3MG/ML)</u>	<u>A077425</u>	<u>002</u>	Aug 29, 2013
<u>AP</u>	MYLAN ASI	<u>60MG/20ML (3MG/ML)</u>	<u>A090212</u>	<u>001</u>	Mar 28, 2014
<u>AP</u>		<u>90MG/30ML (3MG/ML)</u>	<u>A090212</u>	<u>002</u>	Mar 28, 2014
<u>AP</u>	RISING	<u>60MG/20ML (3MG/ML)</u>	<u>A090450</u>	<u>001</u>	Oct 02, 2014
<u>AP</u>		<u>90MG/30ML (3MG/ML)</u>	<u>A090450</u>	<u>002</u>	Oct 02, 2014

AFAMELANOTIDE

IMPLANT; SUBCUTANEOUS

SCENESSE

!	CLIVUNEL INC	16MG	N210797	001	Oct 08, 2019
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PRESCRIPTION DRUG PRODUCT LIST

AFATINIB DIMALEATE

TABLET; ORAL

GILOTRIF

+	BOEHRINGER INGELHEIM	EQ 20MG BASE	N201292 001	Jul 12, 2013
+		EQ 30MG BASE	N201292 002	Jul 12, 2013
+	!	EQ 40MG BASE	N201292 003	Jul 12, 2013

AIR POLYMER-TYPE A

FOAM; INTRAUTERINE

EXEM FOAM KIT

+	!	GISKIT	10ML	N212279 001	Nov 07, 2019
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ALBENDAZOLE

TABLET; ORAL

ALBENDAZOLE

<u>AB</u>		ACTAVIS ELIZABETH	<u>200MG</u>	<u>A208094 001</u>	May 20, 2019
<u>AB</u>	!	DR REDDYS	<u>200MG</u>	<u>A211034 001</u>	Jan 26, 2021
<u>AB</u>		EDENBRIDGE PHARMS	<u>200MG</u>	<u>A211117 001</u>	May 14, 2019
<u>AB</u>		MSN	<u>200MG</u>	<u>A213435 001</u>	Jan 21, 2021
<u>AB</u>		STRIDES PHARMA	<u>200MG</u>	<u>A210011 001</u>	Dec 07, 2018
<u>AB</u>		ZYDUS PHARMS	<u>200MG</u>	<u>A208979 001</u>	Dec 14, 2018

ALBUMIN HUMAN

INJECTABLE; INJECTION

OPTISON

+	!	GE HEALTHCARE	10MG/ML	N020899 001	Dec 31, 1997
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ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATE

<u>AB1</u>		CIPLA	<u>EQ 0.09MG BASE/INH</u>	<u>A209959 001</u>	Apr 08, 2020	
<u>AB1</u>		SANDOZ	<u>EQ 0.09MG BASE/INH</u>	<u>A207085 001</u>	Jun 01, 2021	
		<u>PROVENTIL-HFA</u>				
<u>AB1</u>	+	!	KINDEVA	<u>EQ 0.09MG BASE/INH</u>	<u>N020503 001</u>	Aug 15, 1996
		<u>ALBUTEROL SULFATE</u>				
<u>AB2</u>		LUPIN	<u>EQ 0.09MG BASE/INH</u>	<u>A209954 001</u>	Aug 24, 2020	
		<u>PROAIR HFA</u>				
<u>AB2</u>	+	!	TEVA BRANDED PHARM VENTOLIN HFA	<u>EQ 0.09MG BASE/INH</u>	<u>N021457 001</u>	Oct 29, 2004
BX	+	!	GLAXOSMITHKLINE	EQ 0.09MG BASE/INH	N020983 001	Apr 19, 2001
		POWDER, METERED; INHALATION				
		PROAIR DIGIHALER				
	+		TEVA BRANDED PHARM	EQ 0.09MG BASE/INH	N205636 002	Dec 21, 2018
		PROAIR RESPICLICK				
	+	!	TEVA BRANDED PHARM	EQ 0.09MG BASE/INH	N205636 001	Mar 31, 2015
		SOLUTION; INHALATION				
		<u>ALBUTEROL SULFATE</u>				
<u>AN</u>		LUOXIN AUROVITAS	<u>EQ 0.083% BASE</u>	<u>A206224 001</u>	Oct 17, 2017	
<u>AN</u>		NEPHRON	<u>EQ 0.021% BASE</u>	<u>A076355 002</u>	Mar 31, 2010	
<u>AN</u>	!		<u>EQ 0.042% BASE</u>	<u>A076355 001</u>	Jun 28, 2004	
<u>AN</u>	!		<u>EQ 0.083% BASE</u>	<u>A074880 001</u>	Sep 17, 1997	
<u>AN</u>	!		<u>EQ 0.5% BASE</u>	<u>A075664 001</u>	Jun 26, 2001	
<u>AN</u>	!	RITEDOSE CORP	<u>EQ 0.021% BASE</u>	<u>A214531 001</u>	Dec 28, 2021	
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A214531 002</u>	Dec 28, 2021	
<u>AN</u>			<u>EQ 0.083% BASE</u>	<u>A077839 001</u>	Dec 16, 2008	
<u>AN</u>		SENTISS	<u>EQ 0.5% BASE</u>	<u>A074543 001</u>	Jan 15, 1998	
<u>AN</u>		SUN PHARM	<u>EQ 0.083% BASE</u>	<u>A207857 001</u>	Jul 21, 2017	
<u>AN</u>		WATSON LABS	<u>EQ 0.021% BASE</u>	<u>A077772 001</u>	Sep 25, 2007	
<u>AN</u>			<u>EQ 0.042% BASE</u>	<u>A077772 002</u>	Sep 25, 2007	

SYRUP; ORAL

ALBUTEROL SULFATE

<u>AA</u>		AMNEAL PHARMS	<u>EQ 2MG BASE/5ML</u>	<u>A079241 001</u>	May 12, 2010
<u>AA</u>		CHARTWELL MOLECULAR	<u>EQ 2MG BASE/5ML</u>	<u>A078105 001</u>	Dec 27, 2006
<u>AA</u>		CHARTWELL RX	<u>EQ 2MG BASE/5ML</u>	<u>A077788 001</u>	Jun 26, 2007
<u>AA</u>		COSETTE	<u>EQ 2MG BASE/5ML</u>	<u>A074454 001</u>	Sep 25, 1995
<u>AA</u>		HIKMA	<u>EQ 2MG BASE/5ML</u>	<u>A074749 001</u>	Jan 30, 1998
<u>AA</u>		QUAGEN	<u>EQ 2MG BASE/5ML</u>	<u>A212197 001</u>	Sep 06, 2019
<u>AA</u>	!	TEVA	<u>EQ 2MG BASE/5ML</u>	<u>A073419 001</u>	Mar 30, 1992

TABLET; ORAL

ALBUTEROL SULFATE

<u>AB</u>		AIZANT	<u>EQ 2MG BASE</u>	<u>A210948 001</u>	Mar 15, 2019
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A210948 002</u>	Mar 15, 2019
<u>AB</u>		AMNEAL PHARMS CO	<u>EQ 2MG BASE</u>	<u>A208804 001</u>	May 21, 2018
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A208804 002</u>	May 21, 2018

PRESCRIPTION DRUG PRODUCT LIST

ALBUTEROL SULFATE

TABLET; ORAL

ALBUTEROL SULFATE

<u>AB</u>		DASH PHARMS NATCO	<u>EQ 2MG BASE</u>	<u>A072894 002</u>	Jan 17, 1991
<u>AB</u>	!		<u>EQ 4MG BASE</u>	<u>A072894 001</u>	Jan 17, 1991
<u>AB</u>		RISING	<u>EQ 2MG BASE</u>	<u>A207046 001</u>	Jun 29, 2018
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A207046 002</u>	Jun 29, 2018
<u>AB</u>		SUN PHARM INDUSTRIES	<u>EQ 2MG BASE</u>	<u>A072637 002</u>	Dec 05, 1989
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A072637 001</u>	Dec 05, 1989
<u>AB</u>		VIRTUS PHARM	<u>EQ 2MG BASE</u>	<u>A211397 001</u>	Oct 26, 2018
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A211397 002</u>	Oct 26, 2018
<u>AB</u>		ZYDUS PHARMS	<u>EQ 2MG BASE</u>	<u>A208884 001</u>	Oct 22, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A208884 002</u>	Oct 22, 2020

ALBUTEROL SULFATE; BUDESONIDE

AEROSOL, METERED; INHALATION

AIRSUPRA

+	!	ASTRAZENECA	EQ 90MCG BASE/INH; 80MCG/INH	N214070 001	Jan 10, 2023
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ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

<u>AN</u>		CIPLA	<u>EQ 0.083% BASE; 0.017%</u>	<u>A077559 001</u>	Dec 31, 2007
<u>AN</u>	!	NEPHRON	<u>EQ 0.083% BASE; 0.017%</u>	<u>A076749 001</u>	Dec 31, 2007
<u>AN</u>		RITEDOSE CORP	<u>EQ 0.083% BASE; 0.017%</u>	<u>A202496 001</u>	Oct 01, 2012
<u>AN</u>		SUN PHARM	<u>EQ 0.083% BASE; 0.017%</u>	<u>A207875 001</u>	Aug 07, 2017
		SPRAY, METERED; INHALATION COMBIVENT RESPIMAT			
	+	!	BOEHRINGER INGELHEIM	EQ 0.1MG BASE/INH; 0.02MG/INH	N021747 001 Oct 07, 2011

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ALCLOMETASONE DIPROPIONATE

<u>AB</u>	!	FOUGERA PHARMS	<u>0.05%</u>	<u>A076973 001</u>	Jul 12, 2005
<u>AB</u>		GLENMARK GENERICS	<u>0.05%</u>	<u>A079061 001</u>	Jun 23, 2009
<u>AB</u>		TARO	<u>0.05%</u>	<u>A076587 001</u>	Sep 15, 2005

OINTMENT; TOPICAL

ALCLOMETASONE DIPROPIONATE

<u>AB</u>	!	FOUGERA PHARMS	<u>0.05%</u>	<u>A076884 001</u>	Jul 18, 2005
<u>AB</u>		GLENMARK GENERICS	<u>0.05%</u>	<u>A079227 001</u>	Jul 30, 2009
<u>AB</u>		TARO	<u>0.05%</u>	<u>A076730 001</u>	Jul 29, 2004

ALCOHOL

SOLUTION; INTRA-ARTERIAL

ABLYSINOL

+		BPI LABS	99% (1ML)	N207987 001	Jun 21, 2018
+	!		99% (5ML)	N207987 002	Jun 21, 2018

ALECTINIB HYDROCHLORIDE

CAPSULE; ORAL

ALECENSA

+	!	HOFFMANN-LA ROCHE	EQ 150MG BASE	N208434 001	Dec 11, 2015
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ALENDRONATE SODIUM

SOLUTION; ORAL

ALENDRONATE SODIUM

<u>AA</u>	!	HIKMA	<u>EQ 70MG BASE/75ML</u>	<u>A090520 001</u>	Feb 25, 2013
<u>AA</u>		NOVITIUM PHARMA	<u>EQ 70MG BASE/75ML</u>	<u>A214512 001</u>	May 11, 2023

TABLET; ORAL

ALENDRONATE SODIUM

<u>AB</u>		APOTEX	<u>EQ 5MG BASE</u>	<u>A077982 001</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A077982 002</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 35MG BASE</u>	<u>A077982 003</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 70MG BASE</u>	<u>A077982 004</u>	Aug 04, 2008
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A090124 001</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 35MG BASE</u>	<u>A090124 002</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 70MG BASE</u>	<u>A090124 003</u>	Aug 04, 2008
<u>AB</u>		CIPLA	<u>EQ 5MG BASE</u>	<u>A076768 001</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A076768 002</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 35MG BASE</u>	<u>A076768 003</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A076768 004</u>	Aug 04, 2008
<u>AB</u>			<u>EQ 70MG BASE</u>	<u>A076768 005</u>	Aug 04, 2008
<u>AB</u>		HANGZHOU BINJIANG	<u>EQ 5MG BASE</u>	<u>A090258 001</u>	Sep 24, 2009
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A090258 002</u>	Sep 24, 2009

PRESCRIPTION DRUG PRODUCT LIST

ALENDRONATE SODIUM

TABLET; ORAL

ALENDRONATE SODIUM

<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A090258 003</u>	Sep 24, 2009
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A090258 004</u>	Sep 24, 2009
<u>AB</u>	SUN PHARM	<u>EQ 5MG BASE</u>	<u>A090022 001</u>	Sep 10, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090022 002</u>	Sep 10, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A090022 003</u>	Sep 10, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A090022 004</u>	Sep 10, 2008
<u>AB</u>	WATSON LABS	<u>EQ 35MG BASE</u>	<u>A076984 001</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076984 002</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A076984 003</u>	Aug 04, 2008

FOSAMAX

<u>AB</u>	+! ORGANON	<u>EQ 70MG BASE</u>	<u>N020560 005</u>	Oct 20, 2000
TABLET, EFFERVESCENT; ORAL				
BINOSTO				
	+! RADIUS	EQ 70MG BASE	N202344 001	Mar 12, 2012

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET; ORAL

FOSAMAX PLUS D

	+ ORGANON LLC	EQ 70MG BASE; 2,800 IU	N021762 001	Apr 07, 2005
	+!	EQ 70MG BASE; 5,600 IU	N021762 002	Apr 26, 2007

ALFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ALFENTA

<u>AP</u>	+! RISING	<u>EQ 0.5MG BASE/ML</u>	<u>N019353 001</u>	Dec 29, 1986
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ALFENTANIL

<u>AP</u>	HOSPIRA	<u>EQ 0.5MG BASE/ML</u>	<u>A075221 001</u>	Oct 28, 1999
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ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALFUZOSIN HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>10MG</u>	<u>A079013 001</u>	Jul 18, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A079060 001</u>	Aug 30, 2012
<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A090284 001</u>	Jan 17, 2012
<u>AB</u>	SUN PHARM	<u>10MG</u>	<u>A079057 001</u>	Jul 18, 2011
<u>AB</u>	UNICHEM	<u>10MG</u>	<u>A203192 001</u>	Jan 28, 2016

UROXATRAL

<u>AB</u>	+! CONCORDIA	<u>10MG</u>	<u>N021287 001</u>	Jun 12, 2003
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ALISKIREN HEMIFUMARATE

TABLET; ORAL

ALISKIREN HEMIFUMARATE

<u>AB</u>	ANCHEN PHARMS	<u>EQ 150MG BASE</u>	<u>A206665 001</u>	Mar 22, 2019
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206665 002</u>	Mar 22, 2019

TEKTURNA

<u>AB</u>	+ NODEN PHARMA	<u>EQ 150MG BASE</u>	<u>N021985 001</u>	Mar 05, 2007
<u>AB</u>	+!	<u>EQ 300MG BASE</u>	<u>N021985 002</u>	Mar 05, 2007

ALITRETINOIN

GEL; TOPICAL

PANRETIN

	+! CONCORDIA	EQ 0.1% BASE	N020886 001	Feb 02, 1999
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ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

<u>AB</u>	ACCORD HLTHCARE	<u>100MG</u>	<u>A203154 001</u>	May 06, 2013
<u>AB</u>		<u>300MG</u>	<u>A203154 002</u>	May 06, 2013
<u>AB</u>	CHARTWELL	<u>100MG</u>	<u>A077353 001</u>	Sep 08, 2005
<u>AB</u>		<u>300MG</u>	<u>A077353 002</u>	Sep 08, 2005
<u>AB</u>	HARMAN FINOCHEM	<u>100MG</u>	<u>A214443 001</u>	Mar 07, 2022
<u>AB</u>		<u>300MG</u>	<u>A214443 002</u>	Mar 07, 2022
<u>AB</u>	HETERO LABS LTD V	<u>100MG</u>	<u>A217748 001</u>	Aug 03, 2023
<u>AB</u>		<u>300MG</u>	<u>A217748 002</u>	Aug 03, 2023
<u>AB</u>	INDOCO	<u>100MG</u>	<u>A204467 001</u>	Jul 28, 2016
<u>AB</u>		<u>300MG</u>	<u>A204467 002</u>	Jul 28, 2016
<u>AB</u>	IPCA LABS LTD	<u>100MG</u>	<u>A090637 001</u>	Mar 16, 2011
<u>AB</u>		<u>300MG</u>	<u>A090637 002</u>	Mar 16, 2011
<u>AB</u>	LUPIN LTD	<u>100MG</u>	<u>A211807 001</u>	Dec 14, 2023
<u>AB</u>		<u>300MG</u>	<u>A211807 002</u>	Dec 14, 2023
<u>AB</u>	MYLAN	<u>100MG</u>	<u>A018659 001</u>	Oct 24, 1986
<u>AB</u>		<u>300MG</u>	<u>A018659 002</u>	Oct 24, 1986

PRESCRIPTION DRUG PRODUCT LIST

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

<u>AB</u>	NORTHSTAR HLTHCARE	<u>100MG</u>	<u>A078253 001</u>	Sep 11, 2007
<u>AB</u>		<u>300MG</u>	<u>A078253 002</u>	Sep 11, 2007
<u>AB</u>	SUN PHARM INDUSTRIES	<u>100MG</u>	<u>A071450 002</u>	Jan 09, 1987
<u>AB</u>		<u>300MG</u>	<u>A071450 001</u>	Jan 09, 1987
<u>AB</u>	UNICHEM	<u>100MG</u>	<u>A211820 001</u>	Mar 12, 2019
<u>AB</u>	!	<u>300MG</u>	<u>A211820 002</u>	Mar 12, 2019
<u>AB</u>	VINTAGE PHARMS	<u>100MG</u>	<u>A075798 001</u>	Jun 27, 2003
<u>AB</u>		<u>300MG</u>	<u>A075798 002</u>	Jun 27, 2003
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N018832 002</u>	Sep 28, 1984
<u>AB</u>		<u>300MG</u>	<u>N018877 001</u>	Sep 28, 1984
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A210117 001</u>	Oct 12, 2017
<u>AB</u>		<u>300MG</u>	<u>A210117 002</u>	Oct 12, 2017

LOPURIN

<u>AB</u>	DR REDDYS LA	<u>100MG</u>	<u>A071586 001</u>	Apr 02, 1987
<u>AB</u>		<u>300MG</u>	<u>A071587 001</u>	Apr 02, 1987

ZYLOPRIM

<u>AB</u>	+ CASPER PHARMA LLC	<u>100MG</u>	<u>N016084 001</u>	
<u>AB</u>	+	<u>300MG</u>	<u>N016084 002</u>	
	+	200MG	N016084 003	Aug 04, 2022

ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALLOPURINOL SODIUM

<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A212363 001</u>	Jan 26, 2022
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A076870 001</u>	Aug 26, 2004

ALOPRIM

<u>AP</u>	+! MYLAN	<u>EQ 500MG BASE/VIAL</u>	<u>N020298 001</u>	May 17, 1996
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ALMOTRIPTAN MALATE

TABLET; ORAL

ALMOTRIPTAN MALATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 6.25MG BASE</u>	<u>A205523 001</u>	Mar 03, 2016
<u>AB</u>	!	<u>EQ 12.5MG BASE</u>	<u>A205523 002</u>	Mar 03, 2016
<u>AB</u>	MYLAN	<u>EQ 6.25MG BASE</u>	<u>A205171 001</u>	Nov 09, 2015
<u>AB</u>		<u>EQ 12.5MG BASE</u>	<u>A205171 002</u>	Nov 09, 2015
<u>AB</u>	TEVA PHARMS USA	<u>EQ 6.25MG BASE</u>	<u>A078027 001</u>	Jul 07, 2015
<u>AB</u>		<u>EQ 12.5MG BASE</u>	<u>A078027 002</u>	Jul 07, 2015

ALOGLIPTIN BENZOATE

TABLET; ORAL

NESINA

	+ TAKEDA PHARMS USA	EQ 6.25MG BASE	N022271 001	Jan 25, 2013
	+	EQ 12.5MG BASE	N022271 002	Jan 25, 2013
	+!	EQ 25MG BASE	N022271 003	Jan 25, 2013

ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

KAZANO

	+ TAKEDA PHARMS USA	EQ 12.5MG BASE;500MG	N203414 001	Jan 25, 2013
	+!	EQ 12.5MG BASE;1GM	N203414 002	Jan 25, 2013

ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

OSEN1

	+ TAKEDA PHARMS USA	EQ 12.5MG BASE;EQ 30MG BASE	N022426 005	Jan 25, 2013
	+	EQ 25MG BASE;EQ 15MG BASE	N022426 001	Jan 25, 2013
	+	EQ 25MG BASE;EQ 30MG BASE	N022426 002	Jan 25, 2013
	+!	EQ 25MG BASE;EQ 45MG BASE	N022426 003	Jan 25, 2013

ALOSETRON HYDROCHLORIDE

TABLET; ORAL

ALOSETRON HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>EQ 0.5MG BASE</u>	<u>A206647 001</u>	Dec 22, 2016
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A206647 002</u>	Dec 22, 2016
<u>AB</u>	MANKIND PHARMA	<u>EQ 0.5MG BASE</u>	<u>A213614 001</u>	Sep 09, 2020
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A213614 002</u>	Sep 09, 2020
<u>AB</u>	RISING	<u>EQ 0.5MG BASE</u>	<u>A209180 001</u>	Jan 14, 2019
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A209180 002</u>	Jan 14, 2019

LOTRONEX

<u>AB</u>	+ SEBELA IRELAND LTD	<u>EQ 0.5MG BASE</u>	<u>N021107 002</u>	Dec 23, 2003
<u>AB</u>	+!	<u>EQ 1MG BASE</u>	<u>N021107 001</u>	Feb 09, 2000

PRESCRIPTION DRUG PRODUCT LIST

ALPELISIB

TABLET; ORAL

PIQRAY

+	NOVARTIS	50MG	N212526	001	May 24, 2019
+		150MG	N212526	002	May 24, 2019
+	!	200MG	N212526	003	May 24, 2019

VIJOICE

+	NOVARTIS	50MG	N215039	001	Apr 05, 2022
+		125MG	N215039	002	Apr 05, 2022
+	!	200MG	N215039	003	Apr 05, 2022

ALPHA-TOCOPHEROL ACETATE; ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN K

SOLUTION; INTRAVENOUS

INFUVITE ADULT

+	SANDOZ CANADA INC	2 IU/ML; 40MG/ML; 12MCG/ML; 40 IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1.2MG/ML; 0.72MG/ML; 1.2MG/ML; 660 IU/ML; 0.03MG/ML	N021163	001	May 18, 2000
+	!	2 IU/ML; 40MG/ML; 12MCG/ML; 40 IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1.2MG/ML; 0.72MG/ML; 1.2MG/ML; 660 IU/ML; 30MCG/ML	N021163	002	Jun 16, 2003

ALPRAZOLAM

CONCENTRATE; ORAL

ALPRAZOLAM

!	HIKMA	1MG/ML	A074312	001	Oct 31, 1993
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TABLET; ORAL

ALPRAZOLAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.25MG</u>	<u>A074342</u>	<u>001</u>	Oct 31, 1993
<u>AB</u>		<u>0.5MG</u>	<u>A074342</u>	<u>002</u>	Oct 31, 1993
<u>AB</u>		<u>1MG</u>	<u>A074342</u>	<u>003</u>	Oct 31, 1993
<u>AB</u>		<u>2MG</u>	<u>A074342</u>	<u>004</u>	Oct 31, 1993
<u>AB</u>	APOTEX INC	<u>0.25MG</u>	<u>A077741</u>	<u>001</u>	Jan 19, 2007
<u>AB</u>		<u>0.5MG</u>	<u>A077741</u>	<u>002</u>	Jan 19, 2007
<u>AB</u>		<u>1MG</u>	<u>A077741</u>	<u>003</u>	Jan 19, 2007
<u>AB</u>		<u>2MG</u>	<u>A077741</u>	<u>004</u>	Jan 19, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>0.25MG</u>	<u>A203346</u>	<u>001</u>	Jul 31, 2015
<u>AB</u>		<u>0.5MG</u>	<u>A203346</u>	<u>002</u>	Jul 31, 2015
<u>AB</u>		<u>1MG</u>	<u>A203346</u>	<u>003</u>	Jul 31, 2015
<u>AB</u>		<u>2MG</u>	<u>A203346</u>	<u>004</u>	Jul 31, 2015
<u>AB</u>	BRECKENRIDGE	<u>0.25MG</u>	<u>A207507</u>	<u>001</u>	Jul 09, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A207507</u>	<u>002</u>	Jul 09, 2018
<u>AB</u>		<u>1MG</u>	<u>A207507</u>	<u>003</u>	Jul 09, 2018
<u>AB</u>		<u>2MG</u>	<u>A207507</u>	<u>004</u>	Jul 09, 2018
<u>AB</u>	NATCO	<u>0.25MG</u>	<u>A200739</u>	<u>001</u>	Apr 15, 2015
<u>AB</u>		<u>0.5MG</u>	<u>A200739</u>	<u>002</u>	Apr 15, 2015
<u>AB</u>		<u>1MG</u>	<u>A200739</u>	<u>003</u>	Apr 15, 2015
<u>AB</u>		<u>2MG</u>	<u>A200739</u>	<u>004</u>	Apr 15, 2015
<u>AB</u>	NOVITIUM PHARMA	<u>0.25MG</u>	<u>A074174</u>	<u>001</u>	Oct 19, 1993
<u>AB</u>		<u>0.5MG</u>	<u>A074174</u>	<u>002</u>	Oct 19, 1993
<u>AB</u>		<u>1MG</u>	<u>A074174</u>	<u>003</u>	Oct 19, 1993
<u>AB</u>		<u>2MG</u>	<u>A074174</u>	<u>004</u>	Oct 19, 1993
<u>AB</u>	SANDOZ	<u>0.25MG</u>	<u>A074112</u>	<u>001</u>	Dec 29, 1995
<u>AB</u>		<u>0.5MG</u>	<u>A074112</u>	<u>002</u>	Dec 29, 1995
<u>AB</u>		<u>1MG</u>	<u>A074112</u>	<u>003</u>	Dec 29, 1995
<u>AB</u>		<u>2MG</u>	<u>A074909</u>	<u>001</u>	Mar 25, 1998
<u>AB</u>	STRIDES PHARMA	<u>0.25MG</u>	<u>A090248</u>	<u>001</u>	Sep 17, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090248</u>	<u>002</u>	Sep 17, 2010
<u>AB</u>		<u>1MG</u>	<u>A090248</u>	<u>003</u>	Sep 17, 2010
<u>AB</u>		<u>2MG</u>	<u>A090248</u>	<u>004</u>	Sep 17, 2010
<u>AB</u>	SUN PHARM	<u>0.25MG</u>	<u>A090082</u>	<u>001</u>	Jun 17, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090082</u>	<u>002</u>	Jun 17, 2010
<u>AB</u>		<u>1MG</u>	<u>A090082</u>	<u>003</u>	Jun 17, 2010
<u>AB</u>		<u>2MG</u>	<u>A090082</u>	<u>004</u>	Jun 17, 2010
<u>XANAX</u>					
<u>AB</u>	+	UPJOHN	<u>0.25MG</u>	<u>N018276</u>	<u>001</u>
<u>AB</u>	+		<u>0.5MG</u>	<u>N018276</u>	<u>002</u>
<u>AB</u>	+	!	<u>1MG</u>	<u>N018276</u>	<u>003</u>
<u>AB</u>	+		<u>2MG</u>	<u>N018276</u>	<u>004</u> Nov 27, 1985

PRESCRIPTION DRUG PRODUCT LIST

ALPRAZOLAM

TABLET, EXTENDED RELEASE;ORAL

ALPRAZOLAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.5MG</u>	<u>A078056 001</u>	Feb 13, 2007
<u>AB</u>		<u>1MG</u>	<u>A078056 002</u>	Feb 13, 2007
<u>AB</u>		<u>2MG</u>	<u>A078056 003</u>	Feb 13, 2007
<u>AB</u>		<u>3MG</u>	<u>A078056 004</u>	Feb 13, 2007
<u>AB</u>	AMNEAL PHARMS NY	<u>0.5MG</u>	<u>A078387 001</u>	May 30, 2008
<u>AB</u>		<u>1MG</u>	<u>A078387 002</u>	May 30, 2008
<u>AB</u>		<u>2MG</u>	<u>A078387 003</u>	May 30, 2008
<u>AB</u>		<u>3MG</u>	<u>A078387 004</u>	May 30, 2008
<u>AB</u>	ANCHEN PHARMS	<u>0.5MG</u>	<u>A078469 001</u>	Sep 29, 2011
<u>AB</u>		<u>1MG</u>	<u>A078469 002</u>	Sep 29, 2011
<u>AB</u>		<u>2MG</u>	<u>A078469 003</u>	Sep 29, 2011
<u>AB</u>		<u>3MG</u>	<u>A078469 004</u>	Sep 29, 2011
<u>AB</u>	APOTEX INC	<u>0.5MG</u>	<u>A078449 001</u>	Nov 12, 2008
<u>AB</u>		<u>1MG</u>	<u>A078449 004</u>	Dec 23, 2015
<u>AB</u>		<u>2MG</u>	<u>A078449 002</u>	Nov 12, 2008
<u>AB</u>		<u>3MG</u>	<u>A078449 003</u>	Nov 12, 2008
<u>AB</u>	AUROBINDO PHARMA	<u>0.5MG</u>	<u>A090871 001</u>	Jun 07, 2011
<u>AB</u>		<u>1MG</u>	<u>A090871 002</u>	Jun 07, 2011
<u>AB</u>		<u>2MG</u>	<u>A090871 003</u>	Jun 07, 2011
<u>AB</u>		<u>3MG</u>	<u>A090871 004</u>	Jun 07, 2011

XANAX XR

<u>AB</u>	+ UPJOHN	<u>0.5MG</u>	<u>N021434 001</u>	Jan 17, 2003
<u>AB</u>	+	<u>1MG</u>	<u>N021434 002</u>	Jan 17, 2003
<u>AB</u>	+	<u>2MG</u>	<u>N021434 003</u>	Jan 17, 2003
<u>AB</u>	+	<u>3MG</u>	<u>N021434 004</u>	Jan 17, 2003

TABLET, ORALLY DISINTEGRATING;ORAL

ALPRAZOLAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.25MG</u>	<u>A078561 001</u>	Mar 16, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078561 002</u>	Mar 16, 2010
<u>AB</u>		<u>1MG</u>	<u>A078561 003</u>	Mar 16, 2010
<u>AB</u>		<u>2MG</u>	<u>A078561 004</u>	Mar 16, 2010
<u>AB</u>	PAR PHARM	<u>0.25MG</u>	<u>A078088 001</u>	Jan 09, 2009
<u>AB</u>		<u>0.5MG</u>	<u>A078088 002</u>	Jan 09, 2009
<u>AB</u>	!	<u>1MG</u>	<u>A078088 003</u>	Jan 09, 2009
<u>AB</u>		<u>2MG</u>	<u>A078088 004</u>	Jan 09, 2009

ALPROSTADIL

INJECTABLE; INJECTION

ALPROSTADIL

<u>AP</u>	HIKMA	<u>0.5MG/ML</u>	<u>A074815 001</u>	Jan 20, 1998
<u>AP</u>	MEITHEAL	<u>0.5MG/ML</u>	<u>A075196 001</u>	Apr 30, 1999

CAVERJECT

<u>AP</u>	+ PFIZER	<u>0.01MG/VIAL</u>	<u>N020379 001</u>	Jul 06, 1995
<u>AP</u>	+	<u>0.02MG/VIAL</u>	<u>N020379 002</u>	Jul 06, 1995
<u>AP</u>	+	<u>0.04MG/VIAL</u>	<u>N020379 004</u>	May 19, 1997

EDEX

<u>AP</u>	+ AUXILIUM PHARMS LLC	<u>0.01MG/VIAL</u>	<u>N020649 002</u>	Jun 12, 1997
<u>AP</u>	+	<u>0.02MG/VIAL</u>	<u>N020649 003</u>	Jun 12, 1997
<u>AP</u>	+	<u>0.04MG/VIAL</u>	<u>N020649 004</u>	Jun 12, 1997

PROSTIN VR PEDIATRIC

<u>AP</u>	+	PFIZER	<u>0.5MG/ML</u>	<u>N018484 001</u>
	CAVERJECT IMPULSE			
	PFIZER	0.01MG/VIAL	N021212 001	Jun 11, 2002
		0.02MG/VIAL	N021212 002	Jun 11, 2002
	EDEX			
	+	AUXILIUM PHARMS LLC	0.01MG/VIAL	N020649 005
	+		0.02MG/VIAL	N020649 006
	+		0.04MG/VIAL	N020649 007

SUPPOSITORY; URETHRAL

MUSE

	+	MYLAN SPECIALITY LP	0.25MG	N020700 002	Nov 19, 1996
	+		0.5MG	N020700 003	Nov 19, 1996
	+		1MG	N020700 004	Nov 19, 1996

PRESCRIPTION DRUG PRODUCT LIST

ALVIMOPAN

CAPSULE; ORAL

ALVIMOPAN

<u>AB</u>	HIKMA	<u>12MG</u>	<u>A217753</u>	<u>001</u>	Aug 31, 2023
<u>AB</u>	PAR PHARM	<u>12MG</u>	<u>A216843</u>	<u>001</u>	Jan 24, 2023
<u>AB</u>	! WATSON LABS TEVA	<u>12MG</u>	<u>A208295</u>	<u>001</u>	Dec 19, 2019

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>100MG</u>	<u>A208966</u>	<u>001</u>	Jun 21, 2017
<u>AB</u>	! BIONPHARMA	<u>100MG</u>	<u>A078720</u>	<u>001</u>	May 29, 2008
<u>AB</u>	HERITAGE PHARMA	<u>100MG</u>	<u>A209171</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	HUMANWELL PURACAP	<u>100MG</u>	<u>A214580</u>	<u>001</u>	Dec 20, 2022
<u>AB</u>	RISING	<u>100MG</u>	<u>A210129</u>	<u>001</u>	Mar 02, 2020
<u>AB</u>	RUBICON	<u>100MG</u>	<u>A212044</u>	<u>001</u>	May 21, 2020
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A071293</u>	<u>001</u>	Feb 18, 1987
<u>AB</u>	STRIDES PHARMA	<u>100MG</u>	<u>A209047</u>	<u>001</u>	Jun 07, 2017
<u>AB</u>		<u>100MG</u>	<u>A211354</u>	<u>001</u>	Feb 18, 2022
<u>AB</u>	UPSHER SMITH LABS	<u>100MG</u>	<u>A070589</u>	<u>001</u>	Aug 05, 1986
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A208278</u>	<u>001</u>	May 31, 2016

CAPSULE, EXTENDED RELEASE; ORAL

GOCOVRI

+	ADAMAS OPERATIONS	EQ 68.5MG BASE	N208944	001	Aug 24, 2017
+	!	EQ 137MG BASE	N208944	002	Aug 24, 2017

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

<u>AA</u>	! AUROBINDO PHARMA USA	<u>50MG/5ML</u>	<u>A074170</u>	<u>001</u>	Oct 28, 1994
<u>AA</u>	! CHARTWELL RX	<u>50MG/5ML</u>	<u>A074028</u>	<u>001</u>	Jun 28, 1993
<u>AA</u>	! CMP PHARMA INC	<u>50MG/5ML</u>	<u>A075819</u>	<u>001</u>	Sep 11, 2002
<u>AA</u>	ELYSIUM	<u>50MG/5ML</u>	<u>A214178</u>	<u>001</u>	Aug 20, 2021
<u>AA</u>	! PHARM ASSOC	<u>50MG/5ML</u>	<u>A074509</u>	<u>001</u>	Jul 17, 1995

TABLET; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>100MG</u>	<u>A214284</u>	<u>001</u>	Oct 15, 2020
<u>AB</u>	ATHEM	<u>100MG</u>	<u>A210215</u>	<u>001</u>	Mar 10, 2020
<u>AB</u>	STRIDES PHARMA	<u>100MG</u>	<u>A209035</u>	<u>001</u>	Jun 09, 2017
<u>AB</u>	! UPSHER SMITH LABS	<u>100MG</u>	<u>A076186</u>	<u>001</u>	Dec 16, 2002
<u>AB</u>	WATSON LABS INC	<u>100MG</u>	<u>A208096</u>	<u>001</u>	Dec 15, 2016

TABLET, EXTENDED RELEASE; ORAL

OSMOLEX ER

+	ADAMAS OPERATIONS	EQ 129MG BASE	N209410	001	Feb 16, 2018
+		EQ 193MG BASE	N209410	002	Feb 16, 2018

AMBRISENTAN

TABLET; ORAL

AMBRISENTAN

<u>AB</u>	APOTEX	<u>5MG</u>	<u>A210701</u>	<u>001</u>	May 19, 2022
<u>AB</u>		<u>10MG</u>	<u>A210701</u>	<u>002</u>	May 19, 2022
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A216531</u>	<u>001</u>	Jul 21, 2022
<u>AB</u>		<u>10MG</u>	<u>A216531</u>	<u>002</u>	Jul 21, 2022
<u>AB</u>	CIPLA	<u>5MG</u>	<u>A210715</u>	<u>001</u>	Apr 26, 2019
<u>AB</u>		<u>10MG</u>	<u>A210715</u>	<u>002</u>	Apr 26, 2019
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A208441</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>		<u>10MG</u>	<u>A208441</u>	<u>002</u>	Mar 28, 2019
<u>AB</u>	PAR PHARM INC	<u>5MG</u>	<u>A209509</u>	<u>001</u>	Apr 10, 2019
<u>AB</u>		<u>10MG</u>	<u>A209509</u>	<u>002</u>	Apr 10, 2019
<u>AB</u>	SIGMAPHARM LABS LLC	<u>5MG</u>	<u>A208354</u>	<u>001</u>	Apr 10, 2019
<u>AB</u>		<u>10MG</u>	<u>A208354</u>	<u>002</u>	Apr 10, 2019
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A210784</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>		<u>10MG</u>	<u>A210784</u>	<u>002</u>	Mar 28, 2019
<u>AB</u>	WATSON LABS INC	<u>5MG</u>	<u>A208252</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>		<u>10MG</u>	<u>A208252</u>	<u>002</u>	Mar 28, 2019
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A210058</u>	<u>001</u>	Mar 28, 2019
<u>AB</u>		<u>10MG</u>	<u>A210058</u>	<u>002</u>	Mar 28, 2019

LETAIRIS

<u>AB</u>	+	GILEAD	<u>5MG</u>	<u>N022081</u>	<u>001</u>	Jun 15, 2007
<u>AB</u>	+	!	<u>10MG</u>	<u>N022081</u>	<u>002</u>	Jun 15, 2007

PRESCRIPTION DRUG PRODUCT LIST

AMCINONIDE

CREAM; TOPICAL

AMCINONIDE

AB	!	GENUS LIFESCIENCES	0.1%	A076065	001	May 15, 2003
AB		TARO PHARM INDS	0.1%	A076229	001	May 31, 2002

LOTION; TOPICAL

AMCINONIDE

!	ANDA REPOSITORY	0.1%	A076329	001	Nov 06, 2002
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OINTMENT; TOPICAL

AMCINONIDE

AB	!	ANDA REPOSITORY	0.1%	A076096	001	Nov 19, 2002
AB		TARO PHARM INDS	0.1%	A076367	001	Mar 19, 2003

AMIFAMPRIDINE PHOSPHATE

TABLET; ORAL

FIRDAPSE

+	!	CATALYST PHARMS	EQ 10MG BASE	N208078	001	Nov 28, 2018
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AMIFOSTINE

INJECTABLE; INJECTION

ETHYOL

+	!	CLINIGEN	500MG/VIAL	N020221	001	Dec 08, 1995
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AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

AP	!	AVET LIFESCIENCES	EQ 250MG BASE/ML	A204040	001	Dec 12, 2013
AP		FRESENIUS KABI USA	EQ 250MG BASE/ML	A205604	001	Dec 09, 2015
AP		HIKMA	EQ 250MG BASE/ML	A063315	001	Apr 11, 1994
AP		MEITHEAL	EQ 250MG BASE/ML	A064045	002	Sep 28, 1993
AP		SAGENT PHARMS INC	EQ 250MG BASE/ML	A203323	001	May 12, 2016
!		HIKMA	EQ 50MG BASE/ML	A063313	001	Apr 11, 1994

SUSPENSION, LIPOSOMAL; INHALATION

ARIKAYCE KIT

+	!	INSMED INC	EQ 590MG BASE/8.4ML	N207356	001	Sep 28, 2018
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AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

AB	!	PAR PHARM	5MG	A070346	001	Jan 22, 1986
AB		SIGMAPHARM LABS LLC	5MG	A079133	001	Jan 30, 2009
AB	+	PADAGIS US	5MG	N018200	001	

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB		BARR	EQ 5MG ANHYDROUS; 50MG	A071111	001	May 10, 1988
AB	!	RISING	EQ 5MG ANHYDROUS; 50MG	A073209	001	Oct 31, 1991

AMINO ACIDS

INJECTABLE; INJECTION

AMINO ACIDS

		B BRAUN	15% (150GM/1000ML)	A091112	001	Apr 13, 2012	
			15% (300GM/2000ML)	A091112	002	Apr 13, 2012	
		AMINOSYN II 10% IN PLASTIC CONTAINER					
		ICU MEDICAL INC	10% (10GM/100ML)	N020015	001	Dec 19, 1991	
		AMINOSYN II 15% IN PLASTIC CONTAINER					
		ICU MEDICAL INC	15% (15GM/100ML)	N020041	001	Dec 19, 1991	
		AMINOSYN-PF 10%					
		ICU MEDICAL INC	10% (10GM/100ML)	N019492	002	Oct 17, 1986	
		AMINOSYN-PF 7%					
		ICU MEDICAL INC	7% (7GM/100ML)	N019398	001	Sep 06, 1985	
		CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER					
		BAXTER HLTHCARE	15% (15GM/100ML)	A020512	001	Aug 30, 1996	
		PREMASOL 10% IN PLASTIC CONTAINER					
		BAXTER HLTHCARE	10% (10GM/100ML)	A075880	002	Jun 19, 2003	
		PREMASOL 6% IN PLASTIC CONTAINER					
		BAXTER HLTHCARE	6% (6GM/100ML)	A075880	001	Jun 19, 2003	
		PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER					
	+	!	BAXTER HLTHCARE	20% (20GM/100ML)	N020849	001	Aug 26, 1998
		TRAVASOL 10% IN PLASTIC CONTAINER					
		BAXTER HLTHCARE	10% (10GM/100ML)	N018931	003	Aug 23, 1984	
		TRAVASOL 5.5% IN PLASTIC CONTAINER					
		BAXTER HLTHCARE	5.5% (5.5GM/100ML)	N018931	001	Aug 23, 1984	

PRESCRIPTION DRUG PRODUCT LISTAMINO ACIDS

INJECTABLE; INJECTION

TRAVASOL 8.5% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	8.5% (8.5GM/100ML)	N018931 002	Aug 23, 1984
TROPHAMINE				
+	B BRAUN	6% (6GM/100ML)	N019018 001	Jul 20, 1984
TROPHAMINE 10%				
+	B BRAUN	10% (10GM/100ML)	N019018 003	Sep 07, 1988

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	2.75%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML	N020678 002	Mar 26, 1997
CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	2.75%; 33MG/100ML; 25GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML	N020678 005	Mar 26, 1997
CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	2.75%; 33MG/100ML; 5GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML	N020678 001	Mar 26, 1997
CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	4.25%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020678 009	Mar 26, 1997
CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	4.25%; 33MG/100ML; 20GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020678 011	Mar 26, 1997
CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	4.25%; 33MG/100ML; 25GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020678 012	Mar 26, 1997
CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	4.25%; 33MG/100ML; 5GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	N020678 008	Mar 26, 1997
CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	5%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML	N020678 016	Mar 26, 1997
CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	5%; 33MG/100ML; 15GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML	N020678 017	Mar 26, 1997
CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	5%; 33MG/100ML; 20GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML	N020678 018	Mar 26, 1997
CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	5%; 33MG/100ML; 25GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML	N020678 019	Mar 26, 1997
CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	5%; 33MG/100ML; 35GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML	N020678 021	Mar 26, 1997

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM GLYCEROPHOSPHATE; SOYBEAN OIL

EMULSION; INTRAVENOUS

KABIVEN IN PLASTIC CONTAINER

+	FRESENIUS KABI USA	3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML	N200656 004	Aug 25, 2014
+		; 147MG/100ML; 3.9GM/100ML (1026ML)		
+		3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9GM/100ML (1540ML)	N200656 005	Aug 25, 2014
+		3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9GM/100ML (2053ML)	N200656 006	Aug 25, 2014
+		3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9GM/100ML (2566ML)	N200656 007	Aug 25, 2014
PERIKABIVEN IN PLASTIC CONTAINER				
+	FRESENIUS KABI USA	2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5GM/100ML (1440ML)	N200656 001	Aug 25, 2014
+		2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5GM/100ML (1920ML)	N200656 002	Aug 25, 2014
+		2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5GM/100ML (2400ML)	N200656 003	Aug 25, 2014

PRESCRIPTION DRUG PRODUCT LIST

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER BAXTER HLTHCARE 2.75%;10GM/100ML	N020734 002	Sep 29, 1997
CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER BAXTER HLTHCARE 2.75%;25GM/100ML	N020734 005	Sep 29, 1997
CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER BAXTER HLTHCARE 2.75%;5GM/100ML	N020734 001	Sep 29, 1997
CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER BAXTER HLTHCARE 4.25%;10GM/100ML	N020734 008	Sep 29, 1997
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER BAXTER HLTHCARE 4.25%;20GM/100ML	N020734 010	Sep 29, 1997
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER BAXTER HLTHCARE 4.25%;25GM/100ML	N020734 011	Sep 29, 1997
CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER BAXTER HLTHCARE 4.25%;5GM/100ML	N020734 007	Sep 29, 1997
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER BAXTER HLTHCARE 5%;10GM/100ML	N020734 014	Sep 29, 1997
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER BAXTER HLTHCARE 5%;15GM/100ML	N020734 015	Sep 29, 1997
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER BAXTER HLTHCARE 5%;20GM/100ML	N020734 016	Sep 29, 1997
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER BAXTER HLTHCARE 5%;25GM/100ML	N020734 017	Sep 29, 1997
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER BAXTER HLTHCARE 5%;35GM/100ML	N020734 018	Sep 29, 1997

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMINOCAPROIC ACID

AP LUITPOLD <u>250MG/ML</u>	<u>A071192 001</u>	Dec 01, 1987
AP ! HOSPIRA <u>250MG/ML</u>	<u>A070010 001</u>	Mar 09, 1987

SOLUTION; ORAL

AMICAR

AA +! HIKMA <u>0.25GM/ML</u>	<u>N015230 002</u>	
<u>AMINOCAPROIC ACID</u>		
AA AMNEAL <u>0.25GM/ML</u>	<u>A212780 001</u>	Aug 23, 2019
AA ANNORA PHARMA <u>0.25GM/ML</u>	<u>A216464 001</u>	Nov 04, 2022
AA AUROBINDO PHARMA <u>0.25GM/ML</u>	<u>A216804 001</u>	Sep 26, 2022
AA BELCHER <u>0.25GM/ML</u>	<u>A213825 001</u>	Apr 08, 2021
AA CARNEGIE <u>0.25GM/ML</u>	<u>A214140 001</u>	Jan 26, 2021
AA FLORIDA <u>0.25GM/ML</u>	<u>A215510 001</u>	Mar 23, 2022
AA SUNNY <u>0.25GM/ML</u>	<u>A213213 001</u>	Dec 27, 2023
AA TARO <u>0.25GM/ML</u>	<u>A214458 001</u>	Mar 24, 2023
AA TULEX PHARMS INC <u>0.25GM/ML</u>	<u>A212494 001</u>	Aug 11, 2020

TABLET; ORAL

AMICAR

AB + HIKMA <u>500MG</u>	<u>N015197 001</u>	
AB + <u>1GM</u>	<u>N015197 002</u>	Jun 24, 2004
<u>AMINOCAPROIC ACID</u>		
AB AMNEAL <u>500MG</u>	<u>A212492 001</u>	Nov 26, 2019
AB ANI PHARMS <u>500MG</u>	<u>A211629 001</u>	Dec 14, 2020
AB CARNEGIE <u>500MG</u>	<u>A213928 001</u>	Feb 12, 2021
AB <u>1GM</u>	<u>A213928 002</u>	Feb 12, 2021
AB MSN <u>500MG</u>	<u>A212938 001</u>	Nov 06, 2020
AB <u>1GM</u>	<u>A212938 002</u>	Sep 13, 2023
AB OPTIMUS <u>500MG</u>	<u>A213944 001</u>	Sep 14, 2022
AB <u>1GM</u>	<u>A213944 002</u>	Sep 14, 2022
AB SUNNY <u>500MG</u>	<u>A209060 001</u>	Nov 27, 2018
AB ! <u>1GM</u>	<u>A209060 002</u>	Nov 27, 2018

AMINOLEVULINIC ACID HYDROCHLORIDE

FOR SOLUTION; ORAL

GLEOLAN +! NXDC 1.5GM/VIAL	N208630 001	Jun 06, 2017
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GEL; TOPICAL

AMELUZ +! BIOFRONTERA 10%	N208081 001	May 10, 2016
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SOLUTION; TOPICAL

LEVULAN +! DUSA 20%	N020965 001	Dec 03, 1999
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PRESCRIPTION DRUG PRODUCT LIST

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

! HOSPIRA

25MG/ML

A087242 001 Oct 26, 1983

AMINOSALICYLIC ACID

GRANULE, DELAYED RELEASE; ORAL

PASER

! EUROCEPT PHARMS

4GM/PACKET

A074346 001 Jun 30, 1994

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

AP	ACELLA	50MG/ML	A077610 001	Oct 30, 2008
AP		50MG/ML	A077834 001	Oct 30, 2008
AP	! FRESENIUS KABI USA	50MG/ML	A075761 001	Oct 15, 2002
AP	! GLAND PHARMA LTD	50MG/ML	A077161 001	Apr 20, 2005
AP	HIKMA FARMACEUTICA	50MG/ML	A077234 001	Feb 25, 2008
AP	HOSPIRA	50MG/ML	A203884 001	Nov 25, 2013
AP	! MYLAN INSTITUTIONAL	50MG/ML	A076217 001	Oct 15, 2002

NEXTERONE

+! BAXTER HLTHCARE

150MG/100ML (1.5MG/ML)

N022325 002 Nov 16, 2010

+!

360MG/200ML (1.8MG/ML)

N022325 003 Nov 16, 2010

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

AB	AUROBINDO PHARMA LTD	200MG	A204742 001	Jun 03, 2016
AB	CHARTWELL RX	100MG	A077069 003	Oct 04, 2016
AB		200MG	A077069 001	Apr 08, 2005
AB		400MG	A077069 002	Apr 08, 2005
AB	DR REDDYS LABS SA	100MG	A075389 002	Dec 28, 2017
AB		200MG	A075389 001	Jan 25, 2001
AB		400MG	A075389 003	Dec 28, 2017
AB	RUBICON	100MG	A078578 002	Feb 26, 2021
AB		200MG	A078578 001	Nov 06, 2008
AB		400MG	A078578 003	Feb 26, 2021
AB	TARO	100MG	A075424 002	Dec 18, 2002
AB	!	200MG	A075424 001	Mar 30, 2001
AB		400MG	A076362 001	Nov 29, 2002
AB	TEVA PHARMS	200MG	A074739 001	Nov 30, 1998
AB	UNICHEM	200MG	A213446 001	Jul 21, 2020
AB	ZYDUS PHARMS USA INC	100MG	A079029 002	Jan 06, 2023
AB		200MG	A079029 001	Sep 16, 2008
AB		400MG	A079029 003	Jan 06, 2023

PACERONE

AB	UPSHER SMITH LABS	100MG	A075135 002	Apr 12, 2005
AB		200MG	A075135 001	Apr 30, 1998
AB		400MG	A075135 003	Jul 02, 2020

AMIODARONE HYDROCHLORIDE

TARO

300MG

A076362 002 Dec 02, 2003

AMISULPRIDE

SOLUTION; INTRAVENOUS

BARHEMSYS

+! ACACIA

5MG/2ML (2.5MG/ML)

N209510 001 Feb 26, 2020

+!

10MG/4ML (2.5MG/ML)

N209510 002 Sep 01, 2020

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

AB	ACCORD HLTHCARE	10MG	A202446 001	Jun 04, 2014
AB		25MG	A202446 002	Jun 04, 2014
AB		50MG	A202446 003	Jun 04, 2014
AB		75MG	A202446 004	Jun 04, 2014
AB		100MG	A202446 005	Jun 04, 2014
AB		150MG	A202446 006	Jun 04, 2014
AB	CHEMISTRY HLTH	10MG	A216243 001	Jun 06, 2022
AB		25MG	A216243 002	Jun 06, 2022
AB		50MG	A216243 003	Jun 06, 2022
AB		75MG	A216243 004	Jun 06, 2022
AB		100MG	A216243 005	Jun 06, 2022
AB		150MG	A216243 006	Jun 06, 2022
AB	MANKIND PHARMA	10MG	A213999 001	Feb 19, 2021
AB		25MG	A213999 002	Feb 19, 2021

PRESCRIPTION DRUG PRODUCT LIST

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>		<u>50MG</u>	<u>A213999 003</u>	Feb 19, 2021
<u>AB</u>		<u>75MG</u>	<u>A213999 004</u>	Feb 19, 2021
<u>AB</u>		<u>100MG</u>	<u>A213999 005</u>	Feb 19, 2021
<u>AB</u>		<u>150MG</u>	<u>A213999 006</u>	Feb 19, 2021
<u>AB</u>	RISING	<u>10MG</u>	<u>A217411 001</u>	May 19, 2023
<u>AB</u>		<u>25MG</u>	<u>A217411 002</u>	May 19, 2023
<u>AB</u>		<u>50MG</u>	<u>A217411 003</u>	May 19, 2023
<u>AB</u>		<u>75MG</u>	<u>A217411 004</u>	May 19, 2023
<u>AB</u>		<u>100MG</u>	<u>A217411 005</u>	May 19, 2023
<u>AB</u>		<u>150MG</u>	<u>A217411 006</u>	May 19, 2023
<u>AB</u>	RUBICON	<u>10MG</u>	<u>A215376 001</u>	May 01, 2023
<u>AB</u>		<u>25MG</u>	<u>A215376 002</u>	May 01, 2023
<u>AB</u>		<u>50MG</u>	<u>A215376 003</u>	May 01, 2023
<u>AB</u>		<u>75MG</u>	<u>A215376 004</u>	May 01, 2023
<u>AB</u>		<u>100MG</u>	<u>A215376 005</u>	May 01, 2023
<u>AB</u>		<u>150MG</u>	<u>A215376 006</u>	May 01, 2023
<u>AB</u>	+ SANDOZ	<u>10MG</u>	<u>A085968 004</u>	
<u>AB</u>	+!	<u>25MG</u>	<u>A085968 002</u>	
<u>AB</u>	+	<u>50MG</u>	<u>A085968 001</u>	
<u>AB</u>	+	<u>75MG</u>	<u>A085968 006</u>	
<u>AB</u>	+	<u>100MG</u>	<u>A085968 003</u>	
<u>AB</u>	+	<u>150MG</u>	<u>A085968 005</u>	
<u>AB</u>	SUN PHARM INDS INC	<u>10MG</u>	<u>A089399 002</u>	Jul 14, 1987
<u>AB</u>		<u>25MG</u>	<u>A089399 001</u>	Jul 14, 1987
<u>AB</u>		<u>50MG</u>	<u>A089399 003</u>	Jul 14, 1987
<u>AB</u>		<u>75MG</u>	<u>A089399 004</u>	Jul 14, 1987
<u>AB</u>		<u>100MG</u>	<u>A089399 005</u>	Jul 14, 1987
<u>AB</u>		<u>150MG</u>	<u>A089399 006</u>	Jul 14, 1987
<u>AB</u>	UNICHEM	<u>10MG</u>	<u>A214548 001</u>	May 19, 2021
<u>AB</u>		<u>25MG</u>	<u>A214548 002</u>	May 19, 2021
<u>AB</u>		<u>50MG</u>	<u>A214548 003</u>	May 19, 2021
<u>AB</u>		<u>75MG</u>	<u>A214548 004</u>	May 19, 2021
<u>AB</u>		<u>100MG</u>	<u>A214548 005</u>	May 19, 2021
<u>AB</u>		<u>150MG</u>	<u>A214548 006</u>	May 19, 2021
<u>AB</u>	ZYDUS PHARMS	<u>10MG</u>	<u>A210086 001</u>	Oct 06, 2017
<u>AB</u>		<u>25MG</u>	<u>A210086 002</u>	Oct 06, 2017
<u>AB</u>		<u>50MG</u>	<u>A210086 003</u>	Oct 06, 2017
<u>AB</u>		<u>75MG</u>	<u>A210086 004</u>	Oct 06, 2017
<u>AB</u>		<u>100MG</u>	<u>A210086 005</u>	Oct 06, 2017
<u>AB</u>		<u>150MG</u>	<u>A210086 006</u>	Oct 06, 2017

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MICRO LABS	<u>EQ 12.5MG BASE;5MG</u>	<u>A211925 001</u>	Feb 02, 2022
<u>AB</u>		<u>EQ 25MG BASE;10MG</u>	<u>A211925 002</u>	Feb 02, 2022
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 12.5MG BASE;5MG</u>	<u>A071297 002</u>	Dec 10, 1986
<u>AB</u>	!	<u>EQ 25MG BASE;10MG</u>	<u>A071297 001</u>	Dec 10, 1986

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

	MYLAN	10MG; 2MG	A071443 002	Nov 10, 1988
		10MG; 4MG	A071443 003	Nov 10, 1988
	!	25MG; 2MG	A071443 004	Nov 10, 1988
	!	25MG; 4MG	A071443 005	Nov 10, 1988
	!	50MG; 4MG	A071443 001	Nov 10, 1988

AMLODIPINE BENZOATE

SUSPENSION; ORAL

KATERZIA

+!	AZURITY	EQ 1MG BASE/ML	N211340 001	Jul 08, 2019
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AMLODIPINE BESYLATE

SOLUTION; ORAL

NORLIQVA

+!	CMP DEV LLC	EQ 1MG BASE/ML	N214439 001	Feb 24, 2022
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TABLET; ORAL

AMLODIPINE BESYLATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 2.5MG BASE</u>	<u>A202553 001</u>	Apr 29, 2013
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A202553 002</u>	Apr 29, 2013

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202553 003</u>	Apr 29, 2013
<u>AB</u>	ALKEM	<u>EQ 2.5MG BASE</u>	<u>A078925 001</u>	May 04, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078925 002</u>	May 04, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078925 003</u>	May 04, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 2.5MG BASE</u>	<u>A078021 001</u>	Jul 17, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078021 002</u>	Jul 17, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078021 003</u>	Jul 17, 2007
<u>AB</u>	CHARTWELL RX	<u>EQ 2.5MG BASE</u>	<u>A076692 001</u>	Jul 20, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076692 002</u>	Jul 20, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076692 003</u>	Jul 20, 2007
<u>AB</u>	CHINA RESOURCES	<u>EQ 2.5MG BASE</u>	<u>A090752 003</u>	May 16, 2016
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090752 001</u>	Apr 15, 2011
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090752 002</u>	Apr 15, 2011
<u>AB</u>	CIPLA	<u>EQ 2.5MG BASE</u>	<u>A077073 001</u>	Sep 26, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077073 002</u>	Sep 26, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077073 003</u>	Sep 26, 2007
<u>AB</u>	COREPHARMA	<u>EQ 2.5MG BASE</u>	<u>A076719 001</u>	May 23, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076719 002</u>	May 23, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076719 003</u>	May 23, 2007
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 2.5MG BASE</u>	<u>A078552 001</u>	Apr 08, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078552 002</u>	Apr 08, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078552 003</u>	Apr 08, 2009
<u>AB</u>	INVAGEN PHARMS	<u>EQ 2.5MG BASE</u>	<u>A077955 001</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A206367 001</u>	Dec 10, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077955 002</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A206367 002</u>	Dec 10, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077955 003</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206367 003</u>	Dec 10, 2015
<u>AB</u>	LUPIN	<u>EQ 2.5MG BASE</u>	<u>A078043 001</u>	Jul 12, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078043 002</u>	Jul 12, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078043 003</u>	Jul 12, 2007
<u>AB</u>	ORBION PHARMS	<u>EQ 2.5MG BASE</u>	<u>A078453 001</u>	Jul 02, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078453 002</u>	Jul 02, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078453 003</u>	Jul 02, 2009
<u>AB</u>	OXFORD PHARMS	<u>EQ 2.5MG BASE</u>	<u>A078414 001</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078414 002</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078414 003</u>	Apr 07, 2010
<u>AB</u>	POLYGEN PHARMS	<u>EQ 2.5MG BASE</u>	<u>A207821 001</u>	Jul 11, 2016
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A207821 002</u>	Jul 11, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207821 003</u>	Jul 11, 2016
<u>AB</u>	STRIDES PHARMA	<u>EQ 2.5MG BASE</u>	<u>A077516 001</u>	Jul 11, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077516 002</u>	Jul 11, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077516 003</u>	Jul 11, 2007
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 2.5MG BASE</u>	<u>A077974 001</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077974 002</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077974 003</u>	Jul 09, 2007
<u>AB</u>	TEVA	<u>EQ 2.5MG BASE</u>	<u>A076846 001</u>	Jun 28, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076846 002</u>	Jun 28, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076846 003</u>	Jun 28, 2007
<u>AB</u>	UNICHEM	<u>EQ 2.5MG BASE</u>	<u>A203245 001</u>	Oct 21, 2013
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203245 002</u>	Oct 21, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203245 003</u>	Oct 21, 2013
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 2.5MG BASE</u>	<u>A078226 001</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078226 002</u>	Jul 09, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078226 003</u>	Jul 09, 2007
<u>NORVASC</u>				
<u>AB</u>	+ VIATRIS	<u>EQ 2.5MG BASE</u>	<u>N019787 001</u>	Jul 31, 1992
<u>AB</u>	+	<u>EQ 5MG BASE</u>	<u>N019787 002</u>	Jul 31, 1992
<u>AB</u>	+!	<u>EQ 10MG BASE</u>	<u>N019787 003</u>	Jul 31, 1992

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

<u>AB</u>	APOTEX	<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A205199 001</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A205199 002</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A205199 003</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A205199 004</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A205199 005</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A205199 006</u>	Nov 18, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A205199 007</u>	Nov 18, 2019

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A205199 008</u>	Nov 18, 2019
<u>AB</u>	DR REDDYS	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A203874 001</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A203874 002</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A203874 003</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A203874 004</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A203874 005</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A203874 006</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A203874 007</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A203874 008</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A203874 009</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A203874 010</u>	Mar 07, 2014
<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A203874 011</u>	Mar 07, 2014
<u>AB</u>	MYLAN	<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A200465 004</u>	Nov 29, 2013
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A200465 005</u>	Nov 29, 2013
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A200465 006</u>	Nov 29, 2013
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A200465 007</u>	Nov 29, 2013
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A200465 008</u>	Nov 29, 2013
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A200465 009</u>	Nov 29, 2013
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A200465 010</u>	Nov 29, 2013
<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A200465 011</u>	Nov 29, 2013
<u>AB</u>	ZYDUS PHARMS	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>A207762 001</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>A207762 002</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>A207762 003</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>A207762 004</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>A207762 005</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>A207762 006</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>A207762 007</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>A207762 008</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>A207762 009</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>A207762 010</u>	Jan 11, 2019
<u>AB</u>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>A207762 011</u>	Jan 11, 2019
<u>CADUET</u>				
<u>AB</u>	+	PHARMACIA	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<u>N021540 009</u> Jul 29, 2004
<u>AB</u>	+		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<u>N021540 010</u> Jul 29, 2004
<u>AB</u>	+		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<u>N021540 011</u> Jul 29, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<u>N021540 001</u> Jan 30, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<u>N021540 002</u> Jan 30, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<u>N021540 003</u> Jan 30, 2004
<u>AB</u>	+		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<u>N021540 004</u> Jan 30, 2004
<u>AB</u>	+		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<u>N021540 005</u> Jan 30, 2004
<u>AB</u>	+		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<u>N021540 006</u> Jan 30, 2004
<u>AB</u>	+		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<u>N021540 007</u> Jan 30, 2004
<u>AB</u>	+		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<u>N021540 008</u> Jan 30, 2004

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 2.5MG BASE;10MG</u>	<u>A091431 001</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A091431 002</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A091431 003</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A091431 004</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A091431 005</u>	Dec 30, 2013
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A091431 006</u>	Dec 30, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 2.5MG BASE;10MG</u>	<u>A202239 001</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A202239 002</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A202239 003</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A202239 004</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A202239 005</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202239 006</u>	Sep 05, 2012
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 2.5MG BASE;10MG</u>	<u>A077183 001</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A077183 002</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A077183 003</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A090149 001</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A077183 004</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A090149 002</u>	Jul 05, 2011
<u>AB</u>	LUPIN PHARMS	<u>EQ 2.5MG BASE;10MG</u>	<u>A078466 001</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A078466 002</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A078466 003</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A078466 005</u>	Jul 05, 2011

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A078466 004</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A078466 006</u>	Jul 05, 2011
<u>AB</u>	WATSON LABS	<u>EQ 2.5MG BASE;10MG</u>	<u>A077890 001</u>	Oct 14, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A077890 002</u>	Oct 14, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A077890 003</u>	Oct 14, 2010
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A077890 004</u>	Oct 14, 2010
<u>AB</u>	WATSON LABS INC	<u>EQ 5MG BASE;40MG</u>	<u>A090364 001</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A090364 002</u>	Jul 05, 2011

LOTREL

<u>AB</u>	+	SANDOZ	<u>EQ 2.5MG BASE;10MG</u>	<u>N020364 002</u>	Mar 03, 1995
<u>AB</u>	+		<u>EQ 5MG BASE;10MG</u>	<u>N020364 003</u>	Mar 03, 1995
<u>AB</u>	+		<u>EQ 5MG BASE;20MG</u>	<u>N020364 004</u>	Mar 03, 1995
<u>AB</u>	+		<u>EQ 5MG BASE;40MG</u>	<u>N020364 007</u>	Apr 11, 2006
<u>AB</u>	+		<u>EQ 10MG BASE;20MG</u>	<u>N020364 005</u>	Jun 20, 2002
<u>AB</u>	+		<u>EQ 10MG BASE;40MG</u>	<u>N020364 006</u>	Apr 11, 2006

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE

<u>AB</u>		PAR PHARM INC	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A206137 001</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A206137 002</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 5MG BASE;25MG;40MG</u>	<u>A206137 003</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A206137 004</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 10MG BASE;25MG;40MG</u>	<u>A206137 005</u>	Oct 26, 2016
<u>AB</u>		TEVA PHARMS USA	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A202491 001</u>	Nov 03, 2016
<u>AB</u>			<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A202491 002</u>	Nov 03, 2016
<u>AB</u>			<u>EQ 5MG BASE;25MG;40MG</u>	<u>A202491 003</u>	Nov 03, 2016
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A202491 004</u>	Nov 03, 2016
<u>AB</u>			<u>EQ 10MG BASE;25MG;40MG</u>	<u>A202491 005</u>	Nov 03, 2016
<u>AB</u>		TORRENT	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A203580 001</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A203580 002</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 5MG BASE;25MG;40MG</u>	<u>A203580 003</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A203580 004</u>	Oct 26, 2016
<u>AB</u>			<u>EQ 10MG BASE;25MG;40MG</u>	<u>A203580 005</u>	Oct 26, 2016

TRIBENZOR

<u>AB</u>	+	COSETTE	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>N200175 001</u>	Jul 23, 2010
<u>AB</u>	+		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>N200175 002</u>	Jul 23, 2010
<u>AB</u>	+		<u>EQ 5MG BASE;25MG;40MG</u>	<u>N200175 003</u>	Jul 23, 2010
<u>AB</u>	+		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>N200175 004</u>	Jul 23, 2010
<u>AB</u>	+		<u>EQ 10MG BASE;25MG;40MG</u>	<u>N200175 005</u>	Jul 23, 2010

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET;ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		AUROBINDO PHARMA	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A206180 001</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 5MG BASE;25MG;160MG</u>	<u>A206180 002</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A206180 003</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 10MG BASE;25MG;160MG</u>	<u>A206180 004</u>	Dec 19, 2017
<u>AB</u>			<u>EQ 10MG BASE;25MG;320MG</u>	<u>A206180 005</u>	Dec 19, 2017
<u>AB</u>		LUPIN LTD	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A200797 001</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 5MG BASE;25MG;160MG</u>	<u>A200797 002</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A200797 003</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 10MG BASE;25MG;160MG</u>	<u>A200797 004</u>	Jun 03, 2015
<u>AB</u>			<u>EQ 10MG BASE;25MG;320MG</u>	<u>A200797 005</u>	Jun 03, 2015
<u>AB</u>		STRIDES PHARMA	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>A201087 001</u>	Jun 01, 2015
<u>AB</u>			<u>EQ 5MG BASE;25MG;160MG</u>	<u>A201087 002</u>	Jun 01, 2015
<u>AB</u>			<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>A201087 003</u>	Jun 01, 2015
<u>AB</u>			<u>EQ 10MG BASE;25MG;160MG</u>	<u>A201087 004</u>	Jun 01, 2015
<u>AB</u>			<u>EQ 10MG BASE;25MG;320MG</u>	<u>A201087 005</u>	Jun 01, 2015
<u>EXFORGE HCT</u>					
<u>AB</u>	+	NOVARTIS	<u>EQ 5MG BASE;12.5MG;160MG</u>	<u>N022314 001</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 5MG BASE;25MG;160MG</u>	<u>N022314 002</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 10MG BASE;12.5MG;160MG</u>	<u>N022314 003</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 10MG BASE;25MG;160MG</u>	<u>N022314 004</u>	Apr 30, 2009
<u>AB</u>	+		<u>EQ 10MG BASE;25MG;320MG</u>	<u>N022314 005</u>	Apr 30, 2009

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

AB	AJANTA PHARMA LTD	EQ 5MG BASE;20MG	A207216 001	Oct 28, 2016
AB		EQ 5MG BASE;40MG	A207216 002	Oct 28, 2016
AB		EQ 10MG BASE;20MG	A207216 003	Oct 28, 2016
AB		EQ 10MG BASE;40MG	A207216 004	Oct 28, 2016
AB	ALEMBIC	EQ 5MG BASE;20MG	A207073 001	Jul 17, 2017
AB		EQ 5MG BASE;40MG	A207073 002	Jul 17, 2017
AB		EQ 10MG BASE;20MG	A207073 003	Jul 17, 2017
AB		EQ 10MG BASE;40MG	A207073 004	Jul 17, 2017
AB	ALKEM LABS LTD	EQ 5MG BASE;20MG	A209042 001	Aug 14, 2017
AB		EQ 5MG BASE;40MG	A209042 002	Aug 14, 2017
AB		EQ 10MG BASE;20MG	A209042 003	Aug 14, 2017
AB		EQ 10MG BASE;40MG	A209042 004	Aug 14, 2017
AB	AUROBINDO PHARMA	EQ 5MG BASE;20MG	A206906 001	May 15, 2017
AB		EQ 5MG BASE;40MG	A206906 002	May 15, 2017
AB		EQ 10MG BASE;20MG	A206906 003	May 15, 2017
AB		EQ 10MG BASE;40MG	A206906 004	May 15, 2017
AB	GLENMARK PHARMS LTD	EQ 5MG BASE;20MG	A207807 001	Jul 05, 2017
AB		EQ 5MG BASE;40MG	A207807 002	Jul 05, 2017
AB		EQ 10MG BASE;20MG	A207807 003	Jul 05, 2017
AB		EQ 10MG BASE;40MG	A207807 004	Jul 05, 2017
AB	MACLEODS PHARMS LTD	EQ 5MG BASE;20MG	A206884 001	Oct 26, 2016
AB		EQ 5MG BASE;40MG	A206884 003	Oct 26, 2016
AB		EQ 10MG BASE;20MG	A206884 002	Oct 26, 2016
AB		EQ 10MG BASE;40MG	A206884 004	Oct 26, 2016
AB	MICRO LABS	EQ 5MG BASE;20MG	A207435 001	Nov 02, 2017
AB		EQ 5MG BASE;40MG	A207435 002	Nov 02, 2017
AB		EQ 10MG BASE;20MG	A207435 003	Nov 02, 2017
AB		EQ 10MG BASE;40MG	A207435 004	Nov 02, 2017
AB	ZYDUS PHARMS	EQ 5MG BASE;20MG	A207771 001	Sep 22, 2017
AB		EQ 5MG BASE;40MG	A207771 002	Sep 22, 2017
AB		EQ 10MG BASE;20MG	A207771 003	Sep 22, 2017
AB		EQ 10MG BASE;40MG	A207771 004	Sep 22, 2017
AZOR				
AB	+ COSETTE	EQ 5MG BASE;20MG	N022100 001	Sep 26, 2007
AB	+	EQ 5MG BASE;40MG	N022100 002	Sep 26, 2007
AB	+	EQ 10MG BASE;20MG	N022100 003	Sep 26, 2007
AB	+ !	EQ 10MG BASE;40MG	N022100 004	Sep 26, 2007

AMLODIPINE BESYLATE; PERINDOPRIL ARGININE

TABLET; ORAL

PRESTALIA

+	ADHERA	EQ 2.5MG BASE;3.5MG	N205003 001	Jan 21, 2015
+		EQ 5MG BASE;7MG	N205003 002	Jan 21, 2015
+ !		EQ 10MG BASE;14MG	N205003 003	Jan 21, 2015

AMLODIPINE BESYLATE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND AMLODIPINE

AB	LUPIN LTD	EQ 5MG BASE;40MG	A201586 001	Jan 08, 2014
AB		EQ 5MG BASE;80MG	A201586 003	Jan 08, 2014
AB		EQ 10MG BASE;40MG	A201586 002	Jan 08, 2014
AB		EQ 10MG BASE;80MG	A201586 004	Jan 08, 2014
AB	MYLAN	EQ 5MG BASE;40MG	A202516 001	Aug 26, 2014
AB		EQ 5MG BASE;80MG	A202516 003	Aug 26, 2014
AB		EQ 10MG BASE;40MG	A202516 002	Aug 26, 2014
AB	!	EQ 10MG BASE;80MG	A202516 004	Aug 26, 2014

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

AB	ALEMBIC	EQ 5MG BASE;160MG	A202713 001	Apr 03, 2015
AB		EQ 5MG BASE;320MG	A202713 003	Apr 03, 2015
AB		EQ 10MG BASE;160MG	A202713 002	Apr 03, 2015
AB		EQ 10MG BASE;320MG	A202713 004	Apr 03, 2015
AB	AUROBINDO PHARMA	EQ 5MG BASE;160MG	A206512 001	Apr 22, 2016
AB		EQ 5MG BASE;320MG	A206512 002	Apr 22, 2016
AB		EQ 10MG BASE;160MG	A206512 003	Apr 22, 2016
AB		EQ 10MG BASE;320MG	A206512 004	Apr 22, 2016
AB	HETERO LABS	EQ 5MG BASE;160MG	A205137 001	Sep 16, 2016
AB		EQ 5MG BASE;320MG	A205137 003	Sep 16, 2016
AB		EQ 10MG BASE;160MG	A205137 002	Sep 16, 2016

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A205137 004</u>	Sep 16, 2016
<u>AB</u>	LUPIN	<u>EQ 5MG BASE;160MG</u>	<u>A090245 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090245 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090245 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090245 004</u>	Mar 30, 2015
<u>AB</u>	MYLAN	<u>EQ 5MG BASE;160MG</u>	<u>A090483 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090483 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090483 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090483 004</u>	Mar 30, 2015
<u>AB</u>	NOVEL LABS INC	<u>EQ 5MG BASE;160MG</u>	<u>A202829 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A202829 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A202829 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A202829 004</u>	Mar 30, 2015
<u>AB</u>	STRIDES PHARMA	<u>EQ 5MG BASE;160MG</u>	<u>A090011 001</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090011 003</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090011 002</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090011 004</u>	Mar 28, 2013
<u>EXFORGE</u>				
<u>AB</u>	+ NOVARTIS	<u>EQ 5MG BASE;160MG</u>	<u>N021990 002</u>	Jun 20, 2007
<u>AB</u>	+	<u>EQ 5MG BASE;320MG</u>	<u>N021990 004</u>	Jun 20, 2007
<u>AB</u>	+	<u>EQ 10MG BASE;160MG</u>	<u>N021990 003</u>	Jun 20, 2007
<u>AB</u>	+	<u>EQ 10MG BASE;320MG</u>	<u>N021990 005</u>	Jun 20, 2007

AMMONIA N-13

INJECTABLE; INTRAVENOUS

AMMONIA N 13

<u>AP</u>	3D IMAGING DRUG	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203779 001</u>	Oct 19, 2015
<u>AP</u>	BIOMEDCL RES FDN	<u>48.75mCi-487.5mCi/13ML (3.75-37.5mCi/ML)</u>	<u>A204352 001</u>	May 01, 2015
<u>AP</u>	BRIGHAM WOMENS HOSP	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203783 001</u>	Oct 30, 2014
<u>AP</u>	CARDINAL HLTH 414	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203700 001</u>	Feb 25, 2013
<u>AP</u>	DECATUR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204465 001</u>	Oct 23, 2014
<u>AP</u>	+ FEINSTEIN	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>N022119 001</u>	Aug 23, 2007
<u>AP</u>	GEN HOSP	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A207025 001</u>	Feb 03, 2016
<u>AP</u>	IONETIX	<u>22.5mCi-225mCi/6ML (3.75-37.5mCi/ML)</u>	<u>A210524 001</u>	Dec 21, 2018
<u>AP</u>	JOHNS HOPKINS UNIV	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204514 001</u>	Aug 19, 2014
<u>AP</u>	KREITCHMAN PET CTR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203938 001</u>	Dec 09, 2013
<u>AP</u>	MCPRF	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203321 001</u>	Feb 25, 2013
<u>AP</u>	METHODIST	<u>3.75mCi-260mCi/ML</u>	<u>A215083 001</u>	Jul 09, 2021
<u>AP</u>	MIDWEST MEDCL	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204457 001</u>	Nov 18, 2015
<u>AP</u>	MIPS CRF	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204535 001</u>	Nov 20, 2014
<u>AP</u>	NCM USA BRONX LLC	<u>3.75mCi-260mCi/ML</u>	<u>A204515 001</u>	Feb 04, 2015
<u>AP</u>	NUKEMED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204455 001</u>	Apr 23, 2015
<u>AP</u>	PETNET	<u>30mCi-300mCi (3.75-37.5mCi/ML)</u>	<u>A204510 001</u>	Nov 02, 2015
<u>AP</u>	PRECISION NUCLEAR	<u>3.75mCi-260mCi/ML</u>	<u>A204547 001</u>	Aug 14, 2015
<u>AP</u>	! SOFIE	<u>3.75mCi-260mCi/ML</u>	<u>A203543 001</u>	Dec 14, 2012
<u>AP</u>	UCLA BIOMEDICAL	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203812 001</u>	Jun 27, 2013
<u>AP</u>	UCSF RODIOPHARM	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204496 001</u>	Mar 28, 2014
<u>AP</u>	UNIV ALAHAMA BIRM	<u>15mCi-150mCi/4ML (3.75-37.5mCi/ML)</u>	<u>A211698 001</u>	Nov 03, 2022
<u>AP</u>	UNIV TX SW MEDCTR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A209507 001</u>	Nov 01, 2019
<u>AP</u>	UNIV WISCONSIN	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A211740 001</u>	Sep 09, 2020
<u>AP</u>	WA UNIV SCH MED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204506 001</u>	Feb 07, 2014

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE IN PLASTIC CONTAINER

! HOSPIRA 5MEQ/ML A088366 001 Jun 13, 1984

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	! PADAGIS ISRAEL	<u>EQ 12% BASE</u>	<u>A075774 001</u>	May 01, 2002
<u>AB</u>	TARO	<u>EQ 12% BASE</u>	<u>A075883 001</u>	Apr 10, 2003

LOTION; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	! PADAGIS ISRAEL	<u>EQ 12% BASE</u>	<u>A075570 001</u>	Jun 23, 2004
<u>AB</u>	TARO	<u>EQ 12% BASE</u>	<u>A076216 001</u>	May 28, 2004

PRESCRIPTION DRUG PRODUCT LIST

AMOXAPINE

TABLET; ORAL

AMOXAPINE

WATSON LABS	25MG	A072691 002	Aug 28, 1992
	50MG	A072691 003	Aug 28, 1992
	100MG	A072691 004	Aug 28, 1992
!	150MG	A072691 001	Aug 28, 1992

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>250MG</u>	<u>A065271 001</u>	Nov 09, 2005
<u>AB</u>		<u>500MG</u>	<u>A065271 002</u>	Nov 09, 2005
<u>AB</u>	CHARTWELL	<u>250MG</u>	<u>A062058 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A062058 002</u>	
<u>AB</u>	HIKMA PHARMS	<u>250MG</u>	<u>A065291 001</u>	Feb 05, 2007
<u>AB</u>		<u>500MG</u>	<u>A065291 002</u>	Feb 05, 2007
<u>AB</u>	MICRO LABS	<u>250MG</u>	<u>A207471 001</u>	Jun 24, 2022
<u>AB</u>		<u>500MG</u>	<u>A207471 002</u>	Jun 24, 2022
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A064076 001</u>	Sep 30, 1994
<u>AB</u>		<u>500MG</u>	<u>A064076 002</u>	Sep 30, 1994
<u>AB</u>	TEVA	<u>250MG</u>	<u>A061926 001</u>	
<u>AB</u>	!	<u>500MG</u>	<u>A061926 003</u>	

AMOXIL

<u>AB</u>	US ANTIBIOTICS	<u>250MG</u>	<u>A062216 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A062216 004</u>	

FOR SUSPENSION; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>200MG/5ML</u>	<u>A065334 001</u>	Dec 28, 2006
<u>AB</u>		<u>400MG/5ML</u>	<u>A065334 002</u>	Dec 28, 2006
<u>AB</u>	AUROBINDO PHARMA LTD	<u>125MG/5ML</u>	<u>A204030 001</u>	Sep 15, 2014
<u>AB</u>		<u>250MG/5ML</u>	<u>A204030 002</u>	Sep 15, 2014
<u>AB</u>	DAVA PHARMS INC	<u>125MG/5ML</u>	<u>A062927 001</u>	Nov 25, 1988
<u>AB</u>		<u>250MG/5ML</u>	<u>A062927 002</u>	Nov 25, 1988
<u>AB</u>	HIKMA	<u>125MG/5ML</u>	<u>A065322 002</u>	Jun 19, 2006
<u>AB</u>		<u>200MG/5ML</u>	<u>A065325 002</u>	Jun 19, 2006
<u>AB</u>		<u>250MG/5ML</u>	<u>A065322 001</u>	Jun 19, 2006
<u>AB</u>		<u>400MG/5ML</u>	<u>A065325 001</u>	Jun 19, 2006
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>A065387 001</u>	Mar 26, 2007
<u>AB</u>		<u>200MG/5ML</u>	<u>A065378 001</u>	Mar 26, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065387 002</u>	Mar 26, 2007
<u>AB</u>		<u>400MG/5ML</u>	<u>A065378 002</u>	Mar 26, 2007
<u>AB</u>	TEVA	<u>200MG/5ML</u>	<u>A065119 001</u>	Dec 04, 2002
<u>AB</u>	!	<u>250MG/5ML</u>	<u>A061931 002</u>	
<u>AB</u>	!	<u>400MG/5ML</u>	<u>A065119 002</u>	Dec 04, 2002
<u>AB</u>	WOCKHARDT BIO AG	<u>400MG/5ML</u>	<u>A065319 002</u>	Jun 18, 2007

AMOXICILLIN PEDIATRIC

<u>AB</u>	TEVA	<u>50MG/ML</u>	<u>A061931 003</u>	Dec 01, 1982
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AMOXIL

<u>AB</u>	US ANTIBIOTICS	<u>50MG/ML</u>	<u>A062226 005</u>	
<u>AB</u>		<u>125MG/5ML</u>	<u>A062226 001</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>A062226 002</u>	

LAROTID

<u>AB</u>	US ANTIBIOTICS	<u>125MG/5ML</u>	<u>A062226 003</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>A062226 004</u>	

TABLET; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>A065256 001</u>	Nov 09, 2005
<u>AB</u>		<u>875MG</u>	<u>A065256 002</u>	Nov 09, 2005
<u>AB</u>	HIKMA	<u>875MG</u>	<u>A065255 001</u>	Mar 29, 2006
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065228 001</u>	Jul 13, 2005
<u>AB</u>		<u>875MG</u>	<u>A065228 002</u>	Jul 13, 2005
<u>AB</u>	TEVA	<u>500MG</u>	<u>A065056 001</u>	Sep 18, 2000
<u>AB</u>	!	<u>875MG</u>	<u>A065056 002</u>	Sep 18, 2000

TABLET, CHEWABLE; ORAL

AMOXICILLIN

TEVA	125MG	A064013 002	Sep 11, 1995
!	250MG	A064013 001	Dec 22, 1992

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED REL PELLETS; ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED)

<u>AB</u>	!	RISING	<u>500MG;500MG;30MG</u>	<u>A206006</u>	<u>001</u>	Oct 07, 2016
<u>AB</u>		SANDOZ	<u>500MG;500MG;30MG</u>	<u>A202588</u>	<u>001</u>	Mar 04, 2014

AMOXICILLIN; CLARITHROMYCIN; VONOPRAZAN FUMARATE

CAPSULE, TABLET, TABLET; ORAL

VOQUEZNA TRIPLE PAK

+	!	PHATHOM	500MG;500MG;EQ 20MG BASE	N215152	001	May 03, 2022
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AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>		AUROBINDO PHARMA	<u>125MG/5ML;EQ 31.25MG BASE/5ML</u>	<u>A209371</u>	<u>001</u>	Apr 19, 2019
<u>AB</u>			<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A209371</u>	<u>002</u>	Apr 19, 2019
<u>AB</u>		AUROBINDO PHARMA LTD	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A201090</u>	<u>001</u>	Dec 20, 2011
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A201090</u>	<u>002</u>	Dec 20, 2011
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A201091</u>	<u>001</u>	Dec 20, 2011
<u>AB</u>		DEVA HOLDING AS	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A210374</u>	<u>001</u>	Nov 29, 2023
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A210416</u>	<u>001</u>	Dec 11, 2023
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A209351</u>	<u>001</u>	Sep 13, 2023
<u>AB</u>		HIKMA PHARMS	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065191</u>	<u>002</u>	Jan 25, 2005
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065191</u>	<u>001</u>	Jan 25, 2005
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065373</u>	<u>001</u>	Nov 09, 2007
<u>AB</u>		MICRO LABS	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A217805</u>	<u>001</u>	Dec 08, 2023
<u>AB</u>		MICRO LABS LTD INDIA	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A205187</u>	<u>001</u>	May 21, 2021
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A205187</u>	<u>002</u>	May 21, 2021
<u>AB</u>		SANDOZ	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065066</u>	<u>001</u>	Jun 05, 2002
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065066</u>	<u>002</u>	Jun 05, 2002
<u>AB</u>		SANDOZ INC	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065098</u>	<u>001</u>	Dec 16, 2002
<u>AB</u>			<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065098</u>	<u>002</u>	Dec 16, 2002
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065358</u>	<u>001</u>	Aug 13, 2007
<u>AB</u>		TEVA	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065089</u>	<u>001</u>	May 25, 2004
<u>AB</u>	!		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065089</u>	<u>002</u>	May 25, 2004
<u>AB</u>	!		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065162</u>	<u>001</u>	Mar 12, 2004
<u>AB</u>		WOCKHARDT BIO AG	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A065431</u>	<u>001</u>	Nov 25, 2008
<u>AB</u>			<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065420</u>	<u>001</u>	Dec 02, 2013

AUGMENTIN '125'

<u>AB</u>	+	US ANTIBIOTICS	<u>125MG/5ML;EQ 31.25MG BASE/5ML</u>	<u>N050575</u>	<u>001</u>	Aug 06, 1984
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AUGMENTIN '250'

<u>AB</u>	+	!	US ANTIBIOTICS	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>N050575</u>	<u>002</u>	Aug 06, 1984
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AUGMENTIN ES-600

<u>AB</u>	+	US ANTIBIOTICS	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>N050755</u>	<u>001</u>	Jun 22, 2001
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TABLET; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>		AUROBINDO PHARMA LTD	<u>250MG;EQ 125MG BASE</u>	<u>A091569</u>	<u>001</u>	Jan 20, 2012
<u>AB</u>			<u>500MG;EQ 125MG BASE</u>	<u>A091569</u>	<u>002</u>	Jan 20, 2012
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A091568</u>	<u>001</u>	Jan 20, 2012
<u>AB</u>		DEVA HOLDING AS	<u>500MG;EQ 125MG BASE</u>	<u>A209992</u>	<u>001</u>	Sep 15, 2023
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A209991</u>	<u>001</u>	Sep 15, 2023
<u>AB</u>		HIKMA PHARMS	<u>875MG;EQ 125MG BASE</u>	<u>A203824</u>	<u>001</u>	Aug 23, 2016
<u>AB</u>		MICRO LABS LTD INDIA	<u>250MG;EQ 125MG BASE</u>	<u>A205707</u>	<u>001</u>	Dec 30, 2016
<u>AB</u>			<u>500MG;EQ 125MG BASE</u>	<u>A205707</u>	<u>002</u>	Dec 30, 2016
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A204755</u>	<u>003</u>	Dec 30, 2016
<u>AB</u>	!	SANDOZ	<u>250MG;EQ 125MG BASE</u>	<u>A065189</u>	<u>001</u>	Aug 23, 2005
<u>AB</u>			<u>500MG;EQ 125MG BASE</u>	<u>A065064</u>	<u>001</u>	Mar 15, 2002
<u>AB</u>	!		<u>875MG;EQ 125MG BASE</u>	<u>A065063</u>	<u>001</u>	Mar 14, 2002
<u>AB</u>	!	SANDOZ INC	<u>500MG;EQ 125MG BASE</u>	<u>A065117</u>	<u>001</u>	Nov 27, 2002
<u>AB</u>			<u>875MG;EQ 125MG BASE</u>	<u>A065093</u>	<u>001</u>	Nov 21, 2002
<u>AB</u>		TEVA	<u>500MG;EQ 125MG BASE</u>	<u>A065101</u>	<u>001</u>	Oct 30, 2002
<u>AB</u>		TEVA PHARMS USA	<u>875MG;EQ 125MG BASE</u>	<u>A065096</u>	<u>001</u>	Oct 29, 2002

AUGMENTIN '875'

<u>AB</u>	+	US ANTIBIOTICS	<u>875MG;EQ 125MG BASE</u>	<u>N050720</u>	<u>001</u>	Feb 13, 1996
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TABLET, CHEWABLE; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

		TEVA	200MG;EQ 28.5MG BASE	A065205	001	Feb 09, 2005
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!			400MG;EQ 57MG BASE	A065205	002	Feb 09, 2005
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PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN; CLAVULANATE POTASSIUM

TABLET, EXTENDED RELEASE;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

! SANDOZ

1GM;EQ 62.5MG BASE

A090227 001 Apr 21, 2010

AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN

CAPSULE, DELAYED RELEASE;ORAL

TALICIA

+! REDHILL

250MG;EQ 10MG BASE;12.5MG

N213004 001 Nov 01, 2019

AMOXICILLIN; VONOPRAZAN FUMARATE

CAPSULE, TABLET;ORAL

VOQUEZNA DUAL PAK

+! PHATHOM

500MG;EQ 20MG BASE

N215153 001 May 03, 2022

AMPHETAMINE

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL

ADZENYS XR-ODT

+ NEOS THERAPS

EQ 3.1MG BASE

N204326 001 Jan 27, 2016

+

EQ 6.3MG BASE

N204326 002 Jan 27, 2016

+

EQ 9.4MG BASE

N204326 003 Jan 27, 2016

+

EQ 12.5MG BASE

N204326 004 Jan 27, 2016

+

EQ 15.7MG BASE

N204326 005 Jan 27, 2016

+!

EQ 18.8MG BASE

N204326 006 Jan 27, 2016

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

ADDERALL XR 10AB + TAKEDA PHARMS USA2.5MG;2.5MG;2.5MG;2.5MGN021303 001 Oct 11, 2001ADDERALL XR 15AB + TAKEDA PHARMS USA3.75MG;3.75MG;3.75MG;3.75MGN021303 006 May 22, 2002ADDERALL XR 20AB + TAKEDA PHARMS USA5MG;5MG;5MG;5MGN021303 002 Oct 11, 2001ADDERALL XR 25AB + TAKEDA PHARMS USA6.25MG;6.25MG;6.25MG;6.25MGN021303 004 May 22, 2002ADDERALL XR 30AB +! TAKEDA PHARMS USA7.5MG;7.5MG;7.5MG;7.5MGN021303 003 Oct 11, 2001ADDERALL XR 5AB + TAKEDA PHARMS USA1.25MG;1.25MG;1.25MG;1.25MGN021303 005 May 22, 2002DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATEAB ACTAVIS ELIZABETH1.25MG;1.25MG;1.25MG;1.25MGA077302 001 Jun 22, 2012AB2.5MG;2.5MG;2.5MG;2.5MGA077302 002 Jun 22, 2012AB3.75MG;3.75MG;3.75MG;3.75MGA077302 003 Jun 22, 2012AB5MG;5MG;5MG;5MGA077302 004 Jun 22, 2012AB6.25MG;6.25MG;6.25MG;6.25MGA077302 005 Jun 22, 2012AB7.5MG;7.5MG;7.5MG;7.5MGA077302 006 Jun 22, 2012AB ANI PHARMS1.25MG;1.25MG;1.25MG;1.25MGA205401 001 Jan 22, 2019AB2.5MG;2.5MG;2.5MG;2.5MGA205401 002 Jan 22, 2019AB3.75MG;3.75MG;3.75MG;3.75MGA205401 003 Jan 22, 2019AB5MG;5MG;5MG;5MGA205401 004 Jan 22, 2019AB6.25MG;6.25MG;6.25MG;6.25MGA205401 005 Jan 22, 2019AB7.5MG;7.5MG;7.5MG;7.5MGA205401 006 Jan 22, 2019AB ASCENT PHARMS INC1.25MG;1.25MG;1.25MG;1.25MGA214959 001 Sep 29, 2021AB2.5MG;2.5MG;2.5MG;2.5MGA214959 002 Sep 29, 2021AB3.75MG;3.75MG;3.75MG;3.75MGA214959 003 Sep 29, 2021AB5MG;5MG;5MG;5MGA214959 004 Sep 29, 2021AB6.25MG;6.25MG;6.25MG;6.25MGA214959 005 Sep 29, 2021AB7.5MG;7.5MG;7.5MG;7.5MGA214959 006 Sep 29, 2021AB ELITE LABS INC1.25MG;1.25MG;1.25MG;1.25MGA212037 001 Dec 11, 2019AB2.5MG;2.5MG;2.5MG;2.5MGA212037 002 Dec 11, 2019AB3.75MG;3.75MG;3.75MG;3.75MGA212037 003 Dec 11, 2019AB5MG;5MG;5MG;5MGA212037 004 Dec 11, 2019AB6.25MG;6.25MG;6.25MG;6.25MGA212037 005 Dec 11, 2019AB7.5MG;7.5MG;7.5MG;7.5MGA212037 006 Dec 11, 2019AB GRANULES1.25MG;1.25MG;1.25MG;1.25MGA217027 001 Jan 23, 2023AB2.5MG;2.5MG;2.5MG;2.5MGA217027 002 Jan 23, 2023AB3.75MG;3.75MG;3.75MG;3.75MGA217027 003 Jan 23, 2023AB5MG;5MG;5MG;5MGA217027 004 Jan 23, 2023AB6.25MG;6.25MG;6.25MG;6.25MGA217027 005 Jan 23, 2023AB7.5MG;7.5MG;7.5MG;7.5MGA217027 006 Jan 23, 2023AB IMPAX LABS1.25MG;1.25MG;1.25MG;1.25MGA076852 001 Feb 16, 2016AB2.5MG;2.5MG;2.5MG;2.5MGA076852 002 Feb 16, 2016AB3.75MG;3.75MG;3.75MG;3.75MGA076852 003 Feb 16, 2016AB5MG;5MG;5MG;5MGA076852 004 Feb 16, 2016

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A076852	005	Feb 16, 2016	
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A076852	006	Feb 16, 2016	
AB	LANNETT CO INC	1.25MG; 1.25MG; 1.25MG; 1.25MG	A214403	001	Nov 26, 2021	
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A214403	002	Nov 26, 2021	
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A214403	003	Nov 26, 2021	
AB		5MG; 5MG; 5MG; 5MG	A214403	004	Nov 26, 2021	
AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A214403	005	Nov 26, 2021	
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A214403	006	Nov 26, 2021	
AB	RHODES PHARMS	1.25MG; 1.25MG; 1.25MG; 1.25MG	A210651	001	May 17, 2019	
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A210651	002	May 17, 2019	
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A210651	003	May 17, 2019	
AB		5MG; 5MG; 5MG; 5MG	A210651	004	May 17, 2019	
AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A210651	005	May 17, 2019	
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A210651	006	May 17, 2019	
AB	SPECGX LLC	1.25MG; 1.25MG; 1.25MG; 1.25MG	A211547	001	Apr 22, 2019	
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A211547	002	Apr 22, 2019	
AB		3.125MG; 3.125MG; 3.125MG; 3.125MG	A211546	001	Aug 31, 2023	
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A211547	003	Apr 22, 2019	
AB		5MG; 5MG; 5MG; 5MG	A211547	004	Apr 22, 2019	
AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A211546	002	Aug 31, 2023	
AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A211547	005	Apr 22, 2019	
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A211547	006	Apr 22, 2019	
AB		9.375MG; 9.375MG; 9.375MG; 9.375MG	A211546	003	Aug 31, 2023	
AB		12.5MG; 12.5MG; 12.5MG; 12.5MG	A211546	004	Aug 31, 2023	
AB	SUN PHARM INDS INC	3.125MG; 3.125MG; 3.125MG; 3.125MG	A215997	001	Sep 27, 2023	
AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A215997	002	Sep 27, 2023	
AB		9.375MG; 9.375MG; 9.375MG; 9.375MG	A215997	003	Sep 27, 2023	
AB		12.5MG; 12.5MG; 12.5MG; 12.5MG	A215997	004	Sep 27, 2023	
AB	SUN PHARM INDUSTRIES	1.25MG; 1.25MG; 1.25MG; 1.25MG	A211715	001	May 17, 2019	
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A211715	002	May 17, 2019	
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A211715	003	May 17, 2019	
AB		5MG; 5MG; 5MG; 5MG	A211715	004	May 17, 2019	
AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A211715	005	May 17, 2019	
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A211715	006	May 17, 2019	
AB	TEVA PHARMS USA	3.125MG; 3.125MG; 3.125MG; 3.125MG	A210876	001	Jan 31, 2022	
AB		6.25MG; 6.25MG; 6.25MG; 6.25MG	A210876	002	Jan 31, 2022	
AB		9.375MG; 9.375MG; 9.375MG; 9.375MG	A210876	003	Jan 31, 2022	
AB		12.5MG; 12.5MG; 12.5MG; 12.5MG	A210876	004	Jan 31, 2022	
MYDAYIS						
AB	+	TAKEDA PHARMS USA	3.125MG; 3.125MG; 3.125MG; 3.125MG	N022063	001	Jun 20, 2017
AB	+		6.25MG; 6.25MG; 6.25MG; 6.25MG	N022063	002	Jun 20, 2017
AB	+		9.375MG; 9.375MG; 9.375MG; 9.375MG	N022063	003	Jun 20, 2017
AB	+	!	12.5MG; 12.5MG; 12.5MG; 12.5MG	N022063	004	Jun 20, 2017

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB	ACTAVIS ELIZABETH	1.25MG; 1.25MG; 1.25MG; 1.25MG	A206340	001	Feb 05, 2016
AB		1.875MG; 1.875MG; 1.875MG; 1.875MG	A206340	002	Feb 05, 2016
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A206340	003	Feb 05, 2016
AB		3.125MG; 3.125MG; 3.125MG; 3.125MG	A206340	004	Feb 05, 2016
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A206340	005	Feb 05, 2016
AB		5MG; 5MG; 5MG; 5MG	A206340	006	Feb 05, 2016
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A206340	007	Feb 05, 2016
AB	ALKEM LABS LTD	1.25MG; 1.25MG; 1.25MG; 1.25MG	A217787	001	Oct 06, 2023
AB		1.875MG; 1.875MG; 1.875MG; 1.875MG	A217787	002	Oct 06, 2023
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A217787	003	Oct 06, 2023
AB		3.125MG; 3.125MG; 3.125MG; 3.125MG	A217787	004	Oct 06, 2023
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A217787	005	Oct 06, 2023
AB		5MG; 5MG; 5MG; 5MG	A217787	006	Oct 06, 2023
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A217787	007	Oct 06, 2023
AB	ALVOGEN	1.25MG; 1.25MG; 1.25MG; 1.25MG	A207388	001	Jul 28, 2017
AB		1.875MG; 1.875MG; 1.875MG; 1.875MG	A207388	002	Jul 28, 2017
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A207388	003	Jul 28, 2017
AB		3.125MG; 3.125MG; 3.125MG; 3.125MG	A207388	004	Jul 28, 2017
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A207388	005	Jul 28, 2017
AB		5MG; 5MG; 5MG; 5MG	A207388	006	Jul 28, 2017
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A207388	007	Jul 28, 2017
AB	ASCENT PHARMS INC	1.25MG; 1.25MG; 1.25MG; 1.25MG	A213709	001	Apr 22, 2021
AB		1.875MG; 1.875MG; 1.875MG; 1.875MG	A213709	002	Apr 22, 2021

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET;ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A213709 003</u>	Apr 22, 2021
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A213709 004</u>	Apr 22, 2021
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A213709 005</u>	Apr 22, 2021
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A213709 006</u>	Apr 22, 2021
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A213709 007</u>	Apr 22, 2021
AB	AUROLIFE PHARMA LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A202424 001</u>	Nov 27, 2013
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A202424 002</u>	Nov 27, 2013
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A202424 003</u>	Nov 27, 2013
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A202424 004</u>	Nov 27, 2013
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A202424 005</u>	Nov 27, 2013
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A202424 006</u>	Nov 27, 2013
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A202424 007</u>	Nov 27, 2013
AB	BARR	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040422 001</u>	Feb 11, 2002
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040422 005</u>	Mar 19, 2003
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040422 002</u>	Feb 11, 2002
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040422 006</u>	Mar 19, 2003
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040422 007</u>	Mar 19, 2003
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A040422 003</u>	Feb 11, 2002
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040422 004</u>	Feb 11, 2002
AB	ELITE LABS INC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A211352 001</u>	Dec 07, 2018
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A211352 002</u>	Dec 07, 2018
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A211352 003</u>	Dec 07, 2018
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A211352 004</u>	Dec 07, 2018
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A211352 005</u>	Dec 07, 2018
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A211352 006</u>	Dec 07, 2018
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A211352 007</u>	Dec 07, 2018
AB	EPIC PHARMA LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040444 001</u>	Jun 19, 2002
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040444 005</u>	Nov 03, 2014
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040444 002</u>	Jun 19, 2002
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040444 006</u>	Nov 03, 2014
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040444 007</u>	Nov 03, 2014
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A040444 003</u>	Jun 19, 2002
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040444 004</u>	Jun 19, 2002
AB	GRANULES	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A215771 001</u>	Dec 28, 2021
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A215771 002</u>	Dec 28, 2021
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A215771 003</u>	Dec 28, 2021
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A215771 004</u>	Dec 28, 2021
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A215771 005</u>	Dec 28, 2021
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A215771 006</u>	Dec 28, 2021
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A215771 007</u>	Dec 28, 2021
AB	LANNETT CO INC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A215565 001</u>	Jul 08, 2022
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A215565 002</u>	Jul 08, 2022
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A215565 003</u>	Jul 08, 2022
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A215565 004</u>	Jul 08, 2022
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A215565 005</u>	Jul 08, 2022
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A215565 006</u>	Jul 08, 2022
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A215565 007</u>	Jul 08, 2022
AB	NUVO PHARM	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A209799 001</u>	Dec 28, 2017
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A209799 002</u>	Dec 28, 2017
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A209799 003</u>	Dec 28, 2017
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A209799 004</u>	Dec 28, 2017
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A209799 005</u>	Dec 28, 2017
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A209799 006</u>	Dec 28, 2017
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A209799 007</u>	Dec 28, 2017
AB	RHODES PHARMS	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A213111 001</u>	Jan 13, 2021
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A213111 002</u>	Jan 13, 2021
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A213111 003</u>	Jan 13, 2021
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A213111 004</u>	Jan 13, 2021
AB		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A213111 005</u>	Jan 13, 2021
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A213111 006</u>	Jan 13, 2021
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A213111 007</u>	Jan 13, 2021
AB	SANDOZ	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040439 004</u>	Sep 27, 2002
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040439 001</u>	Jun 14, 2002
AB		<u>5MG;5MG;5MG;5MG</u>	<u>A040439 002</u>	Jun 14, 2002
AB		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040439 003</u>	Jun 14, 2002
AB	SPECGX LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040440 001</u>	Oct 07, 2003
AB		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040440 002</u>	Oct 07, 2003
AB		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040440 003</u>	Oct 07, 2003
AB		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040440 004</u>	Oct 07, 2003

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A040440 005	Oct 07, 2003
AB		5MG; 5MG; 5MG; 5MG	A040440 006	Oct 07, 2003
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A040440 007	Oct 07, 2003
AB	SUN PHARM INDUSTRIES	1.25MG; 1.25MG; 1.25MG; 1.25MG	A040480 001	Sep 09, 2003
AB		1.875MG; 1.875MG; 1.875MG; 1.875MG	A040480 002	Sep 09, 2003
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A040480 003	Sep 09, 2003
AB		3.125MG; 3.125MG; 3.125MG; 3.125MG	A040480 004	Sep 09, 2003
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A040480 005	Sep 09, 2003
AB		5MG; 5MG; 5MG; 5MG	A040480 006	Sep 09, 2003
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A040480 007	Sep 09, 2003
AB	USPHARMA WINDLAS	1.25MG; 1.25MG; 1.25MG; 1.25MG	A210293 001	Apr 03, 2020
AB		1.875MG; 1.875MG; 1.875MG; 1.875MG	A210293 002	Apr 03, 2020
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A210293 003	Apr 03, 2020
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A210293 004	Apr 03, 2020
AB		5MG; 5MG; 5MG; 5MG	A210293 005	Apr 03, 2020
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A210293 006	Apr 03, 2020
AB	ZYDUS PHARMS	1.25MG; 1.25MG; 1.25MG; 1.25MG	A207340 001	Oct 31, 2017
AB		1.875MG; 1.875MG; 1.875MG; 1.875MG	A207340 002	Oct 31, 2017
AB		2.5MG; 2.5MG; 2.5MG; 2.5MG	A207340 003	Oct 31, 2017
AB		3.125MG; 3.125MG; 3.125MG; 3.125MG	A207340 004	Oct 31, 2017
AB		3.75MG; 3.75MG; 3.75MG; 3.75MG	A207340 005	Oct 31, 2017
AB		5MG; 5MG; 5MG; 5MG	A207340 006	Oct 31, 2017
AB		7.5MG; 7.5MG; 7.5MG; 7.5MG	A207340 007	Oct 31, 2017

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

AA	ALKEM LABS LTD	5MG	A213720 001	Oct 27, 2020
AA		10MG	A213720 002	Oct 27, 2020
AA	AMNEAL PHARMS	5MG	A211139 001	Sep 26, 2018
AA		10MG	A211139 002	Sep 26, 2018
AA	AUROLIFE PHARMA LLC	5MG	A211639 001	Apr 17, 2019
AA		10MG	A211639 002	Apr 17, 2019
AA	BIONPHARMA	5MG	A212919 001	Nov 22, 2019
AA		10MG	A212919 002	Nov 22, 2019
AA	CEROVENE INC	5MG	A212582 001	Feb 04, 2020
AA		10MG	A212582 002	Feb 04, 2020
AA	EPIC PHARMA LLC	5MG	A213980 001	Oct 27, 2020
AA		10MG	A213980 002	Oct 27, 2020
AA	GRANULES	5MG	A212619 001	Aug 05, 2019
AA		10MG	A212619 002	Aug 05, 2019
AA	PRINSTON INC	5MG	A211861 001	Mar 11, 2020
AA		10MG	A211861 002	Mar 11, 2020
AA	RHODES PHARMS	5MG	A213852 001	Sep 07, 2021
AA		10MG	A213852 002	Sep 07, 2021
AA	RISE PHARMA	5MG	A212901 001	May 22, 2020
AA		10MG	A212901 002	May 22, 2020
AA	SPECGX LLC	5MG	A213583 001	Jan 22, 2021
AA		10MG	A213583 002	Jan 22, 2021
AA	SUN PHARM INDS INC	5MG	A214574 001	Jan 27, 2021
AA		10MG	A214574 002	Jan 27, 2021
EVEKEO				
AA	ARBOR PHARMS LLC	5MG	A200166 001	Aug 09, 2012
AA	!	10MG	A200166 002	Aug 09, 2012

TABLET, ORALLY DISINTEGRATING; ORAL

EVEKEO ODT

+	AZURITY	5MG	N209905 001	Jan 30, 2019
+		10MG	N209905 002	Jan 30, 2019
+		15MG	N209905 003	Jan 30, 2019
+	!	20MG	N209905 004	Jan 30, 2019

AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE

SUSPENSION, EXTENDED RELEASE; ORAL

DYANAVEL XR

+	TRIS PHARMA INC	2MG/ML; EQ 0.5MG BASE/ML	N208147 001	Oct 19, 2015
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TABLET, EXTENDED RELEASE; ORAL

DYANAVEL XR 10

+	TRIS PHARMA INC	8MG; EQ 2MG BASE	N210526 002	Nov 04, 2021
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PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTRAMPHETAMINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

DYANAVEL XR 15

+ TRIS PHARMA INC 12MG;EQ 3MG BASE N210526 003 Nov 04, 2021

DYANAVEL XR 20

+! TRIS PHARMA INC 16MG;EQ 4MG BASE N210526 004 Nov 04, 2021

DYANAVEL XR 5

+ TRIS PHARMA INC 4MG;EQ 1MG BASE N210526 001 Nov 04, 2021

AMPHOTERICIN B

INJECTABLE;INJECTION

AMPHOTERICIN B

! XGEN PHARMS 50MG/VIAL A063206 001 Apr 29, 1992

INJECTABLE, LIPID COMPLEX;INJECTION

ABELCET

+! LEADIANT BIOSCI INC 5MG/ML N050724 001 Nov 20, 1995

INJECTABLE, LIPOSOMAL;INJECTION

AMBISOME**AB** +! ASTELLAS **50MG/VIAL** **N050740 001** Aug 11, 1997AMPHOTERICIN B**AB** EUGIA PHARMA **50MG/VIAL** **A214010 001** Nov 17, 2022**AB** SPIL **50MG/VIAL** **A212514 001** Dec 14, 2021AMPICILLIN SODIUM

INJECTABLE;INJECTION

AMPICILLIN SODIUM**AP** ACS DOBFAR SPA **EQ 10GM BASE/VIAL** **A090889 001** Apr 03, 2013**AP** ANTIBIOTICE **EQ 250MG BASE/VIAL** **A090354 001** Dec 28, 2009**AP** **EQ 500MG BASE/VIAL** **A090354 002** Dec 28, 2009**AP** **EQ 1GM BASE/VIAL** **A090354 003** Dec 28, 2009**AP** **EQ 2GM BASE/VIAL** **A090354 004** Dec 28, 2009**AP** EUGIA PHARMA **EQ 250MG BASE/VIAL** **A065499 002** Aug 17, 2010

SPECLTS

AP **EQ 500MG BASE/VIAL** **A065499 003** Aug 17, 2010**AP** **EQ 1GM BASE/VIAL** **A065499 004** Aug 17, 2010**AP** **EQ 2GM BASE/VIAL** **A065499 005** Aug 17, 2010**AP** **EQ 10GM BASE/VIAL** **A065493 001** Aug 17, 2010**AP** HQ SPECLT PHARMA **EQ 250MG BASE/VIAL** **A062772 006** Apr 15, 1993**AP** **EQ 500MG BASE/VIAL** **A062772 007** Apr 15, 1993**AP** **EQ 1GM BASE/VIAL** **A062772 001** Apr 15, 1993**AP** **EQ 2GM BASE/VIAL** **A062772 003** Apr 15, 1993**AP** **EQ 10GM BASE/VIAL** **A063142 001** Apr 15, 1993**AP** ISTITUTO BIO ITA **EQ 10GM BASE/VIAL** **A201404 001** Dec 20, 2013

SPA

AP **EQ 250MG BASE/VIAL** **A062719 001** May 12, 1987**AP** **EQ 500MG BASE/VIAL** **A062719 003** May 12, 1987**AP** **EQ 1GM BASE/VIAL** **A062719 002** May 12, 1987**AP** **EQ 2GM BASE/VIAL** **A062797 002** Jul 12, 1993**AP** SAGENT PHARMS INC **EQ 125MG BASE/VIAL** **A090583 001** Nov 27, 2015**AP** **EQ 250MG BASE/VIAL** **A090583 002** Nov 27, 2015**AP** **EQ 500MG BASE/VIAL** **A090583 003** Nov 27, 2015**AP** **EQ 1GM BASE/VIAL** **A090583 004** Nov 27, 2015**AP** **EQ 2GM BASE/VIAL** **A090583 005** Nov 27, 2015**AP** **EQ 10GM BASE/VIAL** **A090581 001** Oct 20, 2015**AP** ! SANDOZ **EQ 125MG BASE/VIAL** **A061395 001****AP** ! **EQ 250MG BASE/VIAL** **A061395 002****AP** ! **EQ 500MG BASE/VIAL** **A061395 003****AP** ! **EQ 1GM BASE/VIAL** **A061395 004****AP** ! **EQ 2GM BASE/VIAL** **A061395 005****AP** ! **EQ 10GM BASE/VIAL** **A061395 006****AP** STERISCIENCE **EQ 1GM BASE/VIAL** **A201025 003** Apr 09, 2014**AP** **EQ 2GM BASE/VIAL** **A201025 004** Apr 09, 2014**AP** **EQ 10GM BASE/VIAL** **A202198 001** Apr 07, 2014

POWDER; INTRAVENOUS

AMPICILLIN SODIUM**AP** ! SANDOZ **EQ 1GM BASE/VIAL** **A062738 001** Feb 19, 1987**AP** ! **EQ 2GM BASE/VIAL** **A062738 002** Feb 19, 1987

PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

<u>AP</u>	ACS DOBFAR	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065406 001</u>	Dec 22, 2009
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065406 002</u>	Dec 22, 2009
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065403 001</u>	Dec 23, 2009
<u>AP</u>	ANTIBIOTICE	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201406 001</u>	Dec 07, 2015
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201406 002</u>	Dec 07, 2015
<u>AP</u>	ASTRAL	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090579 001</u>	Jan 08, 2016
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090579 002</u>	Jan 08, 2016
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090578 001</u>	Jan 11, 2016
<u>AP</u>	EUGIA PHARMA SPECLTS	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090349 001</u>	Sep 20, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090349 002</u>	Sep 20, 2010
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090339 001</u>	Sep 20, 2010
<u>AP</u>	HIKMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065074 001</u>	Mar 19, 2002
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065074 002</u>	Mar 19, 2002
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065076 001</u>	Mar 19, 2002
<u>AP</u>	HQ SPECLT PHARMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065176 001</u>	Nov 30, 2005
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065176 002</u>	Nov 30, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065188 001</u>	Nov 25, 2005
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065222 001</u>	Nov 29, 2005
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065222 002</u>	Nov 29, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065314 001</u>	Nov 27, 2006
<u>AP</u>	SANDOZ	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065241 001</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065310 001</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065241 002</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065310 002</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065240 001</u>	Jul 25, 2006
<u>AP</u>	STERISCIENCE	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201024 001</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201024 002</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A202197 001</u>	Apr 07, 2014
<u>UNASYN</u>				
<u>AP</u>	+! PFIZER	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N050608 002</u>	Dec 31, 1986
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N050608 001</u>	Dec 31, 1986
<u>AP</u>	+!	<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N050608 005</u>	Dec 10, 1993

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A216554 002</u>	Oct 31, 2023
<u>AB</u>	! SANDOZ	<u>EQ 500MG BASE</u>	<u>A064082 002</u>	Aug 29, 1995
	AUROBINDO PHARMA	EQ 250MG BASE	A216554 001	Oct 31, 2023

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

<u>AB</u>	TAKEDA PHARMS USA	<u>EQ 0.5MG BASE</u>	<u>N020333 001</u>	Mar 14, 1997
<u>ANAGRELIDE HYDROCHLORIDE</u>				
<u>AB</u>	IMPAX LABS	<u>EQ 0.5MG BASE</u>	<u>A076910 001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076910 002</u>	Apr 18, 2005
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 0.5MG BASE</u>	<u>A076468 001</u>	Apr 18, 2005
<u>AB</u>	!	<u>EQ 1MG BASE</u>	<u>A076468 002</u>	Apr 18, 2005
<u>AB</u>	TORRENT	<u>EQ 0.5MG BASE</u>	<u>A209151 001</u>	Jun 30, 2017
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A209151 002</u>	Jun 30, 2017

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>1MG</u>	<u>A090568 001</u>	Jun 28, 2010
<u>AB</u>	BELJING YILING	<u>1MG</u>	<u>A206037 001</u>	Nov 09, 2018
<u>AB</u>	CIPLA	<u>1MG</u>	<u>A091164 001</u>	Jun 28, 2010
<u>AB</u>	EUGIA PHARMA	<u>1MG</u>	<u>A212434 001</u>	Jul 24, 2020
<u>AB</u>	KENTON	<u>1MG</u>	<u>A078944 001</u>	Jun 28, 2010
<u>AB</u>	NATCO PHARMA LTD	<u>1MG</u>	<u>A079220 001</u>	Jun 28, 2010
<u>AB</u>	TEVA PHARMS	<u>1MG</u>	<u>A078058 001</u>	Jun 28, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>1MG</u>	<u>A078921 001</u>	Jun 28, 2010

ARIMIDEX

<u>AB</u>	+! ANI PHARMS	<u>1MG</u>	<u>N020541 001</u>	Dec 27, 1995
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PRESCRIPTION DRUG PRODUCT LIST

ANGIOTENSIN II ACETATE

SOLUTION; INTRAVENOUS

GIAPREZA

+! LA JOLLA PHARMA

EQ 0.5MG BASE/ML (EQ 0.5MG BASE/ML)

N209360 003 Dec 23, 2021

+!

EQ 2.5MG BASE/ML (EQ 2.5MG BASE/ML)

N209360 001 Dec 21, 2017

ANIDULAFUNGIN

POWDER; INTRAVENOUS

ERAXIS

+! VICURON HOLDINGS

50MG/VIAL

N021632 001 Feb 17, 2006

+!

100MG/VIAL

N021632 002 Nov 14, 2006

APALUTAMIDE

TABLET; ORAL

ERLEADA

+ JANSSEN BIOTECH

60MG

N210951 001 Feb 14, 2018

+!

240MG

N210951 002 Feb 17, 2023

APIXABAN

TABLET; ORAL

APIXABANAB ACCORD HLTHCARE2.5MGA210180 001 Jul 28, 2020AB5MGA210180 002 Jul 28, 2020AB HETERO LABS LTD V2.5MGA210066 001 Nov 21, 2023AB5MGA210066 002 Nov 21, 2023AB INDOCO2.5MGA209898 001 Sep 11, 2020AB5MGA209898 002 Sep 11, 2020ELIQUISAB + BRISTOL MYERS2.5MGN202155 001 Dec 28, 2012

SQUIBB

AB +!5MGN202155 002 Dec 28, 2012APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYNAP +! MDD US30MG/3ML (10MG/ML)N021264 002 Apr 20, 2004APOMORPHINE HYDROCHLORIDEAP SAGE CHEMS30MG/3ML (10MG/ML)A212025 001 Feb 23, 2022APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

APRACLONIDINE HYDROCHLORIDEAT RISINGEQ 0.5% BASEA077764 001 Mar 12, 2009IOPIDINEAT +! HARROW EYEEQ 0.5% BASEN020258 001 Jul 30, 1993

+!

EQ 1% BASE

N019779 001 Dec 31, 1987

APREMILAST

TABLET; ORAL

APREMILASTAB ALKEM LABS LTD10MGA211761 001 Sep 21, 2021AB20MGA211761 002 Sep 21, 2021AB30MGA211761 003 Sep 21, 2021AB ANNORA10MGA211878 001 Jul 26, 2023AB20MGA211878 002 Jul 26, 2023AB30MGA211878 003 Jul 26, 2023AB GLENMARK PHARMS LTD10MGA211674 001 Oct 16, 2023AB20MGA211674 002 Oct 16, 2023AB30MGA211674 003 Oct 16, 2023AB SHILPA10MGA211774 001 Apr 07, 2023AB20MGA211774 002 Apr 07, 2023AB30MGA211774 003 Apr 07, 2023OTEZLAAB + AMGEN INC10MGN205437 001 Mar 21, 2014AB +20MGN205437 002 Mar 21, 2014AB +!30MGN205437 003 Mar 21, 2014APREPITANT

CAPSULE; ORAL

APREPITANTAB GLENMARK PHARMS SA40MGA207777 001 Oct 12, 2017AB80MGA207777 002 Oct 12, 2017AB125MGA207777 003 Oct 12, 2017AB SANDOZ40MGA090999 001 Sep 24, 2012AB80MGA090999 002 Sep 24, 2012AB125MGA090999 003 Sep 24, 2012

PRESCRIPTION DRUG PRODUCT LIST

APREPITANT

CAPSULE; ORAL

APREPITANT

<u>AB</u>	TORRENT	<u>40MG</u>	<u>A211835 001</u>	Oct 21, 2020
<u>AB</u>		<u>80MG</u>	<u>A211835 002</u>	Oct 21, 2020
<u>AB</u>		<u>125MG</u>	<u>A211835 003</u>	Oct 21, 2020

EMEND

<u>AB</u>	+ MERCK	<u>80MG</u>	<u>N021549 001</u>	Mar 26, 2003
<u>AB</u>	+!	<u>125MG</u>	<u>N021549 002</u>	Mar 26, 2003

EMULSION; INTRAVENOUS

APONVIE

+!	HERON THERAPS INC	32MG/4.4ML (7.2MG/ML)	N216457 001	Sep 16, 2022
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CINVANTI

+!	HERON THERAPS INC	130MG/18ML (7.2MG/ML)	N209296 001	Nov 09, 2017
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FOR SUSPENSION; ORAL

EMEND

+!	MSD MERCK CO	125MG/KIT	N207865 001	Dec 17, 2015
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ARFORMOTEROL TARTRATE

SOLUTION; INHALATION

ARFORMOTEROL TARTRATE

<u>AN</u>	ALEMBIC	<u>EQ 0.015MG BASE/2ML</u>	<u>A214779 001</u>	May 10, 2022
<u>AN</u>	CIPLA	<u>EQ 0.015MG BASE/2ML</u>	<u>A207306 001</u>	Jun 22, 2021
<u>AN</u>	LUPIN	<u>EQ 0.015MG BASE/2ML</u>	<u>A213068 001</u>	Feb 07, 2022
<u>AN</u>	MANKIND PHARMA	<u>EQ 0.015MG BASE/2ML</u>	<u>A216128 001</u>	Nov 15, 2022
<u>AN</u>	RITEDOSE CORP	<u>EQ 0.015MG BASE/2ML</u>	<u>A214736 001</u>	Mar 02, 2022
<u>AN</u>	SLATE RUN PHARMA	<u>EQ 0.015MG BASE/2ML</u>	<u>A213762 001</u>	Jun 22, 2021
<u>AN</u>	SLAYBACK PHARMA LLC	<u>EQ 0.015MG BASE/2ML</u>	<u>A216815 001</u>	Nov 25, 2022
<u>AN</u>	SUN PHARM	<u>EQ 0.015MG BASE/2ML</u>	<u>A215385 001</u>	May 26, 2022
<u>AN</u>	TEVA PHARMS USA	<u>EQ 0.015MG BASE/2ML</u>	<u>A200293 001</u>	Nov 09, 2021

BROVANA

<u>AN</u>	+! LUPIN	<u>EQ 0.015MG BASE/2ML</u>	<u>N021912 001</u>	Oct 06, 2006
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ARGATROBAN

INJECTABLE; INJECTION

ARGATROBAN

<u>AP</u>	AMNEAL PHARMS CO	<u>250MG/2.5ML (100MG/ML)</u>	<u>A206698 001</u>	Jan 26, 2018
<u>AP</u>	CAPLIN	<u>50MG/50ML (1MG/ML)</u>	<u>A214235 001</u>	Jan 21, 2021
<u>AP</u>	FRESENIUS KABI USA	<u>250MG/2.5ML (100MG/ML)</u>	<u>N201811 001</u>	Mar 23, 2015
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/50ML (1MG/ML)</u>	<u>A217848 001</u>	Jul 31, 2023
<u>AP</u>	+! HIKMA PHARM CO LTD	<u>50MG/50ML (1MG/ML)</u>	<u>N203049 002</u>	Sep 30, 2016
<u>AP</u>	+!	<u>250MG/2.5ML (100MG/ML)</u>	<u>N203049 001</u>	Jan 05, 2012
<u>AP</u>	! HOSPIRA	<u>250MG/2.5ML (100MG/ML)</u>	<u>A204120 001</u>	Sep 21, 2016
<u>AP</u>	MYLAN INSTITUTIONAL	<u>250MG/2.5ML (100MG/ML)</u>	<u>A202626 001</u>	Jun 30, 2014
<u>AP</u>	PAR STERILE PRODUCTS	<u>250MG/2.5ML (100MG/ML)</u>	<u>A091665 001</u>	Jun 30, 2014

INJECTABLE; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

<u>AP</u>	GLAND PHARMA LTD	<u>125MG/125ML (1MG/ML)</u>	<u>A205570 001</u>	May 22, 2017
<u>AP</u>	+! SANDOZ	<u>125MG/125ML (1MG/ML)</u>	<u>N022485 001</u>	May 09, 2011

SOLUTION; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

+!	ACCORD HLTHCARE	50MG/50ML (1MG/ML)	N212035 001	Jun 07, 2021
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	EUGIA PHARMA	50MG/50ML (1MG/ML)	N209552 001	Nov 27, 2018
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SPECLTS

ARGININE HYDROCHLORIDE

INJECTABLE; INJECTION

R-GENE 10

+!	PHARMACIA AND UPJOHN	10GM/100ML	N016931 001	
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ARIPIPRAZOLE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

ABILIFY MAINTENA KIT

+	OTSUKA PHARM CO LTD	300MG/VIAL	N202971 001	Feb 28, 2013
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+		300MG	N202971 003	Sep 29, 2014
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+!		400MG/VIAL	N202971 002	Feb 28, 2013
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+		400MG	N202971 004	Sep 29, 2014
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SOLUTION; ORAL

ARIPIPRAZOLE

<u>AA</u>	! AMNEAL PHARMS	<u>1MG/ML</u>	<u>A203906 001</u>	Aug 14, 2015
<u>AA</u>	APOTEX	<u>1MG/ML</u>	<u>A204094 001</u>	Sep 30, 2015
<u>AA</u>	AUROBINDO PHARMA LTD	<u>1MG/ML</u>	<u>A210479 001</u>	Jan 29, 2019

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

SOLUTION;ORAL

ARIPIPIRAZOLE

<u>AA</u>	CHARTWELL RX	<u>1MG/ML</u>	<u>A215595</u>	<u>001</u>	Oct 25, 2022
<u>AA</u>	HETERO LABS LTD III	<u>1MG/ML</u>	<u>A216150</u>	<u>001</u>	Nov 01, 2023
<u>AA</u>	LANNETT CO INC	<u>1MG/ML</u>	<u>A204171</u>	<u>001</u>	Aug 14, 2015
<u>AA</u>	RUBICON	<u>1MG/ML</u>	<u>A216351</u>	<u>001</u>	Oct 27, 2022
<u>AA</u>	VISTAPHARM	<u>1MG/ML</u>	<u>A212870</u>	<u>001</u>	Dec 26, 2019

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ABILIFY ASIMTUFII

+	OTSUKA	720MG/2.4ML (300MG/ML)	N217006	001	Apr 27, 2023
+	!	960MG/3.2ML (300MG/ML)	N217006	002	Apr 27, 2023

TABLET;ORAL

ABILIFY

<u>AB</u>	+	OTSUKA	<u>2MG</u>	<u>N021436</u>	<u>006</u>	Nov 15, 2002
<u>AB</u>	+		<u>5MG</u>	<u>N021436</u>	<u>005</u>	Nov 15, 2002
<u>AB</u>	+	!	<u>10MG</u>	<u>N021436</u>	<u>001</u>	Nov 15, 2002
<u>AB</u>	+		<u>15MG</u>	<u>N021436</u>	<u>002</u>	Nov 15, 2002
<u>AB</u>	+		<u>20MG</u>	<u>N021436</u>	<u>003</u>	Nov 15, 2002
<u>AB</u>	+		<u>30MG</u>	<u>N021436</u>	<u>004</u>	Nov 15, 2002

ARIPIPIRAZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>2MG</u>	<u>A206251</u>	<u>001</u>	Dec 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A206251</u>	<u>002</u>	Dec 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A206251</u>	<u>003</u>	Dec 07, 2016
<u>AB</u>		<u>15MG</u>	<u>A206251</u>	<u>004</u>	Dec 07, 2016
<u>AB</u>		<u>20MG</u>	<u>A206251</u>	<u>005</u>	Dec 07, 2016
<u>AB</u>		<u>30MG</u>	<u>A206251</u>	<u>006</u>	Dec 07, 2016
<u>AB</u>	AJANTA PHARMA LTD	<u>2MG</u>	<u>A206174</u>	<u>001</u>	Sep 12, 2016
<u>AB</u>		<u>5MG</u>	<u>A206174</u>	<u>002</u>	Sep 12, 2016
<u>AB</u>		<u>10MG</u>	<u>A206174</u>	<u>003</u>	Sep 12, 2016
<u>AB</u>		<u>15MG</u>	<u>A206174</u>	<u>004</u>	Sep 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A206174</u>	<u>005</u>	Sep 12, 2016
<u>AB</u>		<u>30MG</u>	<u>A206174</u>	<u>006</u>	Sep 12, 2016
<u>AB</u>	ALEMBIC	<u>2MG</u>	<u>A202101</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A202101</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A202101</u>	<u>003</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A202101</u>	<u>004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A202101</u>	<u>005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A202101</u>	<u>006</u>	Apr 28, 2015
<u>AB</u>	ALKEM LABS LTD	<u>2MG</u>	<u>A207105</u>	<u>001</u>	Feb 21, 2019
<u>AB</u>		<u>5MG</u>	<u>A207105</u>	<u>002</u>	Feb 21, 2019
<u>AB</u>		<u>10MG</u>	<u>A207105</u>	<u>003</u>	Feb 21, 2019
<u>AB</u>		<u>15MG</u>	<u>A207105</u>	<u>004</u>	Feb 21, 2019
<u>AB</u>		<u>20MG</u>	<u>A207105</u>	<u>005</u>	Feb 21, 2019
<u>AB</u>		<u>30MG</u>	<u>A207105</u>	<u>006</u>	Feb 21, 2019
<u>AB</u>	AMNEAL PHARMS	<u>2MG</u>	<u>A204838</u>	<u>001</u>	Jun 17, 2016
<u>AB</u>		<u>5MG</u>	<u>A204838</u>	<u>002</u>	Jun 17, 2016
<u>AB</u>		<u>10MG</u>	<u>A204838</u>	<u>003</u>	Jun 17, 2016
<u>AB</u>		<u>15MG</u>	<u>A204838</u>	<u>004</u>	Jun 17, 2016
<u>AB</u>		<u>20MG</u>	<u>A204838</u>	<u>005</u>	Jun 17, 2016
<u>AB</u>		<u>30MG</u>	<u>A204838</u>	<u>006</u>	Jun 17, 2016
<u>AB</u>	APOTEX	<u>2MG</u>	<u>A078583</u>	<u>001</u>	Jul 24, 2015
<u>AB</u>		<u>5MG</u>	<u>A078583</u>	<u>002</u>	Jul 24, 2015
<u>AB</u>		<u>10MG</u>	<u>A078583</u>	<u>003</u>	Jul 24, 2015
<u>AB</u>		<u>15MG</u>	<u>A078583</u>	<u>004</u>	Jul 24, 2015
<u>AB</u>		<u>20MG</u>	<u>A078583</u>	<u>005</u>	Jul 24, 2015
<u>AB</u>		<u>30MG</u>	<u>A078583</u>	<u>006</u>	Jul 24, 2015
<u>AB</u>	AUROBINDO PHARMA	<u>2MG</u>	<u>A203908</u>	<u>001</u>	Oct 08, 2015
<u>AB</u>		<u>5MG</u>	<u>A203908</u>	<u>002</u>	Oct 08, 2015
<u>AB</u>		<u>10MG</u>	<u>A203908</u>	<u>003</u>	Oct 08, 2015
<u>AB</u>		<u>15MG</u>	<u>A203908</u>	<u>004</u>	Oct 08, 2015
<u>AB</u>		<u>20MG</u>	<u>A203908</u>	<u>005</u>	Oct 08, 2015
<u>AB</u>		<u>30MG</u>	<u>A203908</u>	<u>006</u>	Oct 08, 2015
<u>AB</u>	BRECKENRIDGE	<u>2MG</u>	<u>A091279</u>	<u>001</u>	Jan 09, 2017
<u>AB</u>		<u>5MG</u>	<u>A091279</u>	<u>002</u>	Jan 09, 2017
<u>AB</u>		<u>10MG</u>	<u>A091279</u>	<u>003</u>	Jan 09, 2017
<u>AB</u>		<u>15MG</u>	<u>A091279</u>	<u>004</u>	Jan 09, 2017
<u>AB</u>		<u>20MG</u>	<u>A091279</u>	<u>005</u>	Jan 09, 2017
<u>AB</u>		<u>30MG</u>	<u>A091279</u>	<u>006</u>	Jan 09, 2017
<u>AB</u>	HETERO LABS LTD V	<u>2MG</u>	<u>A205064</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A205064</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A205064</u>	<u>003</u>	Apr 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

TABLET; ORAL

ARIPIPIRAZOLE

<u>AB</u>		<u>15MG</u>	<u>A205064 004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A205064 005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A205064 006</u>	Apr 28, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>2MG</u>	<u>A204111 001</u>	Oct 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A204111 002</u>	Oct 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A204111 003</u>	Oct 07, 2016
<u>AB</u>		<u>15MG</u>	<u>A204111 004</u>	Oct 07, 2016
<u>AB</u>		<u>20MG</u>	<u>A204111 005</u>	Oct 07, 2016
<u>AB</u>		<u>30MG</u>	<u>A204111 006</u>	Oct 07, 2016
<u>AB</u>	ORBION PHARMS	<u>2MG</u>	<u>A202683 001</u>	May 23, 2017
<u>AB</u>		<u>5MG</u>	<u>A202683 002</u>	May 23, 2017
<u>AB</u>		<u>10MG</u>	<u>A202683 003</u>	May 23, 2017
<u>AB</u>		<u>15MG</u>	<u>A202683 004</u>	May 23, 2017
<u>AB</u>		<u>20MG</u>	<u>A202683 005</u>	May 23, 2017
<u>AB</u>		<u>30MG</u>	<u>A202683 006</u>	May 23, 2017
<u>AB</u>	PRINSTON INC	<u>2MG</u>	<u>A205363 001</u>	Dec 04, 2017
<u>AB</u>		<u>5MG</u>	<u>A205363 002</u>	Dec 04, 2017
<u>AB</u>		<u>10MG</u>	<u>A205363 003</u>	Dec 04, 2017
<u>AB</u>		<u>15MG</u>	<u>A205363 004</u>	Dec 04, 2017
<u>AB</u>		<u>20MG</u>	<u>A205363 005</u>	Dec 04, 2017
<u>AB</u>		<u>30MG</u>	<u>A205363 006</u>	Dec 04, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>2MG</u>	<u>A206383 001</u>	Sep 29, 2016
<u>AB</u>		<u>5MG</u>	<u>A206383 002</u>	Sep 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A206383 003</u>	Sep 29, 2016
<u>AB</u>		<u>15MG</u>	<u>A206383 004</u>	Sep 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A206383 005</u>	Sep 29, 2016
<u>AB</u>		<u>30MG</u>	<u>A206383 006</u>	Sep 29, 2016
<u>AB</u>	SUNSHINE	<u>2MG</u>	<u>A213037 001</u>	Oct 02, 2023
<u>AB</u>		<u>5MG</u>	<u>A213037 002</u>	Oct 02, 2023
<u>AB</u>		<u>10MG</u>	<u>A213037 003</u>	Oct 02, 2023
<u>AB</u>	TORRENT	<u>2MG</u>	<u>A201519 001</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A201519 003</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A201519 002</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A201519 004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A201519 005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A201519 006</u>	Apr 28, 2015
<u>AB</u>	UNICHEM	<u>2MG</u>	<u>A203025 001</u>	Dec 01, 2021
<u>AB</u>		<u>5MG</u>	<u>A203025 002</u>	Dec 01, 2021
<u>AB</u>		<u>10MG</u>	<u>A203025 003</u>	Dec 01, 2021
<u>AB</u>		<u>15MG</u>	<u>A203025 004</u>	Dec 01, 2021
<u>AB</u>		<u>20MG</u>	<u>A203025 005</u>	Dec 01, 2021
<u>AB</u>		<u>30MG</u>	<u>A203025 006</u>	Dec 01, 2021
	ABILIFY MYCITE KIT			
	+ OTSUKA	2MG	N207202 001	Nov 13, 2017
	+!	5MG	N207202 002	Nov 13, 2017
	+	10MG	N207202 003	Nov 13, 2017
	+	15MG	N207202 004	Nov 13, 2017
	+	20MG	N207202 005	Nov 13, 2017
	+	30MG	N207202 006	Nov 13, 2017

TABLET, ORALLY DISINTEGRATING; ORAL

ARIPIPIRAZOLE

<u>AB</u>	! ALEMBIC	<u>10MG</u>	<u>A202102 001</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A202102 002</u>	Apr 28, 2015
<u>AB</u>	ORBION PHARMS	<u>10MG</u>	<u>A202547 001</u>	Dec 11, 2017
<u>AB</u>		<u>15MG</u>	<u>A202547 002</u>	Dec 11, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>10MG</u>	<u>A207240 001</u>	Apr 18, 2018
<u>AB</u>		<u>15MG</u>	<u>A207240 002</u>	Apr 18, 2018
<u>AB</u>	SQUARE PHARMS	<u>10MG</u>	<u>A090165 001</u>	Aug 28, 2018
<u>AB</u>		<u>15MG</u>	<u>A090165 002</u>	Aug 28, 2018
		20MG	A090165 003	Aug 28, 2018
		30MG	A090165 004	Aug 28, 2018

ARIPIPIRAZOLE LAUROXIL

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

ARISTADA

	+ ALKERMES INC	441MG/1.6ML (275.63MG/ML)	N207533 001	Oct 05, 2015
	+	662MG/2.4ML (275.83MG/ML)	N207533 002	Oct 05, 2015
	+!	882MG/3.2ML (275.63MG/ML)	N207533 003	Oct 05, 2015
	+	1064MG/3.9ML (272.82MG/ML)	N207533 004	Jun 05, 2017

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE LAUROXIL

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

ARISTADA INITIO KIT

+ ALKERMES INC

675MG/2.4ML

N209830 001 Jun 29, 2018

ARMODAFINIL

TABLET; ORAL

ARMODAFINIL

<u>AB</u>	AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A206069 001</u>	Mar 06, 2018
<u>AB</u>		<u>150MG</u>	<u>A206069 002</u>	Mar 06, 2018
<u>AB</u>		<u>200MG</u>	<u>A206069 004</u>	Dec 07, 2018
<u>AB</u>		<u>250MG</u>	<u>A206069 003</u>	Mar 06, 2018
<u>AB</u>	LUPIN LTD	<u>50MG</u>	<u>A200751 001</u>	Nov 28, 2016
<u>AB</u>		<u>150MG</u>	<u>A200751 003</u>	Nov 28, 2016
<u>AB</u>		<u>200MG</u>	<u>A200751 004</u>	Nov 28, 2016
<u>AB</u>		<u>250MG</u>	<u>A200751 005</u>	Nov 28, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>50MG</u>	<u>A200043 001</u>	Jun 01, 2012
<u>AB</u>		<u>100MG</u>	<u>A200043 004</u>	May 09, 2019
<u>AB</u>		<u>150MG</u>	<u>A200043 002</u>	Jun 01, 2012
<u>AB</u>		<u>200MG</u>	<u>A200043 005</u>	May 09, 2019
<u>AB</u>		<u>250MG</u>	<u>A200043 003</u>	Jun 01, 2012
<u>AB</u>	NATCO PHARMA LTD	<u>50MG</u>	<u>A202768 001</u>	Nov 28, 2016
<u>AB</u>		<u>100MG</u>	<u>A202768 004</u>	Sep 28, 2017
<u>AB</u>		<u>150MG</u>	<u>A202768 002</u>	Nov 28, 2016
<u>AB</u>		<u>200MG</u>	<u>A202768 005</u>	Sep 28, 2017
<u>AB</u>		<u>250MG</u>	<u>A202768 003</u>	Nov 28, 2016
<u>NUVIGIL</u>				
<u>AB</u>	+ CEPHALON	<u>50MG</u>	<u>N021875 001</u>	Jun 15, 2007
<u>AB</u>	+	<u>150MG</u>	<u>N021875 003</u>	Jun 15, 2007
<u>AB</u>	+	<u>200MG</u>	<u>N021875 005</u>	Mar 26, 2009
<u>AB</u>	+!	<u>250MG</u>	<u>N021875 004</u>	Jun 15, 2007

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

ARSENIC TRIOXIDE

<u>AP</u>	AMNEAL	<u>1MG/ML</u>	<u>A210739 001</u>	Jan 25, 2021
<u>AP</u>		<u>2MG/ML</u>	<u>A210739 002</u>	Aug 19, 2021
<u>AP</u>	AMRING PHARMS	<u>1MG/ML</u>	<u>A210802 001</u>	Nov 13, 2018
<u>AP</u>	EUGIA PHARMA	<u>2MG/ML</u>	<u>A214011 001</u>	Oct 15, 2021
<u>AP</u>	! FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A208231 001</u>	Aug 31, 2018
<u>AP</u>	GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A215059 002</u>	Apr 22, 2022
<u>AP</u>		<u>2MG/ML</u>	<u>A215059 001</u>	Oct 07, 2021
<u>AP</u>	INGENUS PHARMS LLC	<u>1MG/ML</u>	<u>A209315 001</u>	Nov 15, 2018
<u>AP</u>	NEXUS	<u>1MG/ML</u>	<u>A209780 001</u>	Nov 15, 2018
<u>AP</u>	ORBICULAR	<u>1MG/ML</u>	<u>A217413 001</u>	Apr 20, 2023
<u>AP</u>		<u>2MG/ML</u>	<u>A217413 002</u>	Apr 20, 2023
<u>AP</u>	PENN LIFE	<u>1MG/ML</u>	<u>A209873 001</u>	May 06, 2019
<u>AP</u>	SANDOZ	<u>2MG/ML</u>	<u>A215359 001</u>	Dec 02, 2021
<u>AP</u>	ZYDUS PHARMS	<u>1MG/ML</u>	<u>A206228 001</u>	Nov 13, 2018
<u>AP</u>		<u>2MG/ML</u>	<u>A206228 002</u>	Aug 30, 2019
<u>TRISENOX</u>				
<u>AP</u>	+! CEPHALON	<u>2MG/ML</u>	<u>N021248 002</u>	Oct 13, 2017

ARTEMETHER; LUMEFANTRINE

TABLET; ORAL

COARTEM

+! NOVARTIS

20MG; 120MG

N022268 001 Apr 07, 2009

ARTESUNATE

POWDER; INTRAVENOUS

ARTESUNATE

+! AMIVAS

110MG/VIAL

N213036 001 May 26, 2020

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ORABLOC

+ PIERREL

4%; EQ 0.009MG BASE/1.8ML (EQ 0.005MG BASE/ML)

N022466 001 Feb 26, 2010

+!

4%; EQ 0.018MG BASE/1.8ML (EQ 0.01MG BASE/ML)

N022466 002 Feb 26, 2010

SEPTOCAINE

+! DEPROCO

4%; EQ 0.0085MG BASE/1.7ML (4%; EQ 0.005MG BASE/ML)

N020971 002 Mar 30, 2006

+!

4%; EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)

N020971 001 Apr 03, 2000

PRESCRIPTION DRUG PRODUCT LIST

ASCIMINIB HYDROCHLORIDE

TABLET; ORAL

SCEMBLIX

+ NOVARTIS

EQ 20MG BASE

N215358 001 Oct 29, 2021

+!

EQ 40MG BASE

N215358 002 Oct 29, 2021

ASCORBIC ACID

SOLUTION; INTRAVENOUS

ASCOR

+! MCGUFF

25,000MG/50ML (500MG/ML)

N209112 001 Oct 02, 2017

ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; TOCOPHEROL ACETATE; VITAMIN A; VITAMIN K

INJECTABLE; INTRAVENOUS

INFUVITE PEDIATRIC

+! SANDOZ CANADA INC

80MG/VIAL; 0.02MG/VIAL; 400 IU/VIAL; 0.001MG/VIAL; 5MG/VIAL; 0.14MG/VIAL; 17MG/VIAL; 1MG/VIAL; 1.4MG/VIAL; 1.2MG/VIAL; 7 IU/VIAL; 2,300 IU/VIAL; 0.2MG/VIAL

N021265 001 Feb 21, 2001

INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)

+! SANDOZ CANADA INC

80MG/VIAL; 0.02MG/VIAL; 400 IU/VIAL; 0.001MG/VIAL; 5MG/VIAL; 0.14MG/VIAL; 17MG/VIAL; 1MG/VIAL; 1.4MG/VIAL; 1.2MG/VIAL; 7 IU/VIAL; 2,300 IU/VIAL; 0.2MG/VIAL

N021265 002 Jan 29, 2004

ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

MOVIPREP**AA** +! SALIX PHARMS **4.7GM; 100GM; 1.015GM; 5.9GM; 2.691GM; 7.5GM** **N021881 001** Aug 02, 2006PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND ASCORBIC**AA** NOVEL LABS INC **4.7GM; 100GM; 1.015GM; 5.9GM; 2.691GM; 7.5GM** **A090145 001** Jan 25, 2012

PLENVU

+! SALIX

7.54GM; 140GM; 2.2GM; 48.11GM; 5.2GM; 9GM

N209381 001 May 04, 2018

ASENAPINE

SYSTEM; TRANSDERMAL

SECUADO

+! HISAMITSU

3.8MG/24HR

N212268 001 Oct 11, 2019

+

5.7MG/24HR

N212268 002 Oct 11, 2019

+

7.6MG/24HR

N212268 003 Oct 11, 2019

ASENAPINE MALEATE

TABLET; SUBLINGUAL

ASENAPINE MALEATE**AB** ALEMBIC **EQ 2.5MG BASE** **A206098 003** Jul 19, 2021**AB** **EQ 5MG BASE** **A206098 001** Dec 10, 2020**AB** **EQ 10MG BASE** **A206098 002** Dec 10, 2020**AB** BRECKENRIDGE **EQ 2.5MG BASE** **A205960 001** Dec 10, 2020**AB** **EQ 5MG BASE** **A205960 003** Mar 08, 2021**AB** **EQ 10MG BASE** **A205960 002** Dec 10, 2020**AB** SIGMAPHARM LABS LLC **EQ 5MG BASE** **A206107 001** Dec 10, 2020**AB** **EQ 10MG BASE** **A206107 002** Dec 10, 2020SAPHRIS**AB** + ALLERGAN **EQ 2.5MG BASE** **N022117 003** Mar 12, 2015**AB** + **EQ 5MG BASE** **N022117 001** Aug 13, 2009**AB** +! **EQ 10MG BASE** **N022117 002** Aug 13, 2009ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE**AA** NOSTRUM LABS INC **325MG; 50MG; 40MG** **A078149 001** Jun 13, 2007LANORINAL**AA** ! LANNETT **325MG; 50MG; 40MG** **A086996 002** Oct 11, 1985

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

! STRIDES PHARMA 325MG; 50MG; 40MG

A204195 001 Sep 22, 2016

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE**AB** LGM PHARMA **325MG; 50MG; 40MG; 30MG** **A075231 001** Nov 30, 2001**AB** ! STEVENS J **325MG; 50MG; 40MG; 30MG** **A074951 001** Aug 31, 1998

PRESCRIPTION DRUG PRODUCT LIST

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENGESIC

GALT PHARMS

385MG; 30MG; 25MG

A075141 001 May 29, 1998

ORPHENGESIC FORTE

! GALT PHARMS

770MG; 60MG; 50MG

A075141 002 May 29, 1998

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

ASPIRIN AND DIPYRIDAMOLE

AB	AMNEAL PHARMS	25MG; 200MG	A206392 001	Mar 08, 2016
AB	BARR	25MG; 200MG	A078804 001	Aug 14, 2009
AB	DR REDDYS	25MG; 200MG	A209048 001	Oct 10, 2018
AB	! GLENMARK PHARMS SA	25MG; 200MG	A210318 001	May 24, 2019
AB	MICRO LABS	25MG; 200MG	A209929 001	Aug 11, 2021
AB	PAR PHARM INC	25MG; 200MG	A207944 001	Jan 18, 2017
AB	SANDOZ	25MG; 200MG	A206739 001	Jan 18, 2017
AB	ZYDUS PHARMS	25MG; 200MG	A206753 001	Aug 29, 2017

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ASPIRIN

! LGM PHARMA

500MG; 5MG

A205479 001 May 28, 2021

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ASPIRIN

AA	EPIC PHARMA LLC	325MG; 4.8355MG	A040910 001	Jul 16, 2020
AA	+! ENDO PHARMS	325MG; 4.8355MG	N007337 007	Aug 05, 2005

ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

AB	AMNEAL	EQ 150MG BASE	A209717 002	Jun 01, 2020
AB		EQ 200MG BASE	A209717 003	Jun 01, 2020
AB		EQ 300MG BASE	A209717 004	Jun 01, 2020
AB	AUROBINDO PHARMA	EQ 100MG BASE	A204806 001	Jun 25, 2018
AB		EQ 150MG BASE	A204806 002	Jun 25, 2018
AB		EQ 200MG BASE	A204806 003	Jun 25, 2018
AB		EQ 300MG BASE	A204806 004	Jun 25, 2018
AB	HETERO LABS LTD III	EQ 150MG BASE	A212278 001	Feb 02, 2022
AB		EQ 200MG BASE	A212278 002	Feb 02, 2022
AB		EQ 300MG BASE	A212278 003	Feb 02, 2022
AB	LAURUS	EQ 150MG BASE	A212579 001	Apr 30, 2021
AB		EQ 200MG BASE	A212579 002	Apr 30, 2021
AB		EQ 300MG BASE	A212579 003	Apr 30, 2021
AB	TEVA PHARMS USA	EQ 100MG BASE	A091673 001	Apr 22, 2014
AB		EQ 150MG BASE	A091673 002	Apr 22, 2014
AB		EQ 200MG BASE	A091673 003	Apr 22, 2014
AB		EQ 300MG BASE	A091673 004	Apr 22, 2014

REYATAZ

AB	+ BRISTOL MYERS SQUIBB	EQ 200MG BASE	N021567 003	Jun 20, 2003
AB	+!	EQ 300MG BASE	N021567 004	Oct 16, 2006

POWDER; ORAL

REYATAZ

+! BRISTOL MYERS
SQUIBB

EQ 50MG BASE/PACKET

N206352 001 Jun 02, 2014

ATAZANAVIR SULFATE; COBICISTAT

TABLET; ORAL

EVOTAZ

+! BRISTOL

EQ 300MG BASE; 150MG

N206353 001 Jan 29, 2015

ATENOLOL

TABLET; ORAL

ATENOLOL

AB	AUROBINDO PHARMA	25MG	A078512 001	Oct 31, 2007
AB		50MG	A078512 002	Oct 31, 2007
AB		100MG	A078512 003	Oct 31, 2007
AB	HLTHCARE	25MG	A073026 002	May 01, 1992
AB		50MG	A073026 003	Sep 17, 1991
AB		100MG	A073026 001	Sep 17, 1991
AB	IPCA LABS LTD	25MG	A077877 001	Dec 27, 2006
AB		50MG	A077877 002	Dec 27, 2006

PRESCRIPTION DRUG PRODUCT LIST

ATENOLOL

TABLET; ORAL

ATENOLOL

<u>AB</u>		<u>100MG</u>	<u>A077877 003</u>	Dec 27, 2006
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A073457 002</u>	Apr 26, 1999
<u>AB</u>		<u>50MG</u>	<u>A073457 003</u>	Jan 24, 1992
<u>AB</u>		<u>100MG</u>	<u>A073457 001</u>	Jan 24, 1992
<u>AB</u>	TEVA	<u>25MG</u>	<u>A074056 003</u>	Jul 19, 2004
<u>AB</u>		<u>50MG</u>	<u>A074056 001</u>	Jan 18, 1995
<u>AB</u>		<u>100MG</u>	<u>A074056 002</u>	Jan 18, 1995
<u>AB</u>	TWI PHARMS	<u>25MG</u>	<u>A072304 002</u>	Jul 31, 1992
<u>AB</u>		<u>50MG</u>	<u>A072304 003</u>	Jul 18, 1988
<u>AB</u>		<u>100MG</u>	<u>A072304 001</u>	Jul 15, 1988
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A213136 001</u>	Nov 21, 2019
<u>AB</u>		<u>50MG</u>	<u>A213136 002</u>	Nov 21, 2019
<u>AB</u>		<u>100MG</u>	<u>A213136 003</u>	Nov 21, 2019
<u>AB</u>	UNIQUE	<u>25MG</u>	<u>A077443 001</u>	Sep 13, 2006
<u>AB</u>		<u>50MG</u>	<u>A077443 002</u>	Sep 13, 2006
<u>AB</u>		<u>100MG</u>	<u>A077443 003</u>	Sep 13, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A076900 001</u>	Jan 28, 2005
<u>AB</u>		<u>50MG</u>	<u>A076900 002</u>	Jan 28, 2005
<u>AB</u>		<u>100MG</u>	<u>A076900 003</u>	Jan 28, 2005

TENORMIN

<u>AB</u>	+	TWI PHARMS	<u>25MG</u>	<u>N018240 004</u>	Apr 09, 1990
<u>AB</u>	+		<u>50MG</u>	<u>N018240 001</u>	
<u>AB</u>	+	!	<u>100MG</u>	<u>N018240 002</u>	

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

<u>AB</u>	NOVITIUM PHARMA	<u>50MG; 25MG</u>	<u>A215560 001</u>	Oct 25, 2021
<u>AB</u>		<u>100MG; 25MG</u>	<u>A215560 002</u>	Oct 25, 2021
<u>AB</u>	TWI PHARMS	<u>50MG; 25MG</u>	<u>A072302 002</u>	May 31, 1990
<u>AB</u>		<u>100MG; 25MG</u>	<u>A072302 001</u>	May 31, 1990
<u>AB</u>	UNICHEM	<u>50MG; 25MG</u>	<u>A213302 001</u>	Nov 25, 2020
<u>AB</u>		<u>100MG; 25MG</u>	<u>A213302 002</u>	Nov 25, 2020
<u>AB</u>	WATSON LABS	<u>50MG; 25MG</u>	<u>A073665 001</u>	Jul 02, 1992
<u>AB</u>		<u>100MG; 25MG</u>	<u>A073665 002</u>	Jul 02, 1992
<u>AB</u>	ZYDUS PHARMS	<u>50MG; 25MG</u>	<u>A210028 001</u>	Mar 08, 2019
<u>AB</u>		<u>100MG; 25MG</u>	<u>A210028 002</u>	Mar 08, 2019

TENORETIC 100

<u>AB</u>	+	!	TWI PHARMS	<u>100MG; 25MG</u>	<u>N018760 001</u>	Jun 08, 1984
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TENORETIC 50

<u>AB</u>	+		TWI PHARMS	<u>50MG; 25MG</u>	<u>N018760 002</u>	Jun 08, 1984
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ATOGEFANT

TABLET; ORAL

QULIPTA

	+	ABBVIE	10MG	N215206 001	Sep 28, 2021
	+		30MG	N215206 002	Sep 28, 2021
	+	!	60MG	N215206 003	Sep 28, 2021

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>10MG</u>	<u>A078983 001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A078983 002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A078983 003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A078983 004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A078983 005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A078983 006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A078983 007</u>	May 30, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A079016 001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079016 002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079016 003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079016 004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079016 005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079016 006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079016 007</u>	May 30, 2017
<u>AB</u>	DR REDDYS	<u>10MG</u>	<u>A090609 001</u>	Feb 23, 2018
<u>AB</u>		<u>18MG</u>	<u>A090609 002</u>	Feb 23, 2018
<u>AB</u>		<u>25MG</u>	<u>A090609 003</u>	Feb 23, 2018
<u>AB</u>		<u>40MG</u>	<u>A090609 004</u>	Feb 23, 2018
<u>AB</u>		<u>60MG</u>	<u>A090609 005</u>	Feb 23, 2018

PRESCRIPTION DRUG PRODUCT LIST

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u>		<u>80MG</u>	<u>A090609</u>	<u>006</u>	Feb 23, 2018
<u>AB</u>		<u>100MG</u>	<u>A090609</u>	<u>007</u>	Feb 23, 2018
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A079019</u>	<u>001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079019</u>	<u>002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079019</u>	<u>003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079019</u>	<u>004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079019</u>	<u>005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079019</u>	<u>006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079019</u>	<u>007</u>	May 30, 2017
<u>AB</u>	HETERO LABS LTD V	<u>10MG</u>	<u>A202682</u>	<u>001</u>	Mar 11, 2021
<u>AB</u>		<u>18MG</u>	<u>A202682</u>	<u>002</u>	Mar 11, 2021
<u>AB</u>		<u>25MG</u>	<u>A202682</u>	<u>003</u>	Mar 11, 2021
<u>AB</u>		<u>40MG</u>	<u>A202682</u>	<u>004</u>	Mar 11, 2021
<u>AB</u>		<u>60MG</u>	<u>A202682</u>	<u>005</u>	Mar 11, 2021
<u>AB</u>		<u>80MG</u>	<u>A202682</u>	<u>006</u>	Mar 11, 2021
<u>AB</u>		<u>100MG</u>	<u>A202682</u>	<u>007</u>	Mar 11, 2021
<u>AB</u>	TEVA PHARMS USA	<u>10MG</u>	<u>A079022</u>	<u>001</u>	May 30, 2017
<u>AB</u>		<u>18MG</u>	<u>A079022</u>	<u>002</u>	May 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A079022</u>	<u>003</u>	May 30, 2017
<u>AB</u>		<u>40MG</u>	<u>A079022</u>	<u>004</u>	May 30, 2017
<u>AB</u>		<u>60MG</u>	<u>A079022</u>	<u>005</u>	May 30, 2017
<u>AB</u>		<u>80MG</u>	<u>A079022</u>	<u>006</u>	May 30, 2017
<u>AB</u>		<u>100MG</u>	<u>A079022</u>	<u>007</u>	May 30, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>10MG</u>	<u>A079017</u>	<u>007</u>	Apr 05, 2023
<u>AB</u>		<u>18MG</u>	<u>A079017</u>	<u>001</u>	Sep 17, 2010
<u>AB</u>		<u>25MG</u>	<u>A079017</u>	<u>002</u>	Sep 17, 2010
<u>AB</u>		<u>40MG</u>	<u>A079017</u>	<u>003</u>	Sep 17, 2010
<u>AB</u>		<u>60MG</u>	<u>A079017</u>	<u>004</u>	Sep 17, 2010
<u>AB</u>		<u>80MG</u>	<u>A079017</u>	<u>005</u>	Sep 17, 2010
<u>AB</u>		<u>100MG</u>	<u>A079017</u>	<u>006</u>	Sep 17, 2010

STRATTERA

<u>AB</u>	+	LILLY	<u>10MG</u>	<u>N021411</u>	<u>002</u>	Nov 26, 2002
<u>AB</u>	+		<u>18MG</u>	<u>N021411</u>	<u>003</u>	Nov 26, 2002
<u>AB</u>	+		<u>25MG</u>	<u>N021411</u>	<u>004</u>	Nov 26, 2002
<u>AB</u>	+		<u>40MG</u>	<u>N021411</u>	<u>005</u>	Nov 26, 2002
<u>AB</u>	+	!	<u>60MG</u>	<u>N021411</u>	<u>006</u>	Nov 26, 2002
<u>AB</u>	+		<u>80MG</u>	<u>N021411</u>	<u>007</u>	Feb 14, 2005
<u>AB</u>	+		<u>100MG</u>	<u>N021411</u>	<u>008</u>	Feb 14, 2005

ATORVASTATIN CALCIUM

SUSPENSION; ORAL

ATORVALIQ

+! CMP DEV LLC

20MG/5ML

N213260 001 Feb 01, 2023

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 10MG BASE</u>	<u>A207687</u>	<u>001</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207687</u>	<u>002</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207687</u>	<u>003</u>	Mar 30, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A207687</u>	<u>004</u>	Mar 30, 2018
<u>AB</u>	ACI	<u>EQ 10MG BASE</u>	<u>A217634</u>	<u>001</u>	Dec 21, 2023
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A217634</u>	<u>002</u>	Dec 21, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217634</u>	<u>003</u>	Dec 21, 2023
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A217634</u>	<u>004</u>	Dec 21, 2023
<u>AB</u>	AGNITIO	<u>EQ 10MG BASE</u>	<u>A214969</u>	<u>001</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214969</u>	<u>002</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214969</u>	<u>003</u>	Sep 02, 2021
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214969</u>	<u>004</u>	Sep 02, 2021
<u>AB</u>	ALKEM LABS LTD	<u>EQ 10MG BASE</u>	<u>A209288</u>	<u>001</u>	Dec 21, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A209288</u>	<u>002</u>	Dec 21, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A209288</u>	<u>003</u>	Dec 21, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A209288</u>	<u>004</u>	Dec 21, 2018
<u>AB</u>	APOTEX INC	<u>EQ 10MG BASE</u>	<u>A090548</u>	<u>001</u>	May 29, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090548</u>	<u>002</u>	May 29, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090548</u>	<u>003</u>	May 29, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A090548</u>	<u>004</u>	May 29, 2012
<u>AB</u>	BIOCON PHARMA	<u>EQ 10MG BASE</u>	<u>A216436</u>	<u>001</u>	Nov 23, 2022
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A216436</u>	<u>002</u>	Nov 23, 2022
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A216436</u>	<u>003</u>	Nov 23, 2022
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A216436</u>	<u>004</u>	Nov 23, 2022

PRESCRIPTION DRUG PRODUCT LIST

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>	CADILA PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A212103 001</u>	Oct 16, 2023
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A212103 002</u>	Oct 16, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A212103 003</u>	Oct 16, 2023
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A212103 004</u>	Oct 16, 2023
<u>AB</u>	DR REDDYS	<u>EQ 10MG BASE</u>	<u>A214659 001</u>	Jul 14, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214659 002</u>	Jul 14, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214659 003</u>	Jul 14, 2021
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214659 004</u>	Jul 14, 2021
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A091650 001</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091650 002</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091650 003</u>	Jul 17, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A202357 001</u>	Jul 17, 2012
<u>AB</u>	GRAVITI PHARMS	<u>EQ 10MG BASE</u>	<u>A209912 001</u>	Jun 18, 2018
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A209912 002</u>	Jun 18, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A209912 003</u>	Jun 18, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A209912 004</u>	Jun 18, 2018
<u>AB</u>	HETERO LABS LTD V	<u>EQ 10MG BASE</u>	<u>A214344 001</u>	Sep 12, 2023
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A214344 002</u>	Sep 12, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214344 003</u>	Sep 12, 2023
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A214344 004</u>	Sep 12, 2023
<u>AB</u>	INVAGEN PHARMS	<u>EQ 10MG BASE</u>	<u>A204846 001</u>	Jan 09, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A204846 002</u>	Jan 09, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A204846 003</u>	Jan 09, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204846 004</u>	Jan 09, 2017
<u>AB</u>	LANNETT CO INC	<u>EQ 10MG BASE</u>	<u>A091624 001</u>	Apr 05, 2013
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091624 002</u>	Apr 05, 2013
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091624 003</u>	Apr 05, 2013
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A091624 004</u>	Apr 05, 2013
<u>AB</u>	LUPIN LTD	<u>EQ 10MG BASE</u>	<u>A204991 001</u>	Mar 06, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A204991 002</u>	Mar 06, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A204991 003</u>	Mar 06, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204991 004</u>	Mar 06, 2019
<u>AB</u>	MANKIND PHARMA	<u>EQ 10MG BASE</u>	<u>A217081 001</u>	Jan 05, 2024
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A217081 002</u>	Jan 05, 2024
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217081 003</u>	Jan 05, 2024
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A217081 004</u>	Jan 05, 2024
<u>AB</u>	MICRO LABS LTD INDIA	<u>EQ 10MG BASE</u>	<u>A205945 001</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205945 002</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205945 003</u>	Nov 07, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A205945 004</u>	Nov 07, 2019
<u>AB</u>	MSN	<u>EQ 10MG BASE</u>	<u>A211933 001</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211933 002</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211933 003</u>	Feb 08, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A211933 004</u>	Feb 08, 2019
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A091226 001</u>	May 29, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A091226 002</u>	May 29, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091226 003</u>	May 29, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A091226 004</u>	May 29, 2012
<u>AB</u>	SANDOZ INC	<u>EQ 10MG BASE</u>	<u>A077575 001</u>	May 29, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077575 002</u>	May 29, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077575 003</u>	May 29, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077575 004</u>	May 29, 2012
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A205519 001</u>	May 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205519 002</u>	May 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205519 003</u>	May 19, 2016
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A205519 004</u>	May 19, 2016
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 10MG BASE</u>	<u>A076477 001</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076477 002</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076477 003</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A076477 004</u>	Nov 30, 2011
<u>AB</u>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A205300 001</u>	Mar 27, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205300 002</u>	Mar 27, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205300 003</u>	Mar 27, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A205300 004</u>	Mar 27, 2017
<u>AB</u>	UMEDICA	<u>EQ 10MG BASE</u>	<u>A213853 001</u>	Aug 19, 2020
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A213853 002</u>	Aug 19, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213853 003</u>	Aug 19, 2020
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A213853 004</u>	Aug 19, 2020
<u>AB</u>	ZYDUS PHARMS	<u>EQ 10MG BASE</u>	<u>A206536 001</u>	Nov 20, 2018

PRESCRIPTION DRUG PRODUCT LIST

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A206536 002</u>	Nov 20, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206536 003</u>	Nov 20, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A206536 004</u>	Nov 20, 2018
<u>LIPITOR</u>				
<u>AB</u>	+	UPJOHN	<u>EQ 10MG BASE</u>	<u>N020702 001</u> Dec 17, 1996
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N020702 002</u> Dec 17, 1996
<u>AB</u>	+		<u>EQ 40MG BASE</u>	<u>N020702 003</u> Dec 17, 1996
<u>AB</u>	+	!	<u>EQ 80MG BASE</u>	<u>N020702 004</u> Apr 07, 2000

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET; ORAL

LYPQOZET

	ALThERA PHARMS	EQ 10MG BASE;10MG	A206084 001	Apr 26, 2017
		EQ 20MG BASE;10MG	A206084 002	Apr 26, 2017
		EQ 40MG BASE;10MG	A206084 003	Apr 26, 2017
	!	EQ 80MG BASE;10MG	A206084 004	Apr 26, 2017

ATOVAQUONE

SUSPENSION; ORAL

ATOVAQUONE

<u>AB</u>	ABHAI LLC	<u>750MG/5ML</u>	<u>A210510 001</u>	May 31, 2019
<u>AB</u>	ABON PHARMS LLC	<u>750MG/5ML</u>	<u>A214272 001</u>	Oct 25, 2021
<u>AB</u>	AMNEAL PHARMS	<u>750MG/5ML</u>	<u>A202960 001</u>	Mar 18, 2014
<u>AB</u>	APOTEX	<u>750MG/5ML</u>	<u>A209750 001</u>	Oct 11, 2017
<u>AB</u>	BIONPHARMA	<u>750MG/5ML</u>	<u>A212918 001</u>	Mar 30, 2021
<u>AB</u>	CHARTWELL RX	<u>750MG/5ML</u>	<u>A207833 001</u>	Apr 28, 2017
<u>AB</u>	GLENMARK PHARMS	<u>750MG/5ML</u>	<u>A209685 001</u>	Nov 21, 2018
<u>AB</u>	HETERO LABS LTD III	<u>750MG/5ML</u>	<u>A210692 001</u>	Oct 11, 2018
<u>AB</u>	LUPIN LTD	<u>750MG/5ML</u>	<u>A209105 001</u>	Sep 11, 2018
<u>MEPRON</u>				
<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>750MG/5ML</u>	<u>N020500 001</u> Feb 08, 1995

ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET; ORAL

ATOVAQUONE AND PROGUANIL HYDROCHLORIDE

<u>AB</u>	GLENMARK GENERICS	<u>62.5MG;25MG</u>	<u>A091211 002</u>	Apr 06, 2015
<u>AB</u>		<u>250MG;100MG</u>	<u>A091211 001</u>	Jan 12, 2011
<u>AB</u>	MYLAN	<u>62.5MG;25MG</u>	<u>A202362 001</u>	May 27, 2014
<u>AB</u>		<u>250MG;100MG</u>	<u>A202362 002</u>	May 27, 2014
<u>MALARONE</u>				
<u>AB</u>	+	GLAXOSMITHKLINE	<u>250MG;100MG</u>	<u>N021078 001</u> Jul 14, 2000
<u>MALARONE PEDIATRIC</u>				
<u>AB</u>	+	GLAXOSMITHKLINE	<u>62.5MG;25MG</u>	<u>N021078 002</u> Jul 14, 2000

ATRAURIUM BESYLATE

INJECTABLE; INJECTION

ATRAURIUM BESYLATE

<u>AP</u>	EUGIA PHARMA	<u>10MG/ML</u>	<u>A206011 001</u>	Apr 08, 2015
<u>AP</u>	!	HIKMA	<u>10MG/ML</u>	<u>A074901 001</u> Jul 18, 1997
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090761 001</u>	Oct 18, 2012
<u>AP</u>	MEITHEAL	<u>10MG/ML</u>	<u>A091489 001</u>	Feb 17, 2012

ATRAURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>	EUGIA PHARMA	<u>10MG/ML</u>	<u>A206010 001</u>	Apr 08, 2015
<u>AP</u>	!	HIKMA	<u>10MG/ML</u>	<u>A074900 001</u> Jul 18, 1997
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090782 001</u>	Oct 18, 2012
<u>AP</u>	MEITHEAL	<u>10MG/ML</u>	<u>A091488 001</u>	Feb 17, 2012

ATROPINE SULFATE

SOLUTION; INTRAVENOUS

ATROPINE SULFATE

<u>AP</u>	ACCORD HLTHCARE	<u>0.25MG/5ML (0.05MG/ML)</u>	<u>A212868 001</u>	Jul 26, 2021
<u>AP</u>	+	<u>0.4MG/ML (0.4MG/ML)</u>	<u>N214652 001</u>	Sep 29, 2020
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A212868 002</u>	Jul 26, 2021
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A212868 003</u>	Jul 26, 2021
<u>AP</u>	+	<u>1MG/ML (1MG/ML)</u>	<u>N214652 002</u>	Sep 29, 2020
<u>AP</u>	AM REGENT	<u>0.4MG/ML (0.4MG/ML)</u>	<u>A216120 001</u>	May 26, 2022
<u>AP</u>		<u>1MG/ML (1MG/ML)</u>	<u>A216120 002</u>	May 26, 2022
<u>AP</u>	AMNEAL	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A215342 001</u>	Jan 26, 2022
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A215342 002</u>	Oct 25, 2023
<u>AP</u>	+	HOSPIRA	<u>0.25MG/5ML (0.05MG/ML)</u>	<u>N021146 002</u> Jul 09, 2001
<u>AP</u>	+		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N021146 001</u> Jul 09, 2001
<u>AP</u>	+		<u>1MG/10ML (0.1MG/ML)</u>	<u>N021146 003</u> Jul 09, 2001

PRESCRIPTION DRUG PRODUCT LIST

ATROPINE SULFATE

SOLUTION;INTRAVENOUS

ATROPINE SULFATE

AP	INTL MEDICATION SYS	1MG/10ML (0.1MG/ML)	A212461 001	Oct 05, 2020
AP	MEDEFIL INC	1MG/10ML (0.1MG/ML)	A214970 001	Nov 04, 2022

SOLUTION;INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, INTRAOSSEOUS, ENDOTRACHEAL

ATROPINE SULFATE

AP	ACCORD HLTHCARE	8MG/20ML (0.4MG/ML)	A213424 001	Mar 19, 2021
AP	+! FRESENIUS KABI USA	8MG/20ML (0.4MG/ML)	N209260 001	Jan 26, 2018
AP	HIKMA	8MG/20ML (0.4MG/ML)	A213561 001	Dec 01, 2021

SOLUTION/DROPS;OPHTHALMIC

ATROPINE SULFATE

AT	! AMNEAL	1%	A214752 001	Jul 14, 2022
AT	APOTEX	1%	A215624 001	Nov 26, 2021
AT	MANKIND PHARMA	1%	A218148 001	Jan 08, 2024
AT	+ RISING	1%	N206289 001	Jul 18, 2014
	+! BAUSCH AND LOMB INC	1%	N213581 001	Mar 15, 2022
	ISOPTO ATROPINE			
	+! ALCON LABS INC	1%	N208151 001	Dec 01, 2016

ATROPINE SULFATE; DIFENOXYN HYDROCHLORIDE

TABLET;ORAL

MOTOFEN

+!	SEBELA IRELAND LTD	0.025MG;1MG	N017744 002	
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ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION;ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

!	HIKMA	0.025MG/5ML;2.5MG/5ML	A087708 001	May 03, 1982
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TABLET;ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

AA	ANI PHARMS	0.025MG;2.5MG	A086727 001	
AA	BAYSHORE PHARMS LLC	0.025MG;2.5MG	A210819 001	Nov 13, 2018
AA	CHARTWELL RX	0.025MG;2.5MG	A207128 001	Oct 21, 2020
AA	LANNETT	0.025MG;2.5MG	A085372 001	
AA	LEADING	0.025MG;2.5MG	A213413 001	Feb 20, 2020
AA	MYLAN	0.025MG;2.5MG	A085762 001	
AA	SPECGX LLC	0.025MG;2.5MG	A213335 001	Oct 06, 2020
AA	WINDER LABS LLC	0.025MG;2.5MG	A211362 001	Jan 27, 2021

LOMOTIL

AA	+! PFIZER	0.025MG;2.5MG	N012462 001	
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ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE;INTRAMUSCULAR

DUODOTE

+!	MMT	2.1MG/0.7ML;600MG/2ML	N021983 001	Sep 28, 2006
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AURANOFIN

CAPSULE;ORAL

RIDAURA

+!	SEBELA IRELAND LTD	3MG	N018689 001	May 24, 1985
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AVACINCAPTAD PEGOL SODIUM

SOLUTION;INTRAVITREAL

IZERVAY

+!	IVERIC BIO	EQ 2MG BASE/0.1ML (EQ 2MG BASE/0.1ML)	N217225 001	Aug 04, 2023
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AVACOPAN

CAPSULE;ORAL

TAVNEOS

+!	CHEMOCENTRYX	10MG	N214487 001	Oct 07, 2021
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AVANAFIL

TABLET;ORAL

STENDRA

+	METUCHEN PHARMS	50MG	N202276 001	Apr 27, 2012
+		100MG	N202276 002	Apr 27, 2012
+!		200MG	N202276 003	Apr 27, 2012

AVAPRITINIB

TABLET;ORAL

AYVAKIT

+	BLUEPRINT MEDICINES	25MG	N212608 004	Jun 16, 2021
+		50MG	N212608 005	Jun 16, 2021
+		100MG	N212608 001	Jan 09, 2020
+		200MG	N212608 002	Jan 09, 2020

PRESCRIPTION DRUG PRODUCT LISTAVAPRITINIB

TABLET; ORAL

AYVAKIT

+!

300MG

N212608 003 Jan 09, 2020

AVATROMBOPAG MALEATE

TABLET; ORAL

DOPTELET

+!

AKARX INC

EQ 20MG BASE

N210238 001 May 21, 2018

AVIBACTAM SODIUM; CEFTAZIDIME

POWDER; INTRAVENOUS

AVYCAZ

+!

ALLERGAN

EQ 0.5GM BASE; 2GM/VIAL

N206494 001 Feb 25, 2015

AXITINIB

TABLET; ORAL

INLYTA

+

PF PRISM CV

1MG

N202324 001 Jan 27, 2012

+!

5MG

N202324 002 Jan 27, 2012

AZACITIDINE

POWDER; INTRAVENOUS, SUBCUTANEOUS

AZACITIDINE

<u>AP</u>	ACCORD HLTHCARE	<u>100MG/VIAL</u>	<u>A207475 001</u>	Jul 02, 2018
<u>AP</u>	ACTAVIS LLC	<u>100MG/VIAL</u>	<u>N208216 001</u>	Apr 29, 2016
<u>AP</u>	AMNEAL	<u>100MG/VIAL</u>	<u>A211549 001</u>	Feb 03, 2022
<u>AP</u>	CIPLA	<u>100MG/VIAL</u>	<u>A209540 001</u>	May 04, 2018
<u>AP</u>	DR REDDYS	<u>100MG/VIAL</u>	<u>A201537 001</u>	Sep 16, 2013
<u>AP</u>	EUGIA PHARMA	<u>100MG/VIAL</u>	<u>A215066 001</u>	Dec 30, 2022
<u>AP</u>	EUROHLTH INTL SARL	<u>100MG/VIAL</u>	<u>A209337 001</u>	Jun 08, 2020
<u>AP</u>	JIANGSU HANSOH PHARM	<u>100MG/VIAL</u>	<u>A215905 001</u>	Jun 28, 2023
<u>AP</u>	MEITHEAL	<u>100MG/VIAL</u>	<u>A212128 001</u>	Nov 02, 2020
<u>AP</u>	NATCO PHARMA LTD	<u>100MG/VIAL</u>	<u>A207234 001</u>	Jun 23, 2017
<u>AP</u>	SHILPA MEDICARE	<u>100MG/VIAL</u>	<u>A207518 001</u>	Sep 29, 2016

VIDAZA

<u>AP</u>	+! BRISTOL-MYERS	<u>100MG/VIAL</u>	<u>N050794 001</u>	May 19, 2004
TABLET; ORAL				
	ONUREG			
	+ BRISTOL	200MG	N214120 001	Sep 01, 2020
	+!	300MG	N214120 002	Sep 01, 2020

AZATHIOPRINE

TABLET; ORAL

AZASAN

<u>AB</u>	AAIPHARMA LLC	<u>25MG</u>	<u>A075252 002</u>	Feb 03, 2003
<u>AB</u>		<u>50MG</u>	<u>A075252 001</u>	Jun 07, 1999
<u>AB</u>		<u>75MG</u>	<u>A075252 003</u>	Feb 03, 2003
<u>AB</u>		<u>100MG</u>	<u>A075252 004</u>	Feb 03, 2003

AZATHIOPRINE

<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A208687 001</u>	Mar 27, 2020
<u>AB</u>		<u>50MG</u>	<u>A208687 002</u>	Mar 27, 2020
<u>AB</u>		<u>75MG</u>	<u>A208687 003</u>	Mar 27, 2020
<u>AB</u>		<u>100MG</u>	<u>A208687 004</u>	Mar 27, 2020
<u>AB</u>	AMNEAL	<u>50MG</u>	<u>A074069 001</u>	Feb 16, 1996
<u>AB</u>		<u>75MG</u>	<u>A074069 002</u>	Nov 02, 2021
<u>AB</u>		<u>100MG</u>	<u>A074069 003</u>	Nov 02, 2021
<u>AB</u>	RISING	<u>50MG</u>	<u>A075568 001</u>	Dec 13, 1999
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A077621 002</u>	Sep 05, 2008
<u>AB</u>		<u>50MG</u>	<u>A077621 001</u>	Mar 15, 2007
<u>AB</u>		<u>75MG</u>	<u>A077621 003</u>	Sep 05, 2008
<u>AB</u>		<u>100MG</u>	<u>A077621 004</u>	Sep 05, 2008

TMURAN

<u>AB</u>	+! SEBELA IRELAND LTD	<u>50MG</u>	<u>N016324 001</u>	
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AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

AZATHIOPRINE SODIUM

!

HIKMA

EQ 100MG BASE/VIAL

A074419 001 Mar 31, 1995

PRESCRIPTION DRUG PRODUCT LIST

AZELAIC ACID

AEROSOL, FOAM;TOPICAL

FINACEA

+! LEO PHARMA AS 15% N207071 001 Jul 29, 2015

CREAM;TOPICAL

AZELEX

+! ALMIRALL 20% N020428 001 Sep 13, 1995

GEL;TOPICAL

AZELAIC ACID

AB	ACTAVIS LABS UT INC	15%	<u>A208011</u>	<u>001</u>	Nov 19, 2018
AB	ENCUBE	15%	<u>A208724</u>	<u>001</u>	Nov 19, 2018
AB	GLENMARK PHARMS	15%	<u>A204637</u>	<u>001</u>	Nov 19, 2018
AB	TARO	15%	<u>A210549</u>	<u>001</u>	Aug 23, 2019

FINACEA

AB	+! LEO PHARMA AS	15%	<u>N021470</u>	<u>001</u>	Dec 24, 2002
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AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AZELASTINE HYDROCHLORIDE

AT	ALEMBIC	0.05%	<u>A209620</u>	<u>001</u>	Mar 20, 2019
AT	APOTEX INC	0.05%	<u>A078621</u>	<u>001</u>	Aug 03, 2009
AT	GLAND PHARMA LTD	0.05%	<u>A210092</u>	<u>001</u>	Feb 25, 2020
AT	! SANDOZ	0.05%	<u>A202305</u>	<u>001</u>	May 31, 2012
AT	SOMERSET THERAPS LLC	0.05%	<u>A207411</u>	<u>001</u>	Mar 29, 2019
AT	SUN PHARM	0.05%	<u>A078738</u>	<u>001</u>	Jun 21, 2010

SPRAY, METERED;NASAL

AZELASTINE HYDROCHLORIDE

AB	ALKEM LABS LTD	0.137MG/SPRAY	<u>A208156</u>	<u>001</u>	Aug 18, 2017
AB	AMNEAL	0.137MG/SPRAY	<u>A204660</u>	<u>001</u>	Aug 28, 2017
AB	! APOTEX INC	0.137MG/SPRAY	<u>A077954</u>	<u>001</u>	Apr 30, 2009
AB	AUROBINDO PHARMA LTD	0.137MG/SPRAY	<u>A212289</u>	<u>001</u>	May 08, 2020
AB	BIONPHARMA	0.137MG/SPRAY	<u>A090176</u>	<u>001</u>	Jul 28, 2015
AB	EPIC PHARMA LLC	0.137MG/SPRAY	<u>A207610</u>	<u>001</u>	May 17, 2019
AB	HIKMA	0.137MG/SPRAY	<u>A091444</u>	<u>001</u>	Oct 24, 2014
AB	SUN PHARM	0.137MG/SPRAY	<u>A090423</u>	<u>001</u>	May 23, 2012
AB	UPSHER SMITH LABS	0.137MG/SPRAY	<u>A202609</u>	<u>001</u>	Mar 17, 2017
AB	ZYDUS PHARMS	0.137MG/SPRAY	<u>A091409</u>	<u>001</u>	Aug 14, 2017

AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE

AB	! APOTEX	0.137MG/SPRAY;0.05MG/SPRAY	<u>A207712</u>	<u>001</u>	Apr 28, 2017
AB	PADAGIS ISRAEL	0.137MG/SPRAY;0.05MG/SPRAY	<u>A208111</u>	<u>001</u>	Feb 18, 2021

DYMISTA

AB	+ MYLAN SPECIALITY LP	0.137MG/SPRAY;0.05MG/SPRAY	<u>N202236</u>	<u>001</u>	May 01, 2012
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AZILSARTAN KAMEDOXOMIL

TABLET;ORAL

EDARBI

+ AZURITY EQ 40MG MEDOXOMIL N200796 001 Feb 25, 2011

+! EQ 80MG MEDOXOMIL N200796 002 Feb 25, 2011

AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE

TABLET;ORAL

EDARBYCLOR

+ AZURITY EQ 40MG MEDOXOMIL;12.5MG N202331 001 Dec 20, 2011

+! EQ 40MG MEDOXOMIL;25MG N202331 002 Dec 20, 2011

AZITHROMYCIN

FOR SUSPENSION;ORAL

AZITHROMYCIN

AB	AMNEAL	EQ 100MG BASE/5ML	<u>A205666</u>	<u>001</u>	Jul 19, 2018
AB		EQ 200MG BASE/5ML	<u>A205666</u>	<u>002</u>	Jul 19, 2018
AB	AUROBINDO PHARMA LTD	EQ 100MG BASE/5ML	<u>A209201</u>	<u>001</u>	Oct 09, 2018
AB		EQ 200MG BASE/5ML	<u>A209201</u>	<u>002</u>	Oct 09, 2018
AB	EPIC PHARMA LLC	EQ 100MG BASE/5ML	<u>A207531</u>	<u>001</u>	Apr 09, 2018
AB		EQ 200MG BASE/5ML	<u>A207531</u>	<u>002</u>	Apr 09, 2018
AB	HAINAN POLY	EQ 100MG BASE/5ML	<u>A217036</u>	<u>001</u>	Jul 27, 2023
AB		EQ 200MG BASE/5ML	<u>A217036</u>	<u>002</u>	Jul 27, 2023
AB	PLIVA	EQ 100MG BASE/5ML	<u>A065246</u>	<u>002</u>	Jul 05, 2006
AB		EQ 200MG BASE/5ML	<u>A065246</u>	<u>001</u>	Jul 05, 2006
AB	ZYDUS	EQ 100MG BASE/5ML	<u>A211147</u>	<u>001</u>	Jul 31, 2018

PRESCRIPTION DRUG PRODUCT LIST

AZITHROMYCIN

FOR SUSPENSION; ORAL

AZITHROMYCINAB EQ 200MG BASE/5ML A211147 002 Jul 31, 2018ZITHROMAXAB + PFIZER EQ 100MG BASE/5ML N050710 001 Oct 19, 1995AB +! EQ 200MG BASE/5ML N050710 002 Oct 19, 1995

+! EQ 1GM BASE/PACKET N050693 001 Sep 28, 1994

INJECTABLE; INJECTION

AZITHROMYCINAP EUGIA PHARMA EQ 500MG BASE/VIAL A203294 001 Jun 19, 2015AP FRESENIUS KABI USA EQ 500MG BASE/VIAL A065179 001 Dec 13, 2005AP GLAND PHARMA LTD EQ 500MG BASE/VIAL A065501 001 Nov 09, 2009AP HOSPIRA EQ 500MG BASE/VIAL A065500 001 Jun 26, 2009AP EQ 500MG BASE/VIAL A065511 001 Jun 26, 2009AP SLATE RUN PHARMA EQ 500MG BASE/VIAL A203412 001 Oct 09, 2018AP SUN PHARM INDS LTD EQ 500MG BASE/VIAL A090923 001 Apr 02, 2013ZITHROMAXAP +! PFIZER EQ 500MG BASE/VIAL N050733 001 Jan 30, 1997

SOLUTION/DROPS; OPHTHALMIC

AZASITE

+! THEA PHARMA 1% N050810 001 Apr 27, 2007

TABLET; ORAL

AZITHROMYCINAB ACI EQ 250MG BASE A215772 001 Jul 15, 2022AB EQ 500MG BASE A215773 001 Jul 12, 2022AB ALEMBIC EQ 250MG BASE A211791 001 Jan 28, 2020AB EQ 500MG BASE A211792 001 Jan 28, 2020AB EQ 600MG BASE A211793 001 Jan 27, 2020AB AUROBINDO PHARMA EQ 250MG BASE A207370 001 Jul 05, 2018

LTD

AB EQ 500MG BASE A207398 001 Jul 05, 2018AB BIONPHARMA EQ 250MG BASE A210000 001 Feb 26, 2019AB EQ 500MG BASE A210001 001 Feb 26, 2019AB EQ 600MG BASE A209999 001 Dec 26, 2018AB CHARTWELL RX EQ 250MG BASE A065404 001 Feb 11, 2008AB EQ 500MG BASE A065405 001 Feb 11, 2008AB EQ 600MG BASE A065302 003 Feb 11, 2008AB CSPC OUYI EQ 250MG BASE A208250 001 Apr 17, 2019AB EQ 500MG BASE A208249 001 Oct 25, 2018AB EQ 600MG BASE A207566 001 Sep 24, 2018AB LUPIN LTD EQ 250MG BASE A065398 001 May 15, 2015AB EQ 500MG BASE A065399 001 May 15, 2015AB EQ 600MG BASE A065400 001 May 15, 2015AB PLIVA EQ 250MG BASE A065225 001 Nov 14, 2005AB EQ 500MG BASE A065223 001 Nov 14, 2005AB ! EQ 600MG BASE A065218 001 Nov 14, 2005AB SANDOZ EQ 250MG BASE A065211 001 Nov 14, 2005AB EQ 500MG BASE A065212 001 Nov 14, 2005AB EQ 600MG BASE A065209 001 Nov 14, 2005AB SUNSHINE EQ 250MG BASE A209045 001 Dec 07, 2018AB EQ 500MG BASE A209044 001 Dec 07, 2018AB EQ 600MG BASE A209043 001 Dec 06, 2018AB TEVA EQ 500MG BASE A065193 001 Nov 14, 2005AB YUNG SHIN PHARM EQ 250MG BASE A211317 001 Jul 26, 2022AB EQ 600MG BASE A211068 001 May 08, 2020ZITHROMAXAB + PFIZER EQ 250MG BASE N050711 001 Jul 18, 1996AB + EQ 500MG BASE N050784 001 May 24, 2002AZTREONAM

FOR SOLUTION; INHALATION

CAYSTON

+! GILEAD 75MG/VIAL N050814 001 Feb 22, 2010

INJECTABLE; INJECTION

AZACTAMAP +! BRISTOL MYERS 1GM/VIAL N050580 002 Dec 31, 1986

SQUIBB

AP +! 2GM/VIAL N050580 003 Dec 31, 1986AZTREONAMAP FRESENIUS KABI USA 1GM/VIAL A065439 002 Jun 18, 2010AP 2GM/VIAL A065439 003 Jun 18, 2010AP HOSPIRA 1GM/VIAL A206517 001 Nov 08, 2021

PRESCRIPTION DRUG PRODUCT LIST

AZTREONAM

INJECTABLE; INJECTION

AZTREONAM

<u>AP</u>		<u>2GM/VIAL</u>	<u>A206517 002</u>	Nov 08, 2021
	FRESENIUS KABI USA	500MG/VIAL	A065439 001	Jun 18, 2010

BACITRACIN

OINTMENT; OPHTHALMIC

BACITRACIN

!	PADAGIS US	500 UNITS/GM	A061212 001	
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BACITRACIN ZINC; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

!	PADAGIS US	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062166 002	
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BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

!	BAUSCH AND LOMB	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A064068 001	Oct 30, 1995
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BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

LUMI-SPORYN

<u>AT</u>	+	CASPER PHARMA LLC	<u>EQ 400 UNITS/GM;EQ 3.5MG BASE/GM;EQ 10,000 UNITS/GM</u>	<u>N050417 001</u>	
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NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC

<u>AT</u>	!	BAUSCH AND LOMB	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064064 001</u>	Oct 30, 1995
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<u>AT</u>		PADAGIS US	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A060764 002</u>	
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BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE

<u>AT</u>		AKORN	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064028 001</u>	Jan 30, 1995
<u>AT</u>	!	BAUSCH AND LOMB	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064046 001</u>	Jan 26, 1995
<u>AT</u>		PADAGIS US	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A065022 001</u>	Feb 27, 2002

BACLOFEN

GRANULES; ORAL

LYVISPAH

+	AMNEAL	5MG/PACKET	N215422 001	Nov 22, 2021
+		10MG/PACKET	N215422 002	Nov 22, 2021
+	!	20MG/PACKET	N215422 003	Nov 22, 2021

INJECTABLE; INTRATHECAL

BACLOFEN

<u>AP</u>		ACIC PHARMS	<u>0.5MG/ML</u>	<u>A216309 001</u>	Aug 10, 2023
<u>AP</u>			<u>2MG/ML</u>	<u>A216309 002</u>	Aug 10, 2023
<u>AP</u>		AMNEAL	<u>0.05MG/ML</u>	<u>A091193 001</u>	May 03, 2016
<u>AP</u>			<u>0.5MG/ML</u>	<u>A091193 002</u>	May 03, 2016
<u>AP</u>			<u>2MG/ML</u>	<u>A091193 003</u>	May 03, 2016
<u>AP</u>		MAIA PHARMS INC	<u>0.05MG/ML</u>	<u>A210777 001</u>	Jan 15, 2021
<u>AP</u>			<u>0.5MG/ML</u>	<u>A210048 001</u>	Sep 11, 2019
<u>AP</u>			<u>1MG/ML</u>	<u>A210315 001</u>	Jul 30, 2019
<u>AP</u>			<u>2MG/ML</u>	<u>A210048 002</u>	Sep 11, 2019
<u>AP</u>		MYLAN LABS LTD	<u>0.5MG/ML</u>	<u>A209592 001</u>	Mar 21, 2018
<u>AP</u>			<u>1MG/ML</u>	<u>A209594 001</u>	Mar 06, 2018
<u>AP</u>			<u>2MG/ML</u>	<u>A209592 002</u>	Mar 21, 2018
<u>AP</u>		RUBICON	<u>0.5MG/ML</u>	<u>A217324 001</u>	Feb 22, 2023
<u>AP</u>			<u>1MG/ML</u>	<u>A217324 002</u>	Feb 22, 2023
<u>AP</u>			<u>2MG/ML</u>	<u>A217324 003</u>	Feb 22, 2023

GABLOFEN

<u>AP</u>	+	!	PIRAMAL CRITICAL	<u>0.05MG/ML</u>	<u>N022462 001</u>	Nov 19, 2010
<u>AP</u>	+			<u>0.5MG/ML</u>	<u>N022462 002</u>	Nov 19, 2010
<u>AP</u>	+			<u>1MG/ML</u>	<u>N022462 004</u>	Jun 22, 2012
<u>AP</u>	+			<u>2MG/ML</u>	<u>N022462 003</u>	Nov 19, 2010

LIORESAL

<u>AP</u>	+	!	AMNEAL	<u>0.05MG/ML</u>	<u>N020075 003</u>	Nov 07, 1996
<u>AP</u>	+			<u>0.5MG/ML</u>	<u>N020075 001</u>	Jun 17, 1992
<u>AP</u>	+			<u>2MG/ML</u>	<u>N020075 002</u>	Jun 17, 1992

SOLUTION; ORAL

OZOBAX DS

+	!	METACEL PHARMS LLC	10MG/5ML	N208193 002	Oct 12, 2023
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PRESCRIPTION DRUG PRODUCT LIST

BACLOFEN

SUSPENSION; ORAL

BACLOFEN

AB	ANI PHARMS	25MG/5ML	A217252 001	Jun 08, 2023
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FLEQSUVY

AB	+! AZURITY	25MG/5ML	N215602 001	Feb 04, 2022
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TABLET; ORAL

BACLOFEN

AB	AUROBINDO PHARMA LTD	10MG	A214099 001	Jul 13, 2021
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AB		20MG	A214099 002	Jul 13, 2021
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AB	BEXIMCO PHARMS USA	5MG	A214114 003	Sep 19, 2023
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AB		10MG	A214114 001	Jul 16, 2021
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AB		20MG	A214114 002	Jul 16, 2021
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AB	EYWA PHARMA	5MG	A211555 003	Nov 30, 2021
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AB		10MG	A211555 001	Feb 01, 2019
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AB		20MG	A211555 002	Feb 01, 2019
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AB	GRAVITI PHARMS	10MG	A217788 001	Jan 10, 2024
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AB		20MG	A217788 002	Jan 10, 2024
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AB	IMPAX	5MG	A077971 003	Jul 07, 2021
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AB		10MG	A077971 001	Oct 26, 2007
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AB		20MG	A077971 002	Oct 26, 2007
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AB	INNOGENIX	5MG	A212378 003	Apr 30, 2021
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AB		10MG	A212378 001	Oct 09, 2020
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AB		20MG	A212378 002	Oct 09, 2020
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AB	IVAX SUB TEVA PHARMS	10MG	A072234 001	Jul 21, 1988
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AB	!	20MG	A072235 001	Jul 21, 1988
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AB	LANNETT CO INC	5MG	A077241 003	Sep 22, 2021
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AB		10MG	A077241 002	Jul 06, 2007
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AB		20MG	A077241 001	Dec 20, 2005
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AB	MANKIND PHARMA	5MG	A215885 001	Jan 25, 2022
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AB		10MG	A215885 002	Jan 25, 2022
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AB		20MG	A215885 003	Jan 25, 2022
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AB	MICRO LABS	5MG	A217687 001	Nov 08, 2023
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AB		10MG	A217687 002	Nov 08, 2023
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AB		20MG	A217687 003	Nov 08, 2023
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AB	NORTHSTAR HLTHCARE	10MG	A078401 002	Sep 18, 2009
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AB		20MG	A078401 001	Sep 18, 2009
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AB	OXFORD PHARMS	10MG	A077088 002	Oct 31, 2007
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AB		20MG	A077088 001	Oct 31, 2007
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AB	RISING	5MG	A214374 001	Mar 05, 2021
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AB		10MG	A214374 002	Mar 05, 2021
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AB		20MG	A214374 003	Mar 05, 2021
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AB	! RUBICON	5MG	A209102 001	Nov 28, 2017
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AB		10MG	A209102 002	Nov 28, 2017
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AB		20MG	A209102 003	Nov 28, 2017
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AB	UNICHEM	5MG	A212067 003	Mar 09, 2023
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AB		10MG	A212067 001	Jul 09, 2020
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AB		20MG	A212067 002	Jul 09, 2020
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AB	UPSHER SMITH LABS	10MG	A074584 001	Aug 19, 1996
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AB		20MG	A074584 002	Aug 19, 1996
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AB	VINTAGE PHARMS	10MG	A077068 002	Aug 30, 2005
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AB		20MG	A077068 001	Aug 30, 2005
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AB	ZYDUS	5MG	A211659 003	Apr 17, 2020
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AB		10MG	A211659 001	Nov 23, 2018
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AB		20MG	A211659 002	Nov 23, 2018
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BALOXAVIR MARBOXIL

FOR SUSPENSION; ORAL

XOFLUZA

	+! GENENTECH INC	2MG/ML	N214410 001	Nov 23, 2020
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TABLET; ORAL

XOFLUZA

	+ GENENTECH INC	40MG	N210854 002	Oct 24, 2018
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	+!	80MG	N210854 003	Mar 18, 2021
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BALSALAZIDE DISODIUM

CAPSULE; ORAL

BALSALAZIDE DISODIUM

AB	APOTEX INC	750MG	A077883 001	Dec 28, 2007
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AB	HIKMA	750MG	A077806 001	Dec 28, 2007
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AB	ZYDUS	750MG	A217592 001	Jun 08, 2023
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PRESCRIPTION DRUG PRODUCT LIST

BALSALAZIDE DISODIUM

CAPSULE; ORAL

COLAZAL

AB	+ !	VALEANT PHARMS INTL	750MG	N020610 001	Jul 18, 2000
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BARICITINIB

TABLET; ORAL

OLUMIANT

+	ELI LILLY AND CO	1MG	N207924 002	Oct 08, 2019
+		2MG	N207924 001	May 31, 2018
+!		4MG	N207924 003	May 10, 2022

BARIUM SULFATE

FOR SUSPENSION; ORAL

E-Z-HD

+!	BRACCO	98% (334GM/BOT)	N208036 001	Jan 11, 2016
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E-Z-PAQUE

+!	BRACCO	96% (169GM/BOT)	N208036 002	Apr 07, 2017
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VARIBAR THIN LIQUID

+!	BRACCO	81% (120GM/BOT)	N208036 004	Apr 30, 2019
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PASTE; ORAL

VARIBAR PUDDING

	BRACCO	40%	N208844 001	Oct 14, 2016
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SUSPENSION; ORAL

ENTERO VU 24%

+!	BRACCO	24% (144GM/600ML)	N208143 008	May 29, 2020
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LIQUID E-Z-PAQUE

+!	BRACCO	60% (213GM/BOT)	N208143 003	Mar 01, 2017
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READI-CAT 2

+!	BRACCO	2% (9GM/BOT)	N208143 001	Jan 15, 2016
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READI-CAT 2 SMOOTHIE

+!	BRACCO	2% (9GM/BOT)	N208143 002	Jan 15, 2016
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TAGITOL V

+!	BRACCO	40% (8GM/BOT)	N208143 005	Aug 04, 2017
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VARIBAR HONEY

+!	BRACCO	40% (100GM/250ML)	N208143 007	Mar 26, 2018
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VARIBAR NECTAR

+!	BRACCO	40% (96GM/240ML)	N208143 004	Jul 07, 2017
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VARIBAR THIN HONEY

+!	BRACCO	40% (100GM/250ML)	N208143 006	Jan 23, 2018
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BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED

TABLET; ORAL

DUAVEE

+!	WYETH PHARMS	EQ 20MG BASE;0.45MG	N022247 001	Oct 03, 2013
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BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

QVAR REDIHALER

+	NORTON WATERFORD	0.04MG/INH	N207921 001	Aug 03, 2017
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+		0.08MG/INH	N207921 002	Aug 03, 2017
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AEROSOL, METERED; NASAL

QNASL

+	TEVA BRANDED PHARM	0.04MG/ACTUATION	N202813 002	Dec 17, 2014
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+!		0.08MG/ACTUATION	N202813 001	Mar 23, 2012
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BEDAQUILINE FUMARATE

TABLET; ORAL

SIRTURO

+	JANSSEN THERAP	EQ 20MG BASE	N204384 002	May 27, 2020
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+!		EQ 100MG BASE	N204384 001	Dec 28, 2012
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BELINOSTAT

POWDER; INTRAVENOUS

BELEODAQ

+!	ACROTECH BIOPHARMA	500MG/VIAL	N206256 001	Jul 03, 2014
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BELUMOSUDIL MESYLATE

TABLET; ORAL

REZUROCK

+!	KADMON PHARMS LLC	EQ 200MG BASE	N214783 001	Jul 16, 2021
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PRESCRIPTION DRUG PRODUCT LIST

BELZUTIFAN

TABLET; ORAL

WELIREG

+! MERCK SHARP DOHME 40MG N215383 001 Aug 13, 2021

BEMPEDOIC ACID

TABLET; ORAL

NEXLETOL

+! ESPERION THERAPS 180MG N211616 001 Feb 21, 2020
INCBEMPEDOIC ACID; EZETIMIBE

TABLET; ORAL

NEXLIZET

+! ESPERION THERAPS 180MG; 10MG N211617 001 Feb 26, 2020
INCBENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A076820 001</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A076820 002</u>	Feb 03, 2006
<u>AB</u>		<u>20MG</u>	<u>A076820 003</u>	Feb 03, 2006
<u>AB</u>		<u>40MG</u>	<u>A076820 004</u>	Feb 03, 2006
<u>AB</u>	ANI PHARMS	<u>5MG</u>	<u>A076333 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076333 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076333 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076333 004</u>	Feb 11, 2004
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A078212 001</u>	May 22, 2008
<u>AB</u>		<u>20MG</u>	<u>A078212 002</u>	May 22, 2008
<u>AB</u>	!	<u>40MG</u>	<u>A078212 003</u>	May 22, 2008
<u>AB</u>	CADILA	<u>5MG</u>	<u>A078848 001</u>	May 23, 2008
<u>AB</u>		<u>10MG</u>	<u>A078848 002</u>	May 23, 2008
<u>AB</u>		<u>20MG</u>	<u>A078848 003</u>	May 23, 2008
<u>AB</u>		<u>40MG</u>	<u>A078848 004</u>	May 23, 2008
<u>AB</u>	CHARTWELL RX	<u>5MG</u>	<u>A076402 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076402 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076402 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076402 004</u>	Feb 11, 2004
<u>AB</u>	COREPHARMA	<u>5MG</u>	<u>A077128 001</u>	Mar 08, 2006
<u>AB</u>		<u>10MG</u>	<u>A077128 002</u>	Mar 08, 2006
<u>AB</u>		<u>20MG</u>	<u>A077128 003</u>	Mar 08, 2006
<u>AB</u>		<u>40MG</u>	<u>A077128 004</u>	Mar 08, 2006
<u>AB</u>	HERITAGE PHARMA	<u>5MG</u>	<u>A076267 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076267 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076267 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076267 004</u>	Feb 11, 2004
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A076118 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076118 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076118 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076118 004</u>	Feb 11, 2004
<u>AB</u>	SUN PHARM INDS LTD	<u>5MG</u>	<u>A076344 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076344 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076344 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076344 004</u>	Feb 11, 2004
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076211 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076211 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076211 003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076211 004</u>	Feb 11, 2004
	<u>LOTENSIN</u>			
<u>AB</u>	+ VALIDUS PHARMS	<u>10MG</u>	<u>N019851 002</u>	Jun 25, 1991
<u>AB</u>	+	<u>20MG</u>	<u>N019851 003</u>	Jun 25, 1991
<u>AB</u>	+	<u>40MG</u>	<u>N019851 004</u>	Jun 25, 1991

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ANI PHARMS	<u>5MG; 6.25MG</u>	<u>A076342 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076342 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A076342 003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076342 004</u>	Feb 11, 2004
<u>AB</u>	APOTEX	<u>5MG; 6.25MG</u>	<u>A078794 001</u>	Aug 21, 2014
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A078794 002</u>	Aug 21, 2014
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A078794 003</u>	Aug 21, 2014
<u>AB</u>		<u>20MG; 25MG</u>	<u>A078794 004</u>	Aug 21, 2014

PRESCRIPTION DRUG PRODUCT LIST

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	SANDOZ	<u>5MG; 6.25MG</u>	<u>A076631 001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076631 002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A076631 003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076631 004</u>	Feb 11, 2004
<u>LOTENSIN HCT</u>				
<u>AB</u>	+ VALIDUS PHARMS	<u>10MG; 12.5MG</u>	<u>N020033 002</u>	May 19, 1992
<u>AB</u>	+	<u>20MG; 12.5MG</u>	<u>N020033 004</u>	May 19, 1992
<u>AB</u>	+!	<u>20MG; 25MG</u>	<u>N020033 003</u>	May 19, 1992

BENDAMUSTINE HYDROCHLORIDE

POWDER; IV (INFUSION)

BENDAMUSTINE HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>25MG/VIAL</u>	<u>A205574 001</u>	Dec 07, 2022
<u>AP</u>		<u>100MG/VIAL</u>	<u>A205574 002</u>	Dec 07, 2022
<u>AP</u>	APOTEX	<u>25MG/VIAL</u>	<u>A204230 001</u>	Jun 05, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A204230 002</u>	Jun 05, 2023
<u>AP</u>	BRECKENRIDGE	<u>25MG/VIAL</u>	<u>A205447 001</u>	Feb 14, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A205447 002</u>	Feb 14, 2023
<u>AP</u>	DR REDDYS	<u>25MG/VIAL</u>	<u>A205376 001</u>	Dec 07, 2022
<u>AP</u>		<u>100MG/VIAL</u>	<u>A205376 002</u>	Dec 07, 2022
<u>AP</u>	EUGIA PHARMA	<u>25MG/VIAL</u>	<u>A214739 001</u>	Jun 05, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A214739 002</u>	Jun 05, 2023
<u>AP</u>	KINDOS	<u>25MG/VIAL</u>	<u>A211001 001</u>	Jun 05, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A211001 002</u>	Jun 05, 2023
<u>TREANDA</u>				
<u>AP</u>	+! CEPHALON	<u>25MG/VIAL</u>	<u>N022249 002</u>	May 01, 2009
<u>AP</u>	+!	<u>100MG/VIAL</u>	<u>N022249 001</u>	Mar 20, 2008

SOLUTION; INTRAVENOUS

BELRAPZO

<u>AP</u>	+! EAGLE PHARMS	<u>100MG/4ML (25MG/ML)</u>	<u>N205580 001</u>	May 15, 2018
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BENDAMUSTINE HYDROCHLORIDE

<u>AP</u>	+! APOTEX	<u>100MG/4ML (25MG/ML)</u>	<u>N215033 001</u>	Dec 07, 2022
<u>AP</u>	+! BAXTER HLTHCARE CORP	<u>100MG/4ML (25MG/ML)</u>	<u>N216078 001</u>	Dec 15, 2022

VIVIMUSTA

+!	SLAYBACK PHARMA LLC	100MG/4ML (25MG/ML)	N212209 001	Dec 07, 2022
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SOLUTION; IV (INFUSION)

BENDEKA

+!	EAGLE PHARMS	100MG/4ML (25MG/ML)	N208194 001	Dec 07, 2015
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BENOXINATE HYDROCHLORIDE; FLUORESCEIN SODIUM

SOLUTION/DROPS; OPHTHALMIC

ALTAFLUOR BENOX

+!	ALTAIRE PHARMS INC	0.4%; 0.25%	N208582 001	Dec 14, 2017
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FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE

+!	BAUSCH LOMB IRELAND	0.4%; 0.3%	N211039 001	Mar 09, 2020
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BENZNIDAZOLE

TABLET; ORAL

BENZNIDAZOLE

+	CHEMO RESEARCH SL	12.5MG	N209570 001	Aug 29, 2017
+!		100MG	N209570 002	Aug 29, 2017

BENZONATATE

CAPSULE; ORAL

BENZONATATE

<u>AA</u>	ACELLA	<u>100MG</u>	<u>A091310 001</u>	Jan 16, 2015
<u>AA</u>		<u>200MG</u>	<u>A091310 002</u>	Jan 16, 2015
<u>AA</u>	ASCENT PHARMS INC	<u>100MG</u>	<u>A211518 001</u>	Feb 22, 2019
<u>AA</u>		<u>200MG</u>	<u>A211518 003</u>	Feb 22, 2019
<u>AA</u>	BIONPHARMA	<u>100MG</u>	<u>A081297 001</u>	Jan 29, 1993
<u>AA</u>		<u>200MG</u>	<u>A081297 002</u>	Oct 30, 2007
<u>AA</u>	CSPC-NBP PHARM	<u>100MG</u>	<u>A202765 002</u>	Aug 25, 2017
<u>AA</u>		<u>200MG</u>	<u>A202765 001</u>	Jul 31, 2015
<u>AA</u>	HERITAGE PHARMS LABS	<u>100MG</u>	<u>A040682 001</u>	Jul 30, 2007
<u>AA</u>		<u>200MG</u>	<u>A040682 002</u>	Jul 30, 2007
<u>AA</u>	! PURACAP PHARM LLC	<u>100MG</u>	<u>A206948 001</u>	Dec 19, 2018
<u>AA</u>	!	<u>200MG</u>	<u>A206948 002</u>	Dec 19, 2018
<u>AA</u>	STRIDES PHARMA	<u>100MG</u>	<u>A091133 001</u>	Jul 30, 2015
<u>AA</u>		<u>200MG</u>	<u>A091133 002</u>	Jul 30, 2015
<u>AA</u>	ZYDUS PHARMS USA	<u>100MG</u>	<u>A040597 001</u>	Jun 08, 2007

PRESCRIPTION DRUG PRODUCT LIST

BENZONATATE

CAPSULE;ORAL

BENZONATATE

AA		200MG	A040597 002	Jun 08, 2007
	<u>TESSALON</u>			
AA	+ PFIZER	100MG	N011210 001	
	BENZONATATE			
	ASCENT PHARMS INC	150MG	A211518 002	Feb 22, 2019

BENZOYL PEROXIDE

CREAM;TOPICAL

EPSOLAY

+!	GALDERMA LABS LP	5%	N214510 001	Apr 22, 2022
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BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL;TOPICAL

ACANYA

AB	+!	BAUSCH	2.5%;EQ 1.2% BASE	N050819 001	Oct 23, 2008
		<u>CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE</u>			
AB		ACTAVIS LABS UT INC	2.5%;EQ 1.2% BASE	A205128 001	Jun 19, 2015
AB		ENCUBE	5%;1.2%	A212433 001	Apr 28, 2021
AB	!	GLENMARK PHARMS	5%;EQ 1% BASE	A209252 001	Mar 14, 2019
AB		MYLAN PHARMS INC	5%;EQ 1% BASE	A065443 001	Aug 11, 2009
AB		PADAGIS ISRAEL	2.5%;EQ 1.2% BASE	A205397 001	Sep 09, 2019
AB			5%;EQ 1% BASE	A202440 001	Sep 21, 2015
AB	!		5%;1.2%	A090979 001	Jun 26, 2012
AB		TARO	2.5%;EQ 1.2% BASE	A206575 001	Aug 19, 2019
AB			3.75%;EQ 1.2% BASE	A208683 001	Jun 05, 2018
AB			5%;EQ 1% BASE	A208776 001	May 25, 2018
AB			5%;1.2%	A206218 001	Dec 15, 2017
AB		ZYDUS PHARMS	5%;1.2%	A210794 001	Dec 28, 2018
		<u>DUAC</u>			
AB	+	STIEFEL	5%;1.2%	N050741 001	Aug 26, 2002
		<u>ONEXTON</u>			
AB	+!	BAUSCH	3.75%;EQ 1.2% BASE	N050819 002	Nov 24, 2014

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL;TOPICAL

BENZAMYCIN

AB	+!	VALEANT INTL	5%;3%	N050557 001	Oct 26, 1984
		<u>ERYTHROMYCIN AND BENZOYL PEROXIDE</u>			
AB		LYNE	5%;3%	A065385 001	Sep 18, 2015

BENZOYL PEROXIDE; TRETINOIN

CREAM;TOPICAL

TWYNEO

+!	GALDERMA LABS LP	3%;0.1%	N214902 001	Jul 26, 2021
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BENZPHETAMINE HYDROCHLORIDE

TABLET;ORAL

BENZPHETAMINE HYDROCHLORIDE

AA		CHARTWELL	50MG	A090473 002	Sep 15, 2010
AA		EPIC PHARMA LLC	50MG	A090346 001	Dec 15, 2015
AA	!	KVK TECH	50MG	A090968 001	Jul 20, 2010
		CHARTWELL	25MG	A090473 001	Sep 15, 2010

BENZTROPINE MESYLATE

INJECTABLE;INJECTION

BENZTROPINE MESYLATE

AP		FRESENIUS KABI USA	1MG/ML	A090233 001	Jul 28, 2009
AP		HIKMA	1MG/ML	A209442 001	Oct 14, 2021
AP	!	HIKMA FARMACEUTICA	1MG/ML	A090287 001	Aug 31, 2009
AP		NAVINTA LLC	1MG/ML	A091525 001	Feb 05, 2013

TABLET;ORAL

BENZTROPINE MESYLATE

AA	!	ASPEN GLOBAL INC	0.5MG	A204713 001	Apr 14, 2015
AA	!		1MG	A204713 002	Apr 14, 2015
AA	!		2MG	A204713 003	Apr 14, 2015
AA		CHARTWELL RX	0.5MG	A081265 003	Nov 29, 2023
AA			1MG	A081265 002	Jan 23, 1992
AA			2MG	A081265 001	Jan 23, 1992
AA		EPIC PHARMA LLC	0.5MG	A072264 001	Feb 27, 1989
AA			1MG	A072265 001	Feb 27, 1989
AA			2MG	A072266 001	Feb 27, 1989
AA		INVAGEN PHARMS	0.5MG	A090294 001	Mar 29, 2010

PRESCRIPTION DRUG PRODUCT LIST

BENZTROPINE MESYLATE

TABLET; ORAL

BENZTROPINE MESYLATE

<u>AA</u>		<u>1MG</u>	<u>A090294 002</u>	Mar 29, 2010
<u>AA</u>		<u>2MG</u>	<u>A090294 003</u>	Mar 29, 2010
<u>AA</u>	LEADING	<u>0.5MG</u>	<u>A090168 001</u>	Nov 28, 2012
<u>AA</u>		<u>1MG</u>	<u>A090168 002</u>	Nov 28, 2012
<u>AA</u>		<u>2MG</u>	<u>A090168 003</u>	Nov 28, 2012
<u>AA</u>	PLIVA	<u>0.5MG</u>	<u>A089058 001</u>	Aug 10, 1988
<u>AA</u>		<u>1MG</u>	<u>A089059 001</u>	Aug 10, 1988
<u>AA</u>		<u>2MG</u>	<u>A089060 001</u>	Aug 10, 1988
<u>AA</u>	VINTAGE	<u>0.5MG</u>	<u>A040715 001</u>	Aug 27, 2007
<u>AA</u>		<u>1MG</u>	<u>A040715 002</u>	Aug 27, 2007
<u>AA</u>		<u>2MG</u>	<u>A040715 003</u>	Aug 27, 2007

BEPOTASTINE BESILATE

SOLUTION/DROPS; OPHTHALMIC

BEPOTASTINE BESILATE

<u>AT</u>	ALEMBIC	<u>1.5%</u>	<u>A214588 001</u>	Apr 05, 2023
<u>AT</u>	APOTEX	<u>1.5%</u>	<u>A206066 001</u>	Mar 05, 2019
<u>AT</u>	MYLAN	<u>1.5%</u>	<u>A206220 001</u>	Mar 18, 2019

BEPREVE

<u>AT</u>	+!	BAUSCH AND LOMB INC	<u>1.5%</u>	<u>N022288 001</u>	Sep 08, 2009
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BEROTRALSTAT HYDROCHLORIDE

CAPSULE; ORAL

ORLADEYO

+	BIOCRYST	EQ 110MG BASE	N214094 001	Dec 03, 2020
+	!	EQ 150MG BASE	N214094 002	Dec 03, 2020

BESIFLOXACIN HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BESIVANCE

+	!	BAUSCH AND LOMB	EQ 0.6% BASE	N022308 001	May 28, 2009
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BETAINE

FOR SOLUTION; ORAL

BETAINE

<u>AB</u>	ETON	<u>1GM/SCOOPFUL</u>	<u>A210508 001</u>	Jan 28, 2022	
<u>AB</u>	NOVITIUM PHARMA	<u>1GM/SCOOPFUL</u>	<u>A214864 001</u>	Nov 23, 2021	
<u>AB</u>	+!	RECORDATI RARE	<u>1GM/SCOOPFUL</u>	<u>N020576 001</u>	Oct 25, 1996

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE

<u>AB</u>	AM REGENT	<u>3MG/ML;EQ 3MG BASE/ML</u>	<u>A090747 001</u>	Jul 31, 2009
<u>AB</u>	HIKMA	<u>3MG/ML;EQ 3MG BASE/ML</u>	<u>A077838 001</u>	Jan 17, 2023

CELESTONE SOLUSPAN

<u>AB</u>	+!	ORGANON	<u>3MG/ML;EQ 3MG BASE/ML</u>	<u>N014602 001</u>
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BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ACTAVIS MID	<u>EQ 0.05% BASE</u>	<u>A070885 001</u>	Feb 03, 1987	
	ATLANTIC				
<u>AB</u>	COSETTE	<u>EQ 0.05% BASE</u>	<u>A210217 001</u>	Oct 12, 2018	
<u>AB</u>	+!	FOUGERA PHARMS	<u>EQ 0.05% BASE</u>	<u>N019137 001</u>	Jun 26, 1984
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A073552 001</u>	Apr 30, 1992	
<u>AB</u>	ZYDUS PHARMS	<u>EQ 0.05% BASE</u>	<u>A208885 001</u>	Jan 11, 2019	

CREAM, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	FOUGERA PHARMS	<u>EQ 0.05% BASE</u>	<u>A076215 001</u>	Dec 09, 2003	
<u>AB</u>	!	GLENMARK GENERICS	<u>EQ 0.05% BASE</u>	<u>A078930 001</u>	Sep 23, 2008
<u>AB</u>	PADAGIS ISRAEL	<u>EQ 0.05% BASE</u>	<u>A076592 001</u>	Dec 09, 2003	
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A076543 001</u>	Dec 09, 2003	

GEL, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	!	FOUGERA PHARMS	<u>EQ 0.05% BASE</u>	<u>A075276 001</u>	May 13, 2003
<u>AB</u>		TARO	<u>EQ 0.05% BASE</u>	<u>A076508 001</u>	Dec 02, 2003

LOTION; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	COSETTE	<u>EQ 0.05% BASE</u>	<u>A071467 001</u>	Aug 10, 1987	
<u>AB</u>	!	FOUGERA PHARMS INC	<u>EQ 0.05% BASE</u>	<u>A070275 001</u>	Aug 12, 1985
<u>AB</u>		PADAGIS US	<u>EQ 0.05% BASE</u>	<u>A072538 001</u>	Jan 31, 1990

PRESCRIPTION DRUG PRODUCT LIST

BETAMETHASONE DIPROPIONATE

LOTION, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

AB	FOUGERA PHARMS	EQ 0.05% BASE	A077111 001	May 21, 2007
AB	HIKMA	EQ 0.05% BASE	A208849 001	Oct 11, 2019
AB	! TARO	EQ 0.05% BASE	A077477 001	May 21, 2007

OINTMENT;TOPICAL

BETAMETHASONE DIPROPIONATE

AB	ACTAVIS MID	EQ 0.05% BASE	A071012 001	Feb 03, 1987
	ATLANTIC			
AB	CADILA	EQ 0.05% BASE	A214048 001	Jul 14, 2020
AB	+! FOUGERA PHARMS INC	EQ 0.05% BASE	N019141 001	Sep 04, 1984
AB	PADAGIS ISRAEL	EQ 0.05% BASE	A215847 001	Apr 12, 2022
AB	TARO	EQ 0.05% BASE	A074271 001	Sep 15, 1994
AB	TASMAN PHARMA	EQ 0.05% BASE	A215186 001	Feb 18, 2022

OINTMENT, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

AB	ACTAVIS MID	EQ 0.05% BASE	A074304 001	Aug 31, 1995
	ATLANTIC			
AB	FOUGERA PHARMS	EQ 0.05% BASE	A075373 001	Jun 22, 1999
AB	LUPIN LTD	EQ 0.05% BASE	A209106 001	Dec 18, 2019
AB	TARO	EQ 0.05% BASE	A076753 001	Oct 12, 2004

DIPROLENE

AB	+! ORGANON	EQ 0.05% BASE	N018741 001	Jul 27, 1983
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SPRAY;TOPICAL

SERNIVO

+!	PRIMUS PHARMS	EQ 0.05% BASE/SPRAY	N208079 001	Feb 05, 2016
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BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE

AEROSOL, FOAM;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE

AB	GLENMARK PHARMS LTD	0.064%;0.005%	A214688 001	Mar 21, 2023
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ENSTILAR

AB	+! LEO PHARMA AS	0.064%;0.005%	N207589 001	Oct 16, 2015
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CREAM;TOPICAL

WYNZORA

+!	MC2	0.064%;0.005%	N213422 001	Jul 20, 2020
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OINTMENT;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE

AB	PADAGIS ISRAEL	0.064%;0.005%	A200174 001	Dec 12, 2014
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TACLONEX

AB	+! LEO PHARMA AS	0.064%;0.005%	N021852 001	Jan 09, 2006
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SUSPENSION;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE

AB	COSETTE	0.064%;0.005%	A210765 001	May 11, 2020
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AB	TARO	0.064%;0.005%	A213269 001	Sep 02, 2020
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CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE

AB	PADAGIS ISRAEL	0.064%;0.005%	A212367 001	Sep 11, 2020
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TACLONEX

AB	+! LEO PHARMA AS	0.064%;0.005%	N022185 001	May 09, 2008
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BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM;TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

AB	ACTAVIS MID	EQ 0.05% BASE;1%	A076002 001	Aug 02, 2002
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	ATLANTIC			
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AB	FOUGERA PHARMS	EQ 0.05% BASE;1%	A075502 001	Jun 05, 2001
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AB	GLENMARK PHARMS	EQ 0.05% BASE;1%	A202894 001	Oct 30, 2015
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AB	! TARO	EQ 0.05% BASE;1%	A075673 001	May 29, 2001
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LOTION;TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

AB	FOUGERA PHARMS	EQ 0.05% BASE;1%	A076516 001	Jun 16, 2005
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AB	! TARO	EQ 0.05% BASE;1%	A076493 001	Jul 28, 2004
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BETAMETHASONE VALERATE

AEROSOL, FOAM;TOPICAL

BETAMETHASONE VALERATE

AB	PADAGIS ISRAEL	0.12%	A078337 001	Nov 26, 2012
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AB	! TARO	0.12%	A208204 001	May 24, 2017
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AB	XIROMED	0.12%	A210639 001	Apr 18, 2023
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CREAM;TOPICAL

BETA-VAL

AB	COSETTE	EQ 0.1% BASE	N018642 001	Mar 24, 1983
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PRESCRIPTION DRUG PRODUCT LIST

BETAMETHASONE VALERATE

CREAM; TOPICAL

BETAMETHASONE VALERATE

AB	+!	FOUGERA PHARMS INC	EQ 0.1% BASE	N018861 001	Aug 31, 1983
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DERMABET

AB		TARO	EQ 0.1% BASE	A072041 001	Jan 06, 1988
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VALNAC

AB		ACTAVIS MID ATLANTIC	EQ 0.1% BASE	A070050 001	Oct 10, 1984
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LOTION; TOPICAL

BETAMETHASONE VALERATE

AB		ANIMA	EQ 0.1% BASE	A070052 001	Jul 31, 1985
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AB	+!	FOUGERA PHARMS INC	EQ 0.1% BASE	N018866 001	Aug 31, 1983
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OINTMENT; TOPICAL

BETA-VAL

AB		COSETTE	EQ 0.1% BASE	A070069 001	Dec 19, 1985
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BETAMETHASONE VALERATE

AB		ACTAVIS MID ATLANTIC	EQ 0.1% BASE	A070051 001	Oct 10, 1984
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AB	+!	FOUGERA PHARMS INC	EQ 0.1% BASE	N018865 001	Aug 31, 1983
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BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

AT		ACELLA	EQ 0.5% BASE	A078694 001	Nov 16, 2009
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AT		MEDIMETRIKS PHARMS	EQ 0.5% BASE	A075630 001	Apr 12, 2001
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BETOPTIC

AT	+!	SANDOZ	EQ 0.5% BASE	N019270 001	Aug 30, 1985
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SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC S**+!** NOVARTIS

EQ 0.25% BASE

N019845 001 Dec 29, 1989

TABLET; ORAL

BETAXOLOL HYDROCHLORIDE

AB		EPIC PHARMA	10MG	A075541 001	Oct 22, 1999
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AB	!		20MG	A075541 002	Oct 22, 1999
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AB		KVK TECH	10MG	A078962 001	Jun 27, 2008
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AB			20MG	A078962 002	Jun 27, 2008
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BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

AA	!	AMNEAL PHARM	5MG	A040855 001	Nov 21, 2007
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AA	!		10MG	A040855 002	Nov 21, 2007
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AA	!		25MG	A040855 003	Nov 21, 2007
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AA	!		50MG	A040855 004	Nov 21, 2007
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AA		CHARTWELL RX	5MG	A040728 002	Oct 26, 2007
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AA			10MG	A040728 003	Oct 26, 2007
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AA			25MG	A040728 004	Oct 26, 2007
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AA			50MG	A040728 001	Oct 26, 2007
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AA		UPSHER SMITH LABS	5MG	A040633 001	Jun 01, 2005
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AA			10MG	A040634 001	Jun 01, 2005
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AA			25MG	A040635 001	Jun 01, 2005
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AA			50MG	A040636 001	Jun 01, 2005
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DUVOID

AA		CHARTWELL RX	10MG	A086262 001	
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AA			25MG	A086263 001	
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AA			50MG	A085882 003	
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BEXAGLIFLOZIN

TABLET; ORAL

BRENZAVVY**+!** THERACOSBIO

20MG

N214373 001 Jan 20, 2023

BEXAROTENE

CAPSULE; ORAL

BEXAROTENE

AB		AMNEAL PHARMS NY	75MG	A210105 001	Sep 04, 2018
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AB		ANI PHARMS	75MG	A209861 001	May 08, 2018
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AB		BIONPHARMA	75MG	A203174 001	Aug 12, 2014
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AB		HIKMA	75MG	A203663 001	Jun 16, 2020
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AB		TEVA PHARMS USA	75MG	A209931 001	Jan 14, 2021
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AB		UPSHER SMITH LABS	75MG	A209886 001	Jul 25, 2018
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TARGRETIN

AB	+!	VALEANT LUXEMBOURG	75MG	N021055 001	Dec 29, 1999
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PRESCRIPTION DRUG PRODUCT LISTBEXAROTENE

GEL; TOPICAL

BEXAROTENE

AB	AMNEAL	1%	A215398 001	Apr 27, 2022
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TARGRETIN

AB	+! BAUSCH	1%	N021056 001	Jun 28, 2000
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BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

AB	ACCORD HLTHCARE	50MG	A078917 001	Jul 06, 2009
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AB	ADAPTIS	50MG	A079089 001	Jul 06, 2009
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AB	APOTEX	50MG	A200274 001	May 21, 2015
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AB	BRECKENRIDGE	50MG	A091011 001	Jun 10, 2015
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AB	SANDOZ	50MG	A078575 001	Jul 06, 2009
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AB	SUN PHARM	50MG	A079110 001	Jul 06, 2009
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AB	WATSON LABS TEVA	50MG	A078634 001	Aug 28, 2009
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CASODEX

AB	+! ANI PHARMS	50MG	N020498 001	Oct 04, 1995
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BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

BIKTARVY

+	GILEAD SCIENCES INC	EQ 30MG BASE;120MG;EQ 15MG BASE	N210251 002	Oct 07, 2021
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+	!	EQ 50MG BASE;200MG;EQ 25MG BASE	N210251 001	Feb 07, 2018
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BIMATOPROST

IMPLANT; OPHTHALMIC

DURYSTA

+	!	ABBVIE	10MCG	N211911 001	Mar 04, 2020
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SOLUTION/DROPS; OPHTHALMIC

BIMATOPROST

AT	!	ALEMBIC	0.03%	A210263 001	Apr 12, 2019
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AT		APOTEX	0.03%	A090449 001	Jul 20, 2015
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AT		EUGIA PHARMA	0.03%	A205537 001	Oct 06, 2022
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AT		GLAND PHARMA LTD	0.03%	A210126 001	Mar 22, 2019
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AT		LUPIN LTD	0.03%	A203991 001	Feb 20, 2015
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AT		MICRO LABS	0.03%	A202505 001	Sep 08, 2020
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AT		SANDOZ	0.03%	A202565 001	May 05, 2015
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AT		SOMERSET THERAPS	0.03%	A207601 001	Jun 19, 2019
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LLC

LUMIGAN

+	!	ABBVIE	0.01%	N022184 001	Aug 31, 2010
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SOLUTION/DROPS; TOPICAL

BIMATOPROST

AT		ALEMBIC	0.03%	A210515 001	Jan 21, 2020
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AT		APOTEX	0.03%	A201894 001	Dec 01, 2014
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AT		HIKMA	0.03%	A203051 001	Oct 09, 2018
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AT		SANDOZ	0.03%	A202719 001	Apr 19, 2016
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LATISSE

AT	+	!	ABBVIE	0.03%	N022369 001	Dec 24, 2008
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BINIMETINIB

TABLET; ORAL

MEKTOVI

+	!	ARRAY BIOPHARMA INC	15MG	N210498 001	Jun 27, 2018
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BIRCH TRITERPENES

GEL; TOPICAL

FILSUVEZ

+	!	AMRYT	10%	N215064 001	Dec 18, 2023
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BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE

AB	PAR PHARM	140MG;125MG;125MG	A205770 001	Mar 06, 2023
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AB	RICONPHARMA LLC	140MG;125MG;125MG	A217511 001	Jul 03, 2023
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PYLERA

AB	+	!	LABS JUVISE	140MG;125MG;125MG	N050786 001	Sep 28, 2006
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PRESCRIPTION DRUG PRODUCT LIST

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE; ORAL

BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE

!	NOSTRUM LABS INC	262.4MG, N/A, N/A; N/A, 250MG, N/A; N/A, N/A, 500MG	A202584	001	Nov 30, 2018
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BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

AB	ALEMBIC	5MG	A204891	001	Jan 11, 2017
AB		10MG	A204891	002	Jan 11, 2017
AB	AUROBINDO PHARMA	5MG	A077910	001	Dec 27, 2006
AB		10MG	A077910	002	Dec 27, 2006
AB	CADILA	5MG	A215680	001	Jul 25, 2022
AB		10MG	A215680	002	Jul 25, 2022
AB	NOVITIUM PHARMA	5MG	A215563	001	Oct 29, 2021
AB		10MG	A215563	002	Oct 29, 2021
AB	PRINSTON INC	5MG	A217368	001	Jul 14, 2023
AB		10MG	A217368	002	Jul 14, 2023
AB	RUBICON	5MG	A075643	001	Nov 16, 2000
AB		10MG	A075643	002	Nov 16, 2000
AB	UNICHEM	5MG	A078635	001	Aug 18, 2009
AB	!	10MG	A078635	002	Aug 18, 2009
AB	UNITED RES LABS	5MG	A075474	001	Oct 25, 2002
AB		10MG	A075474	002	Oct 25, 2002

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

AB	CADILA	2.5MG; 6.25MG	A215666	001	Nov 04, 2022
AB		5MG; 6.25MG	A215666	002	Nov 04, 2022
AB		10MG; 6.25MG	A215666	003	Nov 04, 2022
AB	EDENBRIDGE PHARMS	2.5MG; 6.25MG	A212678	001	Jul 09, 2020
AB		5MG; 6.25MG	A212678	002	Jul 09, 2020
AB		10MG; 6.25MG	A212678	003	Jul 09, 2020
AB	EPIC PHARMA LLC	2.5MG; 6.25MG	A075579	001	Sep 25, 2000
AB		5MG; 6.25MG	A075579	002	Sep 25, 2000
AB		10MG; 6.25MG	A075579	003	Sep 25, 2000
AB	GLENMARK PHARMS LTD	2.5MG; 6.25MG	A215995	001	Jan 26, 2022
AB		5MG; 6.25MG	A215995	002	Jan 26, 2022
AB		10MG; 6.25MG	A215995	003	Jan 26, 2022
AB	MYLAN	2.5MG; 6.25MG	A075768	001	Sep 25, 2000
AB		5MG; 6.25MG	A075768	002	Sep 25, 2000
AB		10MG; 6.25MG	A075768	003	Sep 25, 2000
AB	NOVITIUM PHARMA	2.5MG; 6.25MG	A215562	001	Nov 04, 2021
AB		5MG; 6.25MG	A215562	002	Nov 04, 2021
AB		10MG; 6.25MG	A215562	003	Nov 04, 2021
AB	UNICHEM	2.5MG; 6.25MG	A079106	001	Jul 28, 2010
AB		5MG; 6.25MG	A079106	002	Jul 28, 2010
AB		10MG; 6.25MG	A079106	003	Jul 28, 2010
ZIAC					
AB	+	TEVA BRANDED PHARM	2.5MG; 6.25MG	N020186	003 Mar 26, 1993
AB	+		5MG; 6.25MG	N020186	001 Mar 26, 1993
AB	+	!	10MG; 6.25MG	N020186	002 Mar 26, 1993

BIVALIRUDIN

INJECTABLE; INTRAVENOUS

ANGIOMAX

AP	+	SANDOZ	250MG/VIAL	N020873	001 Dec 15, 2000
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BIVALIRUDIN

AP		ACCORD HLTHCARE	250MG/VIAL	A206551	001 Nov 22, 2017
AP		DR REDDYS	250MG/VIAL	A201577	001 May 26, 2017
AP		EUGIA PHARMA	250MG/VIAL	A205962	001 Jul 27, 2018
AP		FRESENIUS KABI USA	250MG/VIAL	A090189	001 Oct 28, 2016
AP		HOSPIRA	250MG/VIAL	A090811	001 Jul 14, 2015
AP			250MG/VIAL	A090816	001 Jul 14, 2015
AP		MEITHEAL	250MG/VIAL	A091602	001 Jul 16, 2018
AP		MYLAN INSTITUTIONAL	250MG/VIAL	A202471	001 Jun 01, 2018
AP		SHUANGCHENG	250MG/VIAL	A210031	001 Oct 23, 2019
AP		SLATE RUN PHARMA	250MG/VIAL	A213078	001 May 28, 2021

SOLUTION; INTRAVENOUS

ANGIOMAX RTU

+	!	MAIA PHARMS INC	250MG/50ML (5MG/ML)	N211215	001 Jul 25, 2019
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PRESCRIPTION DRUG PRODUCT LIST

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLEOMYCIN SULFATE

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065185 001</u>	Jan 28, 2008
<u>AP</u>	!		<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065185 002</u>	Jan 28, 2008
<u>AP</u>		HIKMA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065042 002</u>	Oct 17, 2001
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065042 001</u>	Oct 17, 2001
<u>AP</u>		HOSPIRA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065031 001</u>	Mar 10, 2000
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065031 002</u>	Mar 10, 2000
<u>AP</u>		MEITHEAL	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A205030 001</u>	Apr 20, 2018
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>A205030 002</u>	Apr 20, 2018
<u>AP</u>		TEVA PHARMS USA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065033 001</u>	Jun 27, 2000
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065033 002</u>	Jun 27, 2000

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

<u>AP</u>		APOTEX	<u>3.5MG/VIAL</u>	<u>A205533 001</u>	May 02, 2022
<u>AP</u>		BAXTER HLTHCARE CORP	<u>3.5MG/VIAL</u>	<u>A213823 001</u>	May 02, 2022
<u>AP</u>		DR REDDYS	<u>3.5MG/VIAL</u>	<u>A202963 001</u>	Jul 26, 2022
<u>AP</u>		EUGIA PHARMA	<u>3.5MG/VIAL</u>	<u>A212825 001</u>	May 02, 2022
<u>AP</u>		FRESENIUS KABI USA	<u>3.5MG/VIAL</u>	<u>A209659 001</u>	May 02, 2022
<u>AP</u>		HOSPIRA	<u>3.5MG/VIAL</u>	<u>A208460 001</u>	Jul 26, 2022
<u>AP</u>		JIANGSU HANSOH PHARM	<u>3.5MG/VIAL</u>	<u>A215011 001</u>	Jul 26, 2022
<u>AP</u>		MEITHEAL	<u>3.5MG/VIAL</u>	<u>A212958 001</u>	Jul 26, 2022
<u>AP</u>		MSN	<u>3.5MG/VIAL</u>	<u>A209622 001</u>	Jul 26, 2022
<u>AP</u>		PHARMASCIENCE INC	<u>3.5MG/VIAL</u>	<u>A208392 001</u>	May 02, 2022
<u>AP</u>		QILU PHARM HAINAN	<u>3.5MG/VIAL</u>	<u>A210824 001</u>	May 02, 2022
<u>AP</u>		SANDOZ	<u>3.5MG/VIAL</u>	<u>A203654 001</u>	Jul 26, 2022
<u>AP</u>		WAVERLEY PHARMA INC	<u>3.5MG/VIAL</u>	<u>A211898 001</u>	Oct 11, 2022
<u>AP</u>		ZYDUS PHARMS	<u>3.5MG/VIAL</u>	<u>A210204 001</u>	May 02, 2022

VELCADE

<u>AP</u>	+	TAKEDA PHARMS USA	<u>3.5MG/VIAL</u>	<u>N021602 001</u>	May 13, 2003
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POWDER; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

+	!	HOSPIRA	1MG/VIAL	N209191 001	May 02, 2022
+	!		2.5MG/VIAL	N209191 002	May 02, 2022

BOSENTAN

TABLET; ORAL

BOSENTAN

<u>AB</u>		SUN PHARM	<u>62.5MG</u>	<u>A209324 001</u>	Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A209324 002</u>	Apr 26, 2019
<u>AB</u>		WATSON LABS INC	<u>62.5MG</u>	<u>A207110 001</u>	Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A207110 002</u>	Apr 26, 2019
<u>AB</u>		ZYDUS PHARMS	<u>62.5MG</u>	<u>A207760 001</u>	Apr 26, 2019
<u>AB</u>			<u>125MG</u>	<u>A207760 002</u>	Apr 26, 2019

TRACLEER

<u>AB</u>	+	ACTELION	<u>62.5MG</u>	<u>N021290 001</u>	Nov 20, 2001
<u>AB</u>	+	!	<u>125MG</u>	<u>N021290 002</u>	Nov 20, 2001

TABLET, FOR SUSPENSION; ORAL

TRACLEER

+	!	ACTELION	32MG	N209279 001	Sep 05, 2017
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BOSUTINIB MONOHYDRATE

CAPSULE; ORAL

BOSULIF

+		PF PRISM CV	EQ 50MG BASE	N217729 001	Sep 26, 2023
+	!		EQ 100MG BASE	N217729 002	Sep 26, 2023

TABLET; ORAL

BOSULIF

+	!	PF PRISM CV	EQ 100MG BASE	N203341 001	Sep 04, 2012
+			EQ 400MG BASE	N203341 003	Oct 27, 2017
+			EQ 500MG BASE	N203341 002	Sep 04, 2012

BREMELANOTIDE ACETATE

SOLUTION; SUBCUTANEOUS

VYLEESI (AUTOINJECTOR)

+	!	PALATIN TECHNOLOGIES	EQ 1.75MG BASE/0.3ML (EQ 1.75MG BASE/0.3 ML)	N210557 001	Jun 21, 2019
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PRESCRIPTION DRUG PRODUCT LIST

BREXANOLONESOLUTION; INTRAVENOUS
ZULRESSO

+! SAGE THERAP 100MG/20ML (5MG/ML) N211371 001 Jun 17, 2019

BREXPIPIRAZOLE

TABLET; ORAL

BREXPIPIRAZOLE

<u>AB</u>	ALKEM LABS LTD	<u>0.25MG</u>	<u>A213782 001</u>	Nov 28, 2023
<u>AB</u>		<u>0.5MG</u>	<u>A213782 002</u>	Nov 28, 2023
<u>AB</u>		<u>1MG</u>	<u>A213782 003</u>	Nov 28, 2023
<u>AB</u>		<u>2MG</u>	<u>A213782 004</u>	Nov 28, 2023
<u>AB</u>		<u>3MG</u>	<u>A213782 005</u>	Nov 28, 2023
<u>AB</u>		<u>4MG</u>	<u>A213782 006</u>	Nov 28, 2023
<u>AB</u>	HETERO LABS LTD V	<u>0.25MG</u>	<u>A213669 001</u>	Nov 20, 2023
<u>AB</u>		<u>0.5MG</u>	<u>A213669 002</u>	Nov 20, 2023
<u>AB</u>		<u>1MG</u>	<u>A213669 003</u>	Nov 20, 2023
<u>AB</u>		<u>2MG</u>	<u>A213669 004</u>	Nov 20, 2023
<u>AB</u>		<u>3MG</u>	<u>A213669 005</u>	Nov 20, 2023
<u>AB</u>		<u>4MG</u>	<u>A213669 006</u>	Nov 20, 2023
<u>AB</u>	OPTIMUS	<u>0.25MG</u>	<u>A213758 001</u>	Oct 25, 2023
<u>AB</u>		<u>0.5MG</u>	<u>A213758 002</u>	Oct 25, 2023
<u>AB</u>		<u>1MG</u>	<u>A213758 003</u>	Oct 25, 2023
<u>AB</u>		<u>2MG</u>	<u>A213758 004</u>	Oct 25, 2023
<u>AB</u>		<u>3MG</u>	<u>A213758 005</u>	Oct 25, 2023
<u>AB</u>		<u>4MG</u>	<u>A213758 006</u>	Oct 25, 2023

REXULTI

<u>AB</u>	+ OTSUKA	<u>0.25MG</u>	<u>N205422 001</u>	Jul 10, 2015
<u>AB</u>	+	<u>0.5MG</u>	<u>N205422 002</u>	Jul 10, 2015
<u>AB</u>	+	<u>1MG</u>	<u>N205422 003</u>	Jul 10, 2015
<u>AB</u>	+!	<u>2MG</u>	<u>N205422 004</u>	Jul 10, 2015
<u>AB</u>	+	<u>3MG</u>	<u>N205422 005</u>	Jul 10, 2015
<u>AB</u>	+	<u>4MG</u>	<u>N205422 006</u>	Jul 10, 2015

BRIGATINIB

TABLET; ORAL

ALUNBRIG

+	TAKEDA PHARMS USA	30MG	N208772 001	Apr 28, 2017
+		90MG	N208772 002	Apr 28, 2017
+!		180MG	N208772 003	Oct 02, 2017

BRILLIANT BLUE G

SOLUTION; OPHTHALMIC

TISSUEBLUE

+! DUTCH OPHTHALMIC 0.025% N209569 001 Dec 20, 2019

BRIMONIDINE TARTRATE

GEL; TOPICAL

BRIMONIDINE TARTRATEAB PADAGIS ISRAEL EQ 0.33% BASE A209158 001 Sep 23, 2021MIRVASOAB +! GALDERMA LABS LP EQ 0.33% BASE N204708 001 Aug 23, 2013

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN PAB +! ABBVIE 0.1% N021770 001 Aug 19, 2005BRIMONIDINE TARTRATEAB ALEMBIC 0.1% A216909 001 Aug 01, 2023AB APOTEX 0.1% A078480 001 Dec 21, 2022ALPHAGAN PAT +! ABBVIE 0.15% N021262 001 Mar 16, 2001BRIMONIDINE TARTRATEAT ALEMBIC 0.15% A215225 001 Mar 29, 2023AT APOTEX 0.15% A078479 001 Jan 31, 2022AT ! BAUSCH AND LOMB 0.2% A076260 001 May 28, 2003AT INDOCO 0.2% A091691 001 Nov 18, 2014AT MICRO LABS 0.2% A217846 001 Nov 22, 2023AT RISING 0.2% A076439 001 Mar 14, 2006AT SANDOZ 0.2% A076254 001 Sep 16, 2003AT 0.2% A078075 001 Jan 30, 2008AT SOMERSET THERAPS 0.2% A208992 001 Mar 11, 2019QOLIANAAT +! SANDOZ 0.15% N021764 001 May 22, 2006

PRESCRIPTION DRUG PRODUCT LIST

BRIMONIDINE TARTRATE; BRINZOLAMIDE

SUSPENSION/DROPS;OPHTHALMIC

SIMBRINZA

+! ALCON LABS INC 0.2%;1%

N204251 001 Apr 19, 2013

BRIMONIDINE TARTRATE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

BRIMONIDINE TARTRATE AND TIMOLOL MALEATE

AB	ALEMBIC	<u>0.2%;EQ 0.5% BASE</u>	<u>A215230 001</u>	Aug 25, 2023
AB	APOTEX	<u>0.2%;EQ 0.5% BASE</u>	<u>A091442 001</u>	Apr 20, 2022
AB	FLORIDA	<u>0.2%;EQ 0.5% BASE</u>	<u>A201949 001</u>	Oct 04, 2022
AB	SANDOZ	<u>0.2%;EQ 0.5% BASE</u>	<u>A091087 001</u>	Apr 04, 2022
AB	SENTISS	<u>0.2%;EQ 0.5% BASE</u>	<u>A091086 001</u>	Oct 31, 2022
AB	UPSHER SMITH LABS	<u>0.2%;EQ 0.5% BASE</u>	<u>A215598 001</u>	Dec 12, 2022

COMBIGAN

AB	+! ABBEVIE	<u>0.2%;EQ 0.5% BASE</u>	<u>N021398 001</u>	Oct 30, 2007
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BRINCIDOFIVIR

SUSPENSION;ORAL

TEMBEXA

+! EMERGENT BIODEFENSE 10MG/ML

N214460 001 Jun 04, 2021

TABLET;ORAL

TEMBEXA

+! EMERGENT BIODEFENSE 100MG

N214461 001 Jun 04, 2021

BRINZOLAMIDE

SUSPENSION/DROPS;OPHTHALMIC

AZOPT

AB	+! SANDOZ	<u>1%</u>	<u>N020816 001</u>	Apr 01, 1998
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BRINZOLAMIDE

AB	BAUSCH AND LOMB	<u>1%</u>	<u>A204884 001</u>	Aug 18, 2021
AB	PADAGIS US	<u>1%</u>	<u>A211914 001</u>	Jul 28, 2023
AB	WATSON LABS INC	<u>1%</u>	<u>A209406 001</u>	Nov 27, 2020

BRIVARACETAM

SOLUTION;INTRAVENOUS

BRIVIACT

+! UCB INC 50MG/5ML (10MG/ML)

N205837 001 May 12, 2016

SOLUTION;ORAL

BRIVIACT

+! UCB INC 10MG/ML

N205838 001 May 12, 2016

TABLET;ORAL

BRIVIACT

+ UCB INC

10MG

N205836 001 May 12, 2016

+

25MG

N205836 002 May 12, 2016

+

50MG

N205836 003 May 12, 2016

+

75MG

N205836 004 May 12, 2016

+!

100MG

N205836 005 May 12, 2016

BROMFENAC SODIUM

SOLUTION/DROPS;OPHTHALMIC

BROMFENAC SODIUM

AB	LUPIN LTD	<u>EQ 0.07% ACID</u>	<u>A206027 001</u>	Nov 22, 2023
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PROLENSA

AB	+! BAUSCH AND LOMB	<u>EQ 0.07% ACID</u>	<u>N203168 001</u>	Apr 05, 2013
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BROMFENAC SODIUM

AT2	! ALEMBIC	<u>EQ 0.09% ACID</u>	<u>A210560 001</u>	Jun 21, 2019
AT2	EUGIA PHARMA	<u>EQ 0.09% ACID</u>	<u>A204813 001</u>	Mar 18, 2022
AT2	GLAND PHARMA LTD	<u>EQ 0.09% ACID</u>	<u>A211029 001</u>	Mar 17, 2020
AT2	LUPIN LTD	<u>EQ 0.09% ACID</u>	<u>A202903 001</u>	Aug 15, 2023
AT2	SENTISS	<u>EQ 0.09% ACID</u>	<u>A203395 001</u>	Jan 22, 2014

BROMSITE

+! SUN PHARM EQ 0.075% ACID

N206911 001 Apr 08, 2016

BROMOCRIPTINE MESYLATE

CAPSULE;ORAL

BROMOCRIPTINE MESYLATE

AB	! MYLAN	<u>EQ 5MG BASE</u>	<u>A077226 001</u>	Apr 04, 2005
AB	ZYDUS PHARMS USA INC	<u>EQ 5MG BASE</u>	<u>A078899 001</u>	Jul 30, 2008

PARLODEL

AB	+ SS PHARMA	<u>EQ 5MG BASE</u>	<u>N017962 002</u>	Mar 01, 1982
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TABLET;ORAL

BROMOCRIPTINE MESYLATE

AB	! PADAGIS US	<u>EQ 2.5MG BASE</u>	<u>A077646 001</u>	Oct 01, 2008
AB	SANDOZ	<u>EQ 2.5MG BASE</u>	<u>A074631 001</u>	Jan 13, 1998

PRESCRIPTION DRUG PRODUCT LIST

BROMOCRIPTINE MESYLATE

TABLET; ORAL

PARLODEL

<u>AB</u>	+	SS PHARMA	<u>EQ 2.5MG BASE</u>	<u>N017962 001</u>	
		CYCLOSET			
	+	VEROSCIENCE	EQ 0.8MG BASE	N020866 001	May 05, 2009

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMFED-DM

<u>AA</u>	+	WOCKHARDT BIO AG	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A088811 001</u>	Jun 07, 1985
		<u>BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE</u>			
<u>AA</u>		ACELLA	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A203375 001</u>	Sep 20, 2016
<u>AA</u>		ALKEM LABS LTD	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A210647 001</u>	Jul 14, 2020
<u>AA</u>		CHARTWELL MOLECULAR	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A213125 001</u>	Apr 17, 2020
<u>AA</u>		DR REDDYS LABS SA	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A207676 001</u>	Dec 04, 2018
<u>AA</u>	!	PADAGIS US	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A205292 001</u>	Jul 15, 2014
<u>AA</u>		PHARM ASSOC	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A202940 001</u>	Jul 21, 2014
<u>AA</u>		TARO	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A205112 001</u>	Feb 27, 2017
<u>AA</u>		WES PHARMA INC	<u>2MG/5ML; 10MG/5ML; 30MG/5ML</u>	<u>A211170 001</u>	Jun 16, 2020

BUDESONIDE

AEROSOL, FOAM; RECTAL

BUDESONIDE

<u>AB</u>		PADAGIS ISRAEL	<u>2MG/ACTUATION</u>	<u>A215328 001</u>	Apr 12, 2023
		<u>UCERIS</u>			
<u>AB</u>	+	SALIX	<u>2MG/ACTUATION</u>	<u>N205613 001</u>	Oct 07, 2014

CAPSULE, DELAYED RELEASE; ORAL

BUDESONIDE

<u>AB</u>		AMNEAL PHARMS	<u>3MG</u>	<u>A206200 001</u>	Jul 31, 2017
<u>AB</u>		AUROBINDO PHARMA USA	<u>3MG</u>	<u>A090410 001</u>	May 16, 2011
<u>AB</u>		DR REDDYS LABS SA	<u>3MG</u>	<u>A206623 001</u>	Apr 08, 2016
<u>AB</u>		RISING	<u>3MG</u>	<u>A207367 001</u>	Apr 07, 2017
<u>AB</u>		ZYDUS PHARMS	<u>3MG</u>	<u>A206134 001</u>	May 04, 2017

ENTOCORT EC

<u>AB</u>	+	PADAGIS US	<u>3MG</u>	<u>N021324 001</u>	Oct 02, 2001
		TARPEYO			
	+	CALLIDITAS	4MG	N215935 001	Dec 15, 2021
		POWDER, METERED; INHALATION			
		PULMICORT FLEXHALER			
	+	ASTRAZENECA	0.08MG/INH	N021949 001	Jul 12, 2006
	+		0.16MG/INH	N021949 002	Jul 12, 2006

SUSPENSION; INHALATION

BUDESONIDE

<u>AN</u>		CIPLA	<u>0.25MG/2ML</u>	<u>A205710 001</u>	Nov 16, 2017
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A205710 002</u>	Nov 16, 2017
<u>AN</u>			<u>1MG/2ML</u>	<u>A205710 003</u>	Nov 16, 2017
<u>AN</u>		EUGIA PHARMA	<u>0.5MG/2ML</u>	<u>A216667 001</u>	Nov 29, 2023
<u>AN</u>		IMPAX LABS INC	<u>0.25MG/2ML</u>	<u>A078404 001</u>	Jul 31, 2012
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A078404 002</u>	Jul 31, 2012
<u>AN</u>		LUPIN	<u>0.5MG/2ML</u>	<u>A210897 001</u>	Nov 09, 2018
<u>AN</u>		NEPHRON	<u>0.25MG/2ML</u>	<u>A078202 001</u>	Mar 30, 2009
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A078202 002</u>	Mar 30, 2009
<u>AN</u>		SANDOZ	<u>0.25MG/2ML</u>	<u>A201966 003</u>	Sep 27, 2013
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A201966 002</u>	Sep 27, 2013
<u>AN</u>			<u>1MG/2ML</u>	<u>A201966 001</u>	Sep 27, 2013
<u>AN</u>		SUN PHARM	<u>0.25MG/2ML</u>	<u>A211922 001</u>	Apr 14, 2021
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A211922 002</u>	Apr 14, 2021
<u>AN</u>			<u>1MG/2ML</u>	<u>A211922 003</u>	Apr 14, 2021
<u>AN</u>		TEVA PHARMS	<u>0.25MG/2ML</u>	<u>A077519 001</u>	Nov 18, 2008
<u>AN</u>			<u>0.5MG/2ML</u>	<u>A077519 002</u>	Nov 18, 2008
<u>AN</u>		TEVA PHARMS USA	<u>1MG/2ML</u>	<u>A204548 001</u>	Mar 08, 2016
		<u>PULMICORT RESPULES</u>			
<u>AN</u>	+	ASTRAZENECA	<u>0.25MG/2ML</u>	<u>N020929 001</u>	Aug 08, 2000
<u>AN</u>	+		<u>0.5MG/2ML</u>	<u>N020929 002</u>	Aug 08, 2000
<u>AN</u>	+		<u>1MG/2ML</u>	<u>N020929 003</u>	Aug 08, 2000

TABLET, EXTENDED RELEASE; ORAL

BUDESONIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>9MG</u>	<u>A205457 001</u>	Jul 03, 2018
<u>AB</u>		MYLAN	<u>9MG</u>	<u>A208851 001</u>	Sep 17, 2020
		<u>UCERIS</u>			
<u>AB</u>	+	SALIX	<u>9MG</u>	<u>N203634 001</u>	Jan 14, 2013

PRESCRIPTION DRUG PRODUCT LISTBUDESONIDE; FORMOTEROL FUMARATEAEROSOL, METERED; INHALATION
SYMBICORT AEROSPHERE

+! ASTRAZENECA 0.16MG/INH; 0.0048MG/INH N216579 001 Apr 28, 2023

BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

AEROSOL, METERED; INHALATION

BREYNA**AB** MYLAN **0.08MG/INH; 0.0045MG/INH** **A211699 001** Mar 15, 2022**AB** **0.16MG/INH; 0.0045MG/INH** **A211699 002** Mar 15, 2022**SYMBICORT****AB** +! ASTRAZENECA **0.08MG/INH; 0.0045MG/INH** **N021929 001** Jul 21, 2006**AB** +! **0.16MG/INH; 0.0045MG/INH** **N021929 002** Jul 21, 2006BUDESONIDE; FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION

BREZTRI AEROSPHERE

+! ASTRAZENECA AB 0.16MG/INH; 0.0048MG/INH; 0.009MG/INH N212122 001 Jul 23, 2020

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE**AP** GLAND PHARMA LTD **0.25MG/ML** **A216434 001** May 26, 2022**AP** HOSPIRA **0.25MG/ML** **A074332 001** Oct 31, 1994**AP** MSN **0.25MG/ML** **A215364 001** Aug 04, 2022**AP** SAGENT **0.25MG/ML** **A074441 001** Jan 27, 1995**AP** ! WEST-WARD PHARMS **0.25MG/ML** **A079196 001** Apr 30, 2008

INT

TABLET; ORAL

BUMETANIDE**AB** AMNEAL PHARMS CO **0.5MG** **A209724 001** Oct 18, 2017**AB** **1MG** **A209724 002** Oct 18, 2017**AB** **2MG** **A209724 003** Oct 18, 2017**AB** HERITAGE PHARMA **0.5MG** **A074225 001** Apr 24, 1995**AB** **1MG** **A074225 002** Apr 24, 1995**AB** **2MG** **A074225 003** Apr 24, 1995**AB** SANDOZ **0.5MG** **A074700 001** Nov 21, 1996**AB** **1MG** **A074700 002** Nov 21, 1996**AB** ! **2MG** **A074700 003** Nov 21, 1996**AB** TARO **0.5MG** **A213458 001** Jul 24, 2023**AB** **1MG** **A213458 002** Jul 24, 2023**AB** **2MG** **A213458 003** Jul 24, 2023**AB** UPSHER SMITH LABS **0.5MG** **A209916 001** Jan 23, 2018**AB** **1MG** **A209916 002** Jan 23, 2018**AB** **2MG** **A209916 003** Jan 23, 2018**AB** ZYDUS PHARMS **0.5MG** **A202900 001** Apr 30, 2018**AB** **1MG** **A202900 002** Apr 30, 2018**AB** **2MG** **A202900 003** Apr 30, 2018**BUMEX****AB** + VALIDUS PHARMS **0.5MG** **N018225 002** Feb 28, 1983**AB** + **1MG** **N018225 001** Feb 28, 1983**AB** + **2MG** **N018225 003** Jun 14, 1985

BUMETANIDE

BX RISING 0.5MG A212019 001 Dec 12, 2019

BX 1MG A212019 002 Dec 12, 2019

BX 2MG A212019 003 Dec 12, 2019

BUPIVACAINE

INJECTABLE, LIPOSOMAL; INJECTION

EXPAREL

+! PACIRA PHARMS INC 133MG/10ML (13.3MG/ML) N022496 001 Oct 28, 2011

+! 266MG/20ML (13.3MG/ML) N022496 002 Oct 28, 2011

SOLUTION, EXTENDED RELEASE; INFILTRATION

POSIMIR

+! INNOCOLL 660MG/5ML (132MG/ML) N204803 001 Feb 01, 2021

BUPIVACAINE HYDROCHLORIDE

IMPLANT; IMPLANTATION

XARACOLL

+! INNOCOLL PHARMS 100MG N209511 001 Aug 28, 2020

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE**AP** EUGIA PHARMA **0.25%** **A207183 001** May 13, 2016**AP** **0.5%** **A207183 002** May 13, 2016**AP** HIKMA PHARMS **0.25%** **A205141 001** Feb 11, 2021

PRESCRIPTION DRUG PRODUCT LIST

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>		<u>0.5%</u>	<u>A205141 002</u>	Feb 11, 2021
<u>AP</u>	HOSPIRA	<u>0.25%</u>	<u>A070583 001</u>	Feb 17, 1987
<u>AP</u>		<u>0.25%</u>	<u>A070590 001</u>	Feb 17, 1987
<u>AP</u>		<u>0.5%</u>	<u>A070584 001</u>	Feb 17, 1986
<u>AP</u>		<u>0.5%</u>	<u>A070597 001</u>	Mar 03, 1987
<u>AP</u>		<u>0.5%</u>	<u>A070609 001</u>	Mar 03, 1987
<u>AP</u>		<u>0.75%</u>	<u>A070585 001</u>	Mar 03, 1987
<u>AP</u>	KINDOS	<u>0.25%</u>	<u>A216039 001</u>	Jun 23, 2023
<u>AP</u>		<u>0.25%</u>	<u>A216040 001</u>	Dec 27, 2023
<u>AP</u>		<u>0.5%</u>	<u>A216039 002</u>	Jun 23, 2023
<u>AP</u>		<u>0.5%</u>	<u>A216040 002</u>	Dec 27, 2023
<u>AP</u>		<u>0.75%</u>	<u>A216040 003</u>	Dec 27, 2023
<u>AP</u>	SOMERSET	<u>0.25%</u>	<u>A217792 001</u>	Nov 20, 2023
<u>AP</u>		<u>0.5%</u>	<u>A217792 002</u>	Nov 20, 2023
<u>AP</u>	STERISCIENCE	<u>0.25%</u>	<u>A091503 001</u>	Oct 18, 2011
<u>AP</u>		<u>0.5%</u>	<u>A091503 002</u>	Oct 18, 2011

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	EUGIA PHARMA	<u>0.25%</u>	<u>A203895 001</u>	Nov 05, 2013
<u>AP</u>		<u>0.5%</u>	<u>A203895 002</u>	Nov 05, 2013
<u>AP</u>		<u>0.75%</u>	<u>A203895 003</u>	Nov 05, 2013
<u>AP</u>	HIKMA PHARMS	<u>0.25%</u>	<u>A204842 001</u>	Feb 11, 2021
<u>AP</u>		<u>0.5%</u>	<u>A204842 002</u>	Feb 11, 2021
<u>AP</u>		<u>0.75%</u>	<u>A204842 003</u>	Feb 11, 2021
<u>AP</u>	STERISCIENCE	<u>0.25%</u>	<u>A091487 002</u>	Oct 18, 2011
<u>AP</u>		<u>0.5%</u>	<u>A091487 001</u>	Oct 18, 2011
<u>AP</u>		<u>0.75%</u>	<u>A091487 003</u>	Oct 18, 2011

MARCAINE HYDROCHLORIDE

<u>AP</u>	+! HOSPIRA	<u>0.25%</u>	<u>N016964 001</u>	
<u>AP</u>	+!	<u>0.5%</u>	<u>N016964 006</u>	

MARCAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	+! HOSPIRA	<u>0.25%</u>	<u>N016964 012</u>	
<u>AP</u>	+!	<u>0.5%</u>	<u>N016964 005</u>	
<u>AP</u>	+!	<u>0.75%</u>	<u>N016964 009</u>	

SENSORCAINE

<u>AP</u>	FRESENIUS KABI USA	<u>0.25%</u>	<u>A070552 001</u>	May 21, 1986
<u>AP</u>	+	<u>0.25%</u>	<u>N018304 001</u>	
<u>AP</u>		<u>0.5%</u>	<u>A070553 001</u>	May 21, 1986
<u>AP</u>	+	<u>0.5%</u>	<u>N018304 002</u>	
<u>AP</u>		<u>0.75%</u>	<u>A070554 001</u>	May 21, 1986
<u>AP</u>	+	<u>0.75%</u>	<u>N018304 003</u>	

INJECTABLE; SPINAL

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>	B BRAUN MEDICAL INC	<u>0.75%</u>	<u>A209087 001</u>	Apr 02, 2019
<u>AP</u>	HOSPIRA	<u>0.75%</u>	<u>A071810 001</u>	Dec 11, 1987
<u>AP</u>	HUONS	<u>0.75%</u>	<u>A212822 001</u>	Dec 30, 2019

MARCAINE

<u>AP</u>	+! HOSPIRA	<u>0.75%</u>	<u>N018692 001</u>	May 04, 1984
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BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

<u>AP</u>	! HOSPIRA	<u>0.5%;0.005MG/ML</u>	<u>A071168 001</u>	Jun 16, 1988
<u>AP</u>		<u>0.5%;0.005MG/ML</u>	<u>A071170 001</u>	Jun 16, 1988
	!	0.25%;0.005MG/ML	A071165 001	Jun 16, 1988
		0.25%;0.005MG/ML	A071167 001	Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE

<u>AP</u>	+! HOSPIRA	<u>0.25%;0.0091MG/ML</u>	<u>N016964 004</u>	
<u>AP</u>	+!	<u>0.5%;0.0091MG/ML</u>	<u>N016964 008</u>	

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE

<u>AP</u>	+! HOSPIRA	<u>0.25%;0.0091MG/ML</u>	<u>N016964 013</u>	
<u>AP</u>	+!	<u>0.5%;0.0091MG/ML</u>	<u>N016964 007</u>	
<u>AP</u>	+!	<u>0.75%;0.0091MG/ML</u>	<u>N016964 010</u>	

SENSORCAINE

<u>AP</u>	FRESENIUS KABI USA	<u>0.25%;0.0091MG/ML</u>	<u>A070966 001</u>	Oct 13, 1987
<u>AP</u>		<u>0.25%;0.0091MG/ML</u>	<u>A070967 001</u>	Oct 13, 1987
<u>AP</u>		<u>0.5%;0.0091MG/ML</u>	<u>A070968 001</u>	Oct 13, 1987
<u>AP</u>	+	<u>0.5%;0.0091MG/ML</u>	<u>N018304 004</u>	Sep 02, 1983

PRESCRIPTION DRUG PRODUCT LISTBUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

SENSORCAINE

AP	+		0.75%;0.0091MG/ML	N018304 005	Sep 02, 1983
		BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE			
	!	SEPTODONT	0.5%;0.0091MG/ML	A077250 001	Sep 27, 2006

BUPIVACAINE; MELOXICAM

SOLUTION, EXTENDED RELEASE; PERIARTICULAR

ZYNRELEF KIT

+	!	HERON THERAPS INC	200MG/7ML (29.25MG/ML); 6MG/7ML (0.88MG/ML)	N211988 002	May 12, 2021
+	!		400MG/14ML (29.25MG/ML); 12MG/14ML (0.88MG/ML)	N211988 004	May 12, 2021

BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

AB		ALVOGEN	5MCG/HR	A207490 001	May 17, 2022
AB			10MCG/HR	A207490 002	May 17, 2022
AB			15MCG/HR	A207490 003	May 17, 2022
AB			20MCG/HR	A207490 004	May 17, 2022
AB		AMNEAL	5MCG/HR	A211586 001	Apr 14, 2020
AB			7.5MCG/HR	A211586 002	Apr 14, 2020
AB			10MCG/HR	A211586 003	Apr 14, 2020
AB			15MCG/HR	A211586 004	Apr 14, 2020
AB			20MCG/HR	A211586 005	Apr 14, 2020
AB		AVEVA	5MCG/HR	A210272 001	Sep 23, 2021
AB			7.5MCG/HR	A210272 002	Sep 23, 2021
AB			10MCG/HR	A210272 003	Sep 23, 2021
AB			15MCG/HR	A210272 004	Sep 23, 2021
AB			20MCG/HR	A210272 005	Sep 23, 2021
AB		WATSON LABS TEVA	5MCG/HR	A204937 001	Nov 20, 2018
AB			7.5MCG/HR	A204937 005	Jun 29, 2021
AB			10MCG/HR	A204937 002	Nov 20, 2018
AB			15MCG/HR	A204937 003	Nov 20, 2018
AB			20MCG/HR	A204937 004	Nov 20, 2018

BUTRANS

AB	+	PURDUE PHARMA LP	5MCG/HR	N021306 001	Jun 30, 2010
AB	+		7.5MCG/HR	N021306 005	Jun 30, 2014
AB	+		10MCG/HR	N021306 002	Jun 30, 2010
AB	+		15MCG/HR	N021306 004	Jul 25, 2013
AB	+		20MCG/HR	N021306 003	Jun 30, 2010

SOLUTION, EXTENDED RELEASE; SUBCUTANEOUS

BRIXADI

+		BRAEBURN	8MG/0.16ML (50MG/ML)	N210136 001	May 23, 2023
+			16MG/0.32ML (50MG/ML)	N210136 002	May 23, 2023
+			24MG/0.48ML (50MG/ML)	N210136 003	May 23, 2023
+			32MG/0.64ML (50MG/ML)	N210136 004	May 23, 2023
+			64MG/0.18ML (356MG/ML)	N210136 005	May 23, 2023
+			96MG/0.27ML (356MG/ML)	N210136 006	May 23, 2023
+	!		128MG/0.36ML (356MG/ML)	N210136 007	May 23, 2023

SUBLOCADE

+		INDIVIOR	100MG/0.5ML (100MG/0.5ML)	N209819 001	Nov 30, 2017
+	!		300MG/1.5ML (200MG/ML)	N209819 002	Nov 30, 2017

BUPRENORPHINE HYDROCHLORIDE

FILM; BUCCAL

BELBUCA

+		BDSI	EQ 0.075MG BASE	N207932 001	Oct 23, 2015
+			EQ 0.15MG BASE	N207932 002	Oct 23, 2015
+			EQ 0.3MG BASE	N207932 003	Oct 23, 2015
+			EQ 0.45MG BASE	N207932 004	Oct 23, 2015
+			EQ 0.6MG BASE	N207932 005	Oct 23, 2015
+			EQ 0.75MG BASE	N207932 006	Oct 23, 2015
+	!		EQ 0.9MG BASE	N207932 007	Oct 23, 2015

INJECTABLE; INJECTION

BUPRENORPHINE HYDROCHLORIDE

AP		AM REGENT	EQ 0.3MG BASE/ML	A078331 001	Mar 27, 2007
AP		HIKMA	EQ 0.3MG BASE/ML	A076931 001	Mar 02, 2005
AP		HOSPIRA	EQ 0.3MG BASE/ML	A074137 001	Jun 03, 1996
AP	!	PAR STERILE PRODUCTS	EQ 0.3MG BASE/ML	A206586 001	Jul 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

BUPRENORPHINE HYDROCHLORIDE

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 2MG BASE</u>	<u>A090819 001</u>	Feb 19, 2015
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A090819 002</u>	Feb 19, 2015
<u>AB</u>	ETHYPHARM	<u>EQ 2MG BASE</u>	<u>A090622 001</u>	Sep 24, 2010
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A090622 002</u>	Sep 24, 2010
<u>AB</u>	HIKMA	<u>EQ 2MG BASE</u>	<u>A078633 001</u>	Oct 08, 2009
<u>AB</u>	!	<u>EQ 8MG BASE</u>	<u>A078633 002</u>	Oct 08, 2009
<u>AB</u>	RHODES PHARMS	<u>EQ 2MG BASE</u>	<u>A207276 001</u>	Mar 27, 2017
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A207276 002</u>	Mar 27, 2017
<u>AB</u>	RUBICON	<u>EQ 2MG BASE</u>	<u>A090279 001</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A090279 002</u>	Jun 10, 2015
<u>AB</u>	SUN PHARM	<u>EQ 2MG BASE</u>	<u>A201760 001</u>	Jan 29, 2016
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A201760 002</u>	Jan 29, 2016

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM;BUCCAL, SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

<u>AB</u>	ALVOGEN	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205954 001</u>	Jan 24, 2019
<u>AB</u>		<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A205954 002</u>	Jan 24, 2019
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205954 003</u>	Jan 24, 2019
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A205954 004</u>	Jan 24, 2019
<u>AB</u>	AVEVA	<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A212756 001</u>	Jun 02, 2022
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A212756 002</u>	Jun 02, 2022
<u>AB</u>	DR REDDYS LABS SA	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205299 001</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A205806 001</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205299 002</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A205806 002</u>	Jun 14, 2018
<u>AB</u>	MYLAN TECHNOLOGIES	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A211785 001</u>	Apr 17, 2020
<u>AB</u>		<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>A211785 002</u>	Apr 17, 2020
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A207607 001</u>	Jun 14, 2018
<u>AB</u>		<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>A207607 002</u>	Jun 14, 2018

SUBOXONE

<u>AB</u>	+ INDIVIOR	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>N022410 001</u>	Aug 30, 2010
<u>AB</u>	+	<u>EQ 4MG BASE;EQ 1MG BASE</u>	<u>N022410 003</u>	Aug 10, 2012
<u>AB</u>	+	<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>N022410 002</u>	Aug 30, 2010
<u>AB</u>	+!	<u>EQ 12MG BASE;EQ 3MG BASE</u>	<u>N022410 004</u>	Aug 10, 2012

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A091422 001</u>	Feb 22, 2013
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A091422 002</u>	Feb 22, 2013
<u>AB</u>	ALKEM LABS LTD	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A214930 001</u>	Jun 15, 2021
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A214930 002</u>	Jun 15, 2021
<u>AB</u>	AMNEAL PHARMS	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A203136 001</u>	Feb 22, 2013
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A203136 002</u>	Feb 22, 2013
<u>AB</u>	ETHYPHARM USA CORP	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A204431 001</u>	Oct 16, 2015
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A204431 002</u>	Oct 16, 2015
<u>AB</u>	HIKMA	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A203326 001</u>	Jun 27, 2014
<u>AB</u>	!	<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A203326 002</u>	Jun 27, 2014
<u>AB</u>	LANNETT CO INC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205022 001</u>	Sep 19, 2016
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205022 002</u>	Sep 19, 2016
<u>AB</u>	RHODES PHARMS	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A205601 001</u>	Mar 30, 2020
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A205601 002</u>	Mar 30, 2020
<u>AB</u>	SPECGX LLC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A207000 001</u>	Dec 13, 2017
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A207000 002</u>	Dec 13, 2017
<u>AB</u>	SUN PHARM	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A201633 001</u>	Aug 05, 2016
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A201633 002</u>	Aug 05, 2016
<u>AB</u>	WES PHARMA INC	<u>EQ 2MG BASE;EQ 0.5MG BASE</u>	<u>A209069 001</u>	Jul 17, 2020
<u>AB</u>		<u>EQ 8MG BASE;EQ 2MG BASE</u>	<u>A209069 002</u>	Jul 17, 2020

ZUBSOLV

+	OREXO US INC	EQ 0.7MG BASE;EQ 0.18MG BASE	N204242 006	Oct 04, 2016
+		EQ 1.4MG BASE;EQ 0.36MG BASE	N204242 001	Jul 03, 2013
+		EQ 2.9MG BASE;EQ 0.71MG BASE	N204242 005	Jun 04, 2015
+		EQ 5.7MG BASE;EQ 1.4MG BASE	N204242 002	Jul 03, 2013
+		EQ 8.6MG BASE;EQ 2.1MG BASE	N204242 003	Dec 11, 2014
+	!	EQ 11.4MG BASE;EQ 2.9MG BASE	N204242 004	Dec 11, 2014

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

APLENZIN

+	BAUSCH	174MG	N022108	001	Apr 23, 2008
+		348MG	N022108	002	Apr 23, 2008
+	!	522MG	N022108	003	Apr 23, 2008

BUPROPION HYDROCHLORIDE

TABLET;ORAL

BUPROPION HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>75MG</u>	<u>A203013</u>	<u>001</u>	Jun 08, 2018
<u>AB</u>		<u>100MG</u>	<u>A203013</u>	<u>002</u>	Jun 08, 2018
<u>AB</u>	APNAR PHARMA LP	<u>75MG</u>	<u>A075584</u>	<u>001</u>	Feb 07, 2000
<u>AB</u>		<u>100MG</u>	<u>A075584</u>	<u>002</u>	Feb 07, 2000
<u>AB</u>	APOTEX INC	<u>75MG</u>	<u>A076143</u>	<u>001</u>	Jan 17, 2006
<u>AB</u>	!	<u>100MG</u>	<u>A076143</u>	<u>002</u>	Jan 17, 2006
<u>AB</u>	CADILA PHARMS LTD	<u>75MG</u>	<u>A208606</u>	<u>001</u>	Jan 16, 2020
<u>AB</u>		<u>100MG</u>	<u>A208606</u>	<u>002</u>	Jan 16, 2020
<u>AB</u>	HERITAGE PHARMA	<u>75MG</u>	<u>A206975</u>	<u>001</u>	Aug 19, 2016
<u>AB</u>		<u>100MG</u>	<u>A206975</u>	<u>002</u>	Aug 19, 2016
<u>AB</u>	INVAGEN PHARMS	<u>75MG</u>	<u>A207389</u>	<u>001</u>	Sep 18, 2017
<u>AB</u>		<u>100MG</u>	<u>A207389</u>	<u>002</u>	Sep 18, 2017
<u>AB</u>	MICRO LABS	<u>75MG</u>	<u>A207403</u>	<u>001</u>	Apr 17, 2020
<u>AB</u>		<u>100MG</u>	<u>A207403</u>	<u>002</u>	Apr 17, 2020

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

<u>AB1</u>	ACTAVIS LABS FL INC	<u>100MG</u>	<u>A079095</u>	<u>001</u>	Mar 24, 2009
<u>AB1</u>		<u>150MG</u>	<u>A079095</u>	<u>002</u>	Mar 24, 2009
<u>AB1</u>		<u>200MG</u>	<u>A079095</u>	<u>003</u>	Mar 24, 2009
<u>AB1</u>	ANCHEN PHARMS	<u>100MG</u>	<u>A091459</u>	<u>001</u>	Jun 09, 2011
<u>AB1</u>		<u>150MG</u>	<u>A091459</u>	<u>002</u>	Jun 09, 2011
<u>AB1</u>		<u>200MG</u>	<u>A091459</u>	<u>003</u>	Jun 09, 2011
<u>AB1</u>	ANNORA PHARMA	<u>100MG</u>	<u>A216800</u>	<u>001</u>	May 31, 2023
<u>AB1</u>		<u>150MG</u>	<u>A216800</u>	<u>002</u>	May 31, 2023
<u>AB1</u>		<u>200MG</u>	<u>A216800</u>	<u>003</u>	May 31, 2023
<u>AB1</u>	EPIC PHARMA LLC	<u>100MG</u>	<u>A075932</u>	<u>001</u>	Nov 25, 2003
<u>AB1</u>		<u>150MG</u>	<u>A075932</u>	<u>002</u>	Mar 22, 2004
<u>AB1</u>		<u>200MG</u>	<u>A075932</u>	<u>003</u>	Jun 22, 2005
<u>AB1</u>	IMPAX LABS	<u>100MG</u>	<u>A075913</u>	<u>001</u>	Jan 28, 2004
<u>AB1</u>		<u>150MG</u>	<u>A075913</u>	<u>002</u>	Mar 22, 2004
<u>AB1</u>	INVAGEN PHARMS	<u>100MG</u>	<u>A206674</u>	<u>001</u>	Feb 09, 2016
<u>AB1</u>		<u>150MG</u>	<u>A206674</u>	<u>002</u>	Feb 09, 2016
<u>AB1</u>		<u>200MG</u>	<u>A206674</u>	<u>003</u>	Feb 09, 2016
<u>AB1</u>	PRINSTON INC	<u>100MG</u>	<u>A202304</u>	<u>001</u>	May 26, 2015
<u>AB1</u>		<u>150MG</u>	<u>A202304</u>	<u>002</u>	May 26, 2015
<u>AB1</u>		<u>200MG</u>	<u>A202304</u>	<u>003</u>	May 26, 2015
<u>AB1</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A205794</u>	<u>001</u>	Mar 01, 2016
<u>AB1</u>		<u>150MG</u>	<u>A205794</u>	<u>002</u>	Mar 01, 2016
<u>AB1</u>		<u>200MG</u>	<u>A205794</u>	<u>003</u>	Mar 01, 2016
<u>AB1</u>	SUN PHARM	<u>150MG</u>	<u>A078866</u>	<u>002</u>	Apr 06, 2010
<u>AB1</u>		<u>200MG</u>	<u>A078866</u>	<u>003</u>	Apr 06, 2010
<u>AB1</u>	YICHANG HUMANWELL	<u>100MG</u>	<u>A211347</u>	<u>001</u>	Oct 16, 2018
<u>AB1</u>		<u>150MG</u>	<u>A211347</u>	<u>002</u>	Oct 16, 2018
<u>AB1</u>		<u>200MG</u>	<u>A211347</u>	<u>003</u>	Oct 16, 2018

WELLBUTRIN SR

<u>AB1</u>	+	GLAXOSMITHKLINE	<u>100MG</u>	<u>N020358</u>	<u>002</u>	Oct 04, 1996
<u>AB1</u>	+		<u>150MG</u>	<u>N020358</u>	<u>003</u>	Oct 04, 1996
<u>AB1</u>	+	!	<u>200MG</u>	<u>N020358</u>	<u>004</u>	Jun 14, 2002

BUPROPION HYDROCHLORIDE

<u>AB2</u>	!	ACTAVIS LABS FL INC	<u>150MG</u>	<u>A079094</u>	<u>001</u>	Mar 24, 2009
<u>AB2</u>		ANCHEN PHARMS	<u>150MG</u>	<u>A091520</u>	<u>001</u>	Jun 09, 2011
<u>AB2</u>		IMPAX LABS	<u>150MG</u>	<u>A075914</u>	<u>001</u>	May 27, 2004
<u>AB2</u>		SANDOZ	<u>150MG</u>	<u>A077475</u>	<u>001</u>	Mar 12, 2008
<u>AB2</u>		SCIEGEN PHARMS INC	<u>150MG</u>	<u>A206122</u>	<u>001</u>	Aug 17, 2016
<u>AB2</u>		YICHANG HUMANWELL	<u>150MG</u>	<u>A216766</u>	<u>001</u>	Jan 09, 2023
<u>AB3</u>		ACCORD HLTHCARE	<u>150MG</u>	<u>A210497</u>	<u>001</u>	Oct 31, 2018
<u>AB3</u>			<u>300MG</u>	<u>A210497</u>	<u>002</u>	Oct 31, 2018
<u>AB3</u>		ACTAVIS LABS FL INC	<u>150MG</u>	<u>A077715</u>	<u>001</u>	Nov 26, 2008
<u>AB3</u>		ANBISON LAB	<u>150MG</u>	<u>A207224</u>	<u>001</u>	Jun 30, 2017
<u>AB3</u>			<u>300MG</u>	<u>A207224</u>	<u>002</u>	Jun 30, 2017
<u>AB3</u>		GRANULES	<u>150MG</u>	<u>A215568</u>	<u>001</u>	Feb 02, 2022
<u>AB3</u>			<u>300MG</u>	<u>A215568</u>	<u>002</u>	Feb 02, 2022
<u>AB3</u>		GRAVITI PHARMS	<u>150MG</u>	<u>A211020</u>	<u>001</u>	Jan 28, 2019

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

<u>AB3</u>		<u>300MG</u>	<u>A211020 002</u>	Jan 28, 2019
<u>AB3</u>	INVAGEN PHARMS	<u>150MG</u>	<u>A206556 001</u>	Aug 26, 2016
<u>AB3</u>		<u>300MG</u>	<u>A206556 002</u>	Aug 26, 2016
<u>AB3</u>	LUPIN LTD	<u>150MG</u>	<u>A090693 001</u>	Apr 06, 2017
<u>AB3</u>		<u>300MG</u>	<u>A090693 002</u>	Apr 06, 2017
<u>AB3</u>	SCIEGEN PHARMS INC	<u>150MG</u>	<u>A207479 001</u>	Apr 12, 2017
<u>AB3</u>		<u>300MG</u>	<u>A207479 002</u>	Apr 12, 2017
<u>AB3</u>	SCINOPHARM TAIWAN	<u>150MG</u>	<u>A210081 001</u>	Nov 03, 2017
<u>AB3</u>		<u>300MG</u>	<u>A210081 002</u>	Nov 03, 2017
<u>AB3</u>	SINOTHERAPEUTICS INC	<u>150MG</u>	<u>A208652 001</u>	Aug 21, 2017
<u>AB3</u>		<u>300MG</u>	<u>A208652 002</u>	Aug 21, 2017
<u>AB3</u>	SUN PHARM	<u>150MG</u>	<u>A200216 001</u>	Nov 30, 2020
<u>AB3</u>		<u>300MG</u>	<u>A203650 001</u>	Dec 31, 2020
<u>AB3</u>	WATSON LABS INC	<u>150MG</u>	<u>A077285 001</u>	Nov 26, 2008
<u>AB3</u>		<u>300MG</u>	<u>A077285 002</u>	Aug 15, 2008
<u>AB3</u>	WOCKHARDT LTD	<u>150MG</u>	<u>A202189 001</u>	Nov 21, 2012
<u>AB3</u>		<u>300MG</u>	<u>A202189 002</u>	Jan 28, 2022
<u>AB3</u>	YICHANG HUMANWELL	<u>150MG</u>	<u>A210015 001</u>	Jun 14, 2018
<u>AB3</u>		<u>300MG</u>	<u>A210015 002</u>	Jun 14, 2018
<u>AB3</u>	ZHEJIANG JUTAI PHARM	<u>150MG</u>	<u>A211200 002</u>	Apr 29, 2020
<u>AB3</u>		<u>300MG</u>	<u>A211200 001</u>	Sep 05, 2019
<u>AB3</u>	ZYDUS PHARMS	<u>150MG</u>	<u>A201567 002</u>	Jul 23, 2018
<u>AB3</u>		<u>300MG</u>	<u>A201567 001</u>	Jan 17, 2014

WELLBUTRIN XL

<u>AB3</u>	+ BAUSCH	<u>150MG</u>	<u>N021515 001</u>	Aug 28, 2003
<u>AB3</u>	+!	<u>300MG</u>	<u>N021515 002</u>	Aug 28, 2003
	FORFIVO XL			
	+! TWI PHARMS	450MG	N022497 001	Nov 10, 2011

BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

AUVELITY

+!	AXSOME	105MG;45MG	N215430 001	Aug 18, 2022
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BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CONTRAVE

+!	NALPROPION	90MG;8MG	N200063 001	Sep 10, 2014
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BUSPIRONE HYDROCHLORIDE

TABLET;ORAL

BUSPIRONE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A202557 001</u>	Dec 30, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A202557 002</u>	Dec 30, 2014
<u>AB</u>		<u>10MG</u>	<u>A202557 003</u>	Dec 30, 2014
<u>AB</u>	!	<u>15MG</u>	<u>A202557 004</u>	Dec 30, 2014
<u>AB</u>		<u>30MG</u>	<u>A202557 005</u>	Dec 30, 2014
<u>AB</u>	AMNEAL PHARMS CO	<u>5MG</u>	<u>A208829 001</u>	May 24, 2017
<u>AB</u>		<u>7.5MG</u>	<u>A208829 002</u>	May 24, 2017
<u>AB</u>		<u>10MG</u>	<u>A208829 003</u>	May 24, 2017
<u>AB</u>		<u>15MG</u>	<u>A208829 004</u>	May 24, 2017
<u>AB</u>		<u>30MG</u>	<u>A208829 005</u>	May 24, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A078246 001</u>	Feb 27, 2009
<u>AB</u>		<u>7.5MG</u>	<u>A078246 005</u>	Feb 21, 2020
<u>AB</u>		<u>10MG</u>	<u>A078246 002</u>	Feb 27, 2009
<u>AB</u>		<u>15MG</u>	<u>A078246 003</u>	Feb 27, 2009
<u>AB</u>		<u>30MG</u>	<u>A078246 004</u>	Feb 27, 2009
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A208972 001</u>	Apr 16, 2019
<u>AB</u>		<u>7.5MG</u>	<u>A208972 002</u>	Apr 16, 2019
<u>AB</u>		<u>10MG</u>	<u>A208972 003</u>	Apr 16, 2019
<u>AB</u>		<u>15MG</u>	<u>A208972 004</u>	Apr 16, 2019
<u>AB</u>		<u>30MG</u>	<u>A208972 005</u>	Apr 16, 2019
<u>AB</u>	HERITAGE PHARMA	<u>5MG</u>	<u>A204582 001</u>	Sep 18, 2015
<u>AB</u>		<u>10MG</u>	<u>A204582 002</u>	Sep 18, 2015
<u>AB</u>		<u>15MG</u>	<u>A204582 003</u>	Sep 18, 2015
<u>AB</u>		<u>30MG</u>	<u>A204582 004</u>	Sep 18, 2015
<u>AB</u>	IMPAX LABS INC	<u>5MG</u>	<u>A074253 001</u>	Mar 28, 2001
<u>AB</u>		<u>10MG</u>	<u>A074253 002</u>	Mar 28, 2001

PRESCRIPTION DRUG PRODUCT LIST

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

<u>AB</u>		<u>15MG</u>	<u>A074253</u>	<u>003</u>	Mar 13, 2002
<u>AB</u>	INVENTIA HLTHCARE	<u>5MG</u>	<u>A209696</u>	<u>001</u>	May 03, 2018
<u>AB</u>		<u>7.5MG</u>	<u>A209696</u>	<u>002</u>	May 03, 2018
<u>AB</u>		<u>10MG</u>	<u>A209696</u>	<u>003</u>	May 03, 2018
<u>AB</u>		<u>15MG</u>	<u>A209696</u>	<u>004</u>	May 03, 2018
<u>AB</u>		<u>30MG</u>	<u>A209696</u>	<u>005</u>	May 03, 2018
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A076008</u>	<u>003</u>	Mar 01, 2002
<u>AB</u>		<u>7.5MG</u>	<u>A076008</u>	<u>002</u>	Jul 08, 2013
<u>AB</u>		<u>10MG</u>	<u>A076008</u>	<u>004</u>	Mar 01, 2002
<u>AB</u>		<u>15MG</u>	<u>A076008</u>	<u>005</u>	Mar 28, 2001
<u>AB</u>		<u>30MG</u>	<u>A076008</u>	<u>001</u>	Jun 28, 2001
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A075388</u>	<u>001</u>	May 09, 2002
<u>AB</u>		<u>10MG</u>	<u>A075388</u>	<u>002</u>	May 09, 2002
<u>AB</u>		<u>15MG</u>	<u>A075388</u>	<u>003</u>	May 09, 2002
<u>AB</u>		<u>30MG</u>	<u>A078302</u>	<u>001</u>	Dec 17, 2007
<u>AB</u>	RUBICON	<u>5MG</u>	<u>A075521</u>	<u>001</u>	Apr 05, 2002
<u>AB</u>		<u>7.5MG</u>	<u>A075521</u>	<u>004</u>	Mar 16, 2021
<u>AB</u>		<u>10MG</u>	<u>A075521</u>	<u>002</u>	Apr 05, 2002
<u>AB</u>		<u>15MG</u>	<u>A075521</u>	<u>003</u>	Apr 05, 2002
<u>AB</u>		<u>30MG</u>	<u>A075521</u>	<u>005</u>	Mar 16, 2021
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A202330</u>	<u>001</u>	Aug 25, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A202330</u>	<u>005</u>	Feb 17, 2017
<u>AB</u>		<u>10MG</u>	<u>A202330</u>	<u>002</u>	Aug 25, 2014
<u>AB</u>		<u>15MG</u>	<u>A202330</u>	<u>003</u>	Aug 25, 2014
<u>AB</u>		<u>30MG</u>	<u>A202330</u>	<u>004</u>	Aug 25, 2014
<u>AB</u>	TEVA	<u>5MG</u>	<u>A075022</u>	<u>001</u>	Feb 28, 2002
<u>AB</u>		<u>10MG</u>	<u>A075022</u>	<u>002</u>	Feb 28, 2002
<u>AB</u>		<u>15MG</u>	<u>A075022</u>	<u>003</u>	Feb 28, 2002
<u>AB</u>		<u>30MG</u>	<u>A075022</u>	<u>004</u>	Mar 25, 2004
<u>AB</u>	UNICHEM	<u>5MG</u>	<u>A210907</u>	<u>001</u>	Nov 14, 2019
<u>AB</u>		<u>10MG</u>	<u>A210907</u>	<u>002</u>	Nov 14, 2019
<u>AB</u>		<u>15MG</u>	<u>A210907</u>	<u>003</u>	Nov 14, 2019
<u>AB</u>		<u>30MG</u>	<u>A210907</u>	<u>004</u>	Nov 14, 2019
<u>AB</u>	YILING	<u>5MG</u>	<u>A202087</u>	<u>001</u>	Dec 16, 2015
<u>AB</u>		<u>10MG</u>	<u>A202087</u>	<u>002</u>	Dec 16, 2015
<u>AB</u>		<u>15MG</u>	<u>A202087</u>	<u>003</u>	Dec 16, 2015
<u>AB</u>		<u>30MG</u>	<u>A202087</u>	<u>004</u>	Dec 16, 2015
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A078888</u>	<u>001</u>	Feb 07, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A078888</u>	<u>005</u>	Mar 21, 2023
<u>AB</u>		<u>10MG</u>	<u>A078888</u>	<u>002</u>	Feb 07, 2014
<u>AB</u>		<u>15MG</u>	<u>A078888</u>	<u>003</u>	Feb 07, 2014
<u>AB</u>		<u>30MG</u>	<u>A078888</u>	<u>004</u>	Feb 07, 2014

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

<u>AP</u>	ACCORD HLTHCARE INC	<u>6MG/ML</u>	<u>A210148</u>	<u>001</u>	Feb 22, 2019
<u>AP</u>	AMNEAL	<u>6MG/ML</u>	<u>A209580</u>	<u>001</u>	Dec 18, 2017
<u>AP</u>	APOTEX	<u>6MG/ML</u>	<u>A210448</u>	<u>001</u>	May 07, 2019
<u>AP</u>	HOSPIRA	<u>6MG/ML</u>	<u>A205672</u>	<u>001</u>	Jul 31, 2018
<u>AP</u>	MEITHEAL	<u>6MG/ML</u>	<u>A212127</u>	<u>001</u>	Oct 23, 2020
<u>AP</u>	MYLAN INSTITUTIONAL	<u>6MG/ML</u>	<u>A208536</u>	<u>001</u>	Nov 20, 2017
<u>AP</u>	NEXUS	<u>6MG/ML</u>	<u>A207794</u>	<u>001</u>	Jan 14, 2019
<u>AP</u>	PHARMASCIENCE INC	<u>6MG/ML</u>	<u>A207050</u>	<u>001</u>	Mar 24, 2017
<u>AP</u>	SHILPA	<u>6MG/ML</u>	<u>A210931</u>	<u>001</u>	Apr 18, 2019

BUSULFEX

<u>AP</u>	+	OTSUKA PHARM	<u>6MG/ML</u>	<u>N020954</u>	<u>001</u>	Feb 04, 1999
TABLET; ORAL						
MYLERAN						
	+	WAYLIS THERAP	2MG	N009386	001	

BUTOCONAZOLE NITRATE

CREAM; VAGINAL

GYNAZOLE-1

!	PADAGIS ISRAEL	2%	A200923	001	May 18, 2012
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PRESCRIPTION DRUG PRODUCT LIST

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

<u>AP</u>	HIKMA	<u>2MG/ML</u>	<u>A075046 001</u>	Aug 12, 1998
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A078400 001</u>	May 01, 2009
<u>AP</u>		<u>2MG/ML</u>	<u>A078400 002</u>	May 01, 2009

BUTORPHANOL TARTRATE PRESERVATIVE FREE

<u>AP</u>	HIKMA	<u>1MG/ML</u>	<u>A075045 001</u>	Aug 12, 1998
<u>AP</u>		<u>2MG/ML</u>	<u>A075045 002</u>	Aug 12, 1998
<u>AP</u>	! HOSPIRA	<u>1MG/ML</u>	<u>A074626 001</u>	Jan 23, 1997
<u>AP</u>	!	<u>2MG/ML</u>	<u>A074626 002</u>	Jan 23, 1997

SPRAY, METERED; NASAL

BUTORPHANOL TARTRATE

<u>AB</u>	APOTEX	<u>1MG/SPRAY</u>	<u>A075499 001</u>	Dec 04, 2002
<u>AB</u>	HIKMA	<u>1MG/SPRAY</u>	<u>A075824 001</u>	Mar 12, 2002
<u>AB</u>	! RISING	<u>1MG/SPRAY</u>	<u>A075759 001</u>	Aug 08, 2001

CABAZITAXEL

SOLUTION; INTRAVENOUS

CABAZITAXEL

<u>AP</u>	ACCORD HLTHCARE	<u>60MG/1.5ML (40MG/ML)</u>	<u>A207693 001</u>	Oct 26, 2022
<u>AP</u>	DR REDDYS	<u>60MG/1.5ML (40MG/ML)</u>	<u>A207718 001</u>	Feb 10, 2023

JEVTANA KIT

<u>AP</u>	+! SANOFI AVENTIS US	<u>60MG/1.5ML (40MG/ML)</u>	<u>N201023 001</u>	Jun 17, 2010
	CABAZITAXEL			
	+! ACCORD HLTHCARE	60MG/3ML (20MG/ML)	N207949 001	Dec 29, 2021

CABERGOLINE

TABLET; ORAL

CABERGOLINE

<u>AB</u>	! INGENUS PHARMS LLC	<u>0.5MG</u>	<u>A204735 001</u>	Aug 01, 2018
<u>AB</u>	IVAX SUB TEVA	<u>0.5MG</u>	<u>A077750 001</u>	Mar 07, 2007
	PHARMS			
<u>AB</u>	STRIDES PHARMA	<u>0.5MG</u>	<u>A076310 001</u>	Dec 29, 2005

CABOTEGRAVIR

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

APRETUDE

	+! VIIV HLTHCARE	600MG/3ML (200MG/ML)	N215499 001	Dec 20, 2021
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CABOTEGRAVIR SODIUM

TABLET; ORAL

VOCABRIA

	+! VIIV HLTHCARE	EQ 30MG BASE	N212887 001	Jan 21, 2021
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CABOTEGRAVIR; RILPIVIRINE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

CABENUVA KIT

	+! VIIV HLTHCARE	400MG/2ML (200MG/ML); 600MG/2ML (300MG/ML)	N212888 001	Jan 21, 2021
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	+!	600MG/3ML (200MG/ML); 900MG/3ML (300MG/ML)	N212888 002	Jan 21, 2021
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CABOZANTINIB S-MALATE

CAPSULE; ORAL

COMETRIQ

	+ EXELIXIS	EQ 20MG BASE	N203756 001	Nov 29, 2012
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	+!	EQ 80MG BASE	N203756 002	Nov 29, 2012
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TABLET; ORAL

CABOMETYX

	+ EXELIXIS INC	EQ 20MG BASE	N208692 001	Apr 25, 2016
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	+	EQ 40MG BASE	N208692 002	Apr 25, 2016
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	+!	EQ 60MG BASE	N208692 003	Apr 25, 2016
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CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAF CIT

<u>AP</u>	+! HIKMA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>N020793 001</u>	Sep 21, 1999
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CAFFEINE CITRATE

<u>AP</u>	AM REGENT	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077906 001</u>	May 15, 2007
<u>AP</u>	EUGIA PHARMA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A205013 001</u>	Sep 22, 2015
<u>AP</u>	EXELA PHARMA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077233 001</u>	Sep 21, 2006
	SCIENCE			
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077997 001</u>	Jul 20, 2007
<u>AP</u>	MICRO LABS	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A207400 001</u>	Dec 14, 2017
<u>AP</u>	SAGENT PHARMS	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A090827 001</u>	Aug 29, 2012

PRESCRIPTION DRUG PRODUCT LIST

CAFFEINE CITRATE

SOLUTION;ORAL

CAFFEINE CITRATE

AA	!	EXELA PHARMA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077304 001</u>	Sep 21, 2006
AA		FRESENIUS KABI USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A078002 001</u>	Jan 31, 2008
AA		MICRO LABS	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A213202 001</u>	Dec 16, 2019
AA		SAGENT PHARMS	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A091102 001</u>	Aug 29, 2012
AA		SUN PHARM	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A090357 001</u>	Sep 30, 2009

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY;RECTAL

MIGERGOT

+	!	COSETTE	100MG;2MG	A086557 001	Oct 04, 1983
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TABLET;ORAL

ERGOTAMINE TARTRATE AND CAFFEINE

!		MIKART	100MG;1MG	A040590 001	Sep 16, 2005
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CALCIFEDIOL

CAPSULE, EXTENDED RELEASE;ORAL

RAYALDEE

+	!	EIRGEN	0.03MG	N208010 001	Jun 17, 2016
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CALCIPOTRIENE

AEROSOL, FOAM;TOPICAL

SORILUX

+	!	MAYNE PHARMA	0.005%	N022563 001	Oct 06, 2010
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CREAM;TOPICAL

CALCIPOTRIENE

AB		GLENMARK PHARMS	<u>0.005%</u>	<u>A205772 001</u>	Jun 09, 2015
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DOVONEX

AB	+	!	LEO PHARMA AS	<u>0.005%</u>	<u>N020554 001</u>	Jul 22, 1996
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OINTMENT;TOPICAL

CALCIPOTRIENE

AB	!		GLENMARK PHARMS INC	<u>0.005%</u>	<u>A090633 001</u>	Mar 24, 2010
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DOVONEX

AB	+		LEO PHARMA AS	<u>0.005%</u>	<u>N020273 001</u>	Dec 29, 1993
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SOLUTION;TOPICAL

CALCIPOTRIENE

AT			CHARTWELL RX	<u>0.005%</u>	<u>A207163 001</u>	Dec 26, 2017
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AT	!		COSETTE	<u>0.005%</u>	<u>A078468 001</u>	Mar 24, 2011
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AT			FOUGERA PHARMS	<u>0.005%</u>	<u>A078305 001</u>	May 06, 2008
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AT			HIKMA	<u>0.005%</u>	<u>A077579 001</u>	Nov 19, 2009
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CALCITONIN SALMON

INJECTABLE;INJECTION

CALCITONIN-SALMON

AP			CUSTOPHARM INC	<u>200 IU/ML</u>	<u>A212416 001</u>	May 14, 2021
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AP			PAR STERILE PRODUCTS	<u>200 IU/ML</u>	<u>A209358 001</u>	Nov 10, 2021
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MIACALCIN

AP	+	!	MYLAN IRELAND LTD	<u>200 IU/ML</u>	<u>N017808 002</u>	Mar 29, 1991
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SPRAY, METERED;NASAL

CALCITONIN-SALMON

!			APOTEX INC	200 IU/SPRAY	A076396 001	Nov 17, 2008
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CALCITRIOL

CAPSULE;ORAL

CALCITRIOL

AB			AMNEAL PHARMS	<u>0.25MCG</u>	<u>A203289 002</u>	Jun 14, 2017
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AB				<u>0.5MCG</u>	<u>A203289 001</u>	Jun 14, 2017
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AB			BIONPHARMA	<u>0.25MCG</u>	<u>A091174 001</u>	May 24, 2013
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AB				<u>0.5MCG</u>	<u>A091174 002</u>	May 24, 2013
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AB			HIKMA	<u>0.25MCG</u>	<u>A076917 001</u>	Mar 27, 2006
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AB			STRIDES PHARMA	<u>0.25MCG</u>	<u>A091356 001</u>	Dec 12, 2014
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AB	!			<u>0.5MCG</u>	<u>A091356 002</u>	Dec 12, 2014
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AB			SUN PHARM	<u>0.25MCG</u>	<u>A204556 001</u>	Feb 21, 2019
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AB				<u>0.5MCG</u>	<u>A204556 002</u>	Feb 21, 2019
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AB			TEVA	<u>0.25MCG</u>	<u>A075765 001</u>	Oct 12, 2001
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AB				<u>0.5MCG</u>	<u>A075765 002</u>	Oct 12, 2001
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ROCALTROL

AB	+		SS PHARMA	<u>0.25MCG</u>	<u>N018044 001</u>	
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AB	+			<u>0.5MCG</u>	<u>N018044 002</u>	
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INJECTABLE;INJECTION

CALCITRIOL

AP			GLAND PHARMA LTD	<u>0.001MG/ML</u>	<u>A211030 001</u>	Feb 03, 2020
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PRESCRIPTION DRUG PRODUCT LIST

CALCITRIOL

INJECTABLE; INJECTION

CALCITRIOL

AP ! LONG GROVE PHARMS **0.001MG/ML** **A078066 001** Jan 29, 2008
 OINTMENT; TOPICAL
 VECTICAL

+! GALDERMA LABS LP 3MCG/GM N022087 001 Jan 23, 2009

SOLUTION; ORAL

CALCITRIOL

AA HIKMA **1MCG/ML** **A076242 001** Jul 18, 2003

AA PATRIN **1MCG/ML** **A203973 001** Jul 28, 2023

AA RISING **1MCG/ML** **A209798 001** Nov 21, 2018

ROCALTROL

AA +! SS PHARMA **1MCG/ML** **N021068 001** Nov 20, 1998

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

AB CHARTWELL RX **667MG** **A091312 001** Jun 01, 2012

AB HERITAGE PHARMS INC **667MG** **A202315 001** Jun 29, 2015

AB HIKMA **667MG** **A077728 001** Feb 26, 2008

AB INVAGEN PHARMS **667MG** **A203135 001** Feb 07, 2013

AB LUPIN LTD **667MG** **A202127 001** Jul 09, 2015

AB NOSTRUM LABS INC **667MG** **A203179 001** Oct 26, 2015

AB SQUARE PHARMS **667MG** **A217205 001** Mar 13, 2023

AB SUVEN PHARMS **667MG** **A211038 001** Feb 21, 2020

PHOSLO GELCAPS

AB +! FRESENIUS MEDCL **667MG** **N021160 003** Apr 02, 2001

TABLET; ORAL

CALCIUM ACETATE

AB ! CHARTWELL MOLECULAR **667MG** **A202420 001** Feb 05, 2013

AB HERITAGE PHARMS INC **667MG** **A202885 001** Jan 22, 2015

AB PADAGIS US **667MG** **A091561 001** Apr 13, 2011

CALCIUM CHLORIDE

INJECTABLE; INJECTION

CALCIUM CHLORIDE 10%

AP AM REGENT **100MG/ML** **A209088 001** Jul 27, 2017

AP INTL MEDICATION SYS **100MG/ML** **A203477 001** May 09, 2018

AP MEDEFIL INC **100MG/ML** **A211553 001** May 01, 2019

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

AP +! HOSPIRA **100MG/ML** **N021117 001** Jan 28, 2000

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

BSS PLUS

+! ALCON 0.154MG/ML; 0.92MG/ML; 0.184MG/ML; 0.2MG/M L; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/M L N018469 001

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.0 5GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML ; 7.07GM/1000ML (5000ML) N021703 011 Oct 10, 2008

PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML) N021703 006 Oct 25, 2006

PRISMASOL BGK 2/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.0 3GM/1000ML; 0.157GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML (5000ML) N021703 002 Oct 25, 2006

PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 5.15GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; 0.157GM/1000ML; 3.09GM/100 0ML; 6.46GM/1000ML (5000ML) N021703 003 Oct 25, 2006

PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.4 4GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML (5000ML) N021703 015 Oct 10, 2008

PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09 N021703 004 Oct 25, 2006

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER

GM/1000ML; 6.46GM/1000ML (5000ML)

PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER

+	!	BAXTER HLTHCARE	N/A/1000ML; N/A/1000ML; 5.4GM/1000ML; 2.44	N021703	014	Oct 10, 2008
		CORP	GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46			
			GM/1000ML (5000ML)			

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5	N018883	001	Nov 30, 1984
			67MG/100ML; 392MG/100ML			

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5	N018883	004	Nov 30, 1984
			38MG/100ML; 448MG/100ML			

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5	N020171	001	Aug 19, 1992
			38MG/100ML; 448MG/100ML			

DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5	N018883	002	Nov 30, 1984
			67MG/100ML; 392MG/100ML			

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5	N018883	005	Nov 30, 1984
			38MG/100ML; 448MG/100ML			

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5	N020171	002	Aug 19, 1992
			38MG/100ML; 448MG/100ML			

DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 5	N018883	003	Nov 30, 1984
			567MG/100ML; 392MG/100ML			

DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5	N018883	006	Nov 30, 1984
			538MG/100ML; 448MG/100ML			

DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	+	FRESENIUS MEDCL	18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5	N020171	003	Aug 19, 1992
			538MG/100ML; 448MG/100ML			

DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5	N020183	001	Dec 04, 1992
			38MG/100ML; 448MG/100ML			

DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5	N020183	002	Dec 04, 1992
			38MG/100ML; 448MG/100ML			

DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5	N020183	004	Dec 04, 1992
			538MG/100ML; 448MG/100ML			

DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5	N017512	012	Jan 10, 1989
			38MG/100ML; 448MG/100ML			

DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5	N017512	013	Jul 11, 1990
			38MG/100ML; 448MG/100ML			

DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 5	N017512	014	Jul 11, 1990
			38MG/100ML; 448MG/100ML			

DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	18.3MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5	N017512	015	Jul 11, 1990
			538MG/100ML; 448MG/100ML			

DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5	N017512	004	
			38MG/100ML; 448MG/100ML			

AT	+		25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5	N020163	001	Dec 04, 1992
			38MG/100ML; 448MG/100ML			

DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5	N017512	005	
			38MG/100ML; 448MG/100ML			

AT	+		25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5	N020163	002	Dec 04, 1992
			38MG/100ML; 448MG/100ML			

DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5	N017512	006	
			538MG/100ML; 448MG/100ML			

AT	+		25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5	N020163	003	Dec 04, 1992
			538MG/100ML; 448MG/100ML			

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INTRATHECAL

ELLIOTTS B SOLUTION

+! LUKARE MEDICAL LLC 0.2MG/ML;0.8MG/ML;0.3MG/ML;0.3MG/ML;1.9 MG/ML;7.3MG/ML;0.2MG/ML N020577 001 Sep 27, 1996

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP + ICU MEDICAL INC 20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML N017608 001

DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER

AP B BRAUN 20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML N019634 003 Feb 24, 1988

LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML N016679 001

POTASSIUM CHLORIDE 15MEO IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;254MG/100ML;600MG/ 100ML;310MG/100ML N019367 006 Apr 05, 1985

POTASSIUM CHLORIDE 20MEO IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;179MG/100ML;600MG/ 100ML;310MG/100ML N019367 004 Apr 05, 1985

AP 20MG/100ML;5GM/100ML;328MG/100ML;600MG/ 100ML;310MG/100ML N019367 005 Apr 05, 1985

AP + ICU MEDICAL INC 20MG/100ML;5GM/100ML;179MG/100ML;600MG/ 100ML;310MG/100ML N019685 002 Oct 17, 1988

POTASSIUM CHLORIDE 30MEO IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;254MG/100ML;600MG/ 100ML;310MG/100ML N019367 007 Apr 05, 1985

POTASSIUM CHLORIDE 40MEO IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;328MG/100ML;600MG/ 100ML;310MG/100ML N019367 008 Apr 05, 1985

DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER

B BRAUN 10MG/100ML;2.5GM/100ML;15MG/100ML;300MG /100ML;160MG/100ML N019634 001 Feb 24, 1988

POTASSIUM CHLORIDE 10MEO IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML;5GM/100ML;105MG/100ML;600MG/ 100ML;310MG/100ML N019367 002 Apr 05, 1985

20MG/100ML;5GM/100ML;179MG/100ML;600MG/ 100ML;310MG/100ML N019367 003 Apr 05, 1985

POTASSIUM CHLORIDE 5MEO IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML;5GM/100ML;105MG/100ML;600MG/ 100ML;310MG/100ML N019367 001 Apr 05, 1985

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

+! HOSPIRA 16.5MG/ML;25.4MG/ML;74.6MG/ML;121MG/ML; 16.1MG/ML N018895 001 Jul 20, 1984

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

SOLUTION; IRRIGATION

BALANCED SALT

AT B BRAUN 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6 .4MG/ML;1.7MG/ML A091387 001 Feb 03, 2010

BSS

AT +! ALCON 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6 .4MG/ML;1.7MG/ML N020742 001 Dec 10, 1997

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

INJECTABLE; INJECTION

PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML;3.05GM/1000ML;0.314GM/1000ML N207026 002 Jan 13, 2015
CORP ;2.21GM/1000ML;6.95GM/1000ML;0.187GM/10
00ML (5000ML)

PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 3.68GM/1000ML;3.05GM/1000ML;0.314GM/10 0ML N207026 001 Jan 13, 2015
CORP ;3.09GM/1000ML;6.34GM/1000ML;0.187GM/10

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

SOLUTION; PERFUSION, CARDIAC

CARDIOPLEGIC IN PLASTIC CONTAINER

<u>AT</u>	BAXTER HLTHCARE	<u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u>	<u>A075323 001</u>	Apr 21, 2000
<u>AT</u>	FRESENIUS KABI USA	<u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u>	<u>A214623 001</u>	Feb 18, 2022

PLEGISOL IN PLASTIC CONTAINER

<u>AT</u>	+! HOSPIRA	<u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u>	<u>N018608 001</u>	Feb 26, 1982
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CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N020002 001</u>	Apr 17, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N016693 001</u>	
<u>AP</u>	ICU MEDICAL INC	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N018251 001</u>	

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

<u>AT</u>	B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N018156 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N018495 001</u>	Feb 19, 1982
<u>AT</u>	ICU MEDICAL INC	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N017635 001</u>	

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N019632 001</u>	Feb 29, 1988
<u>AP</u>	+! BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N016682 001</u>	
<u>AP</u>	FRESENIUS KABI USA	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>A209338 001</u>	Jan 28, 2019
<u>AP</u>	ICU MEDICAL INC	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N017641 001</u>	

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AT</u>	+! B BRAUN	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N018681 001</u>	Dec 27, 1982
<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N018494 001</u>	Feb 19, 1982
<u>AT</u>	+!	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N018921 001</u>	Apr 03, 1984
<u>AT</u>	+! ICU MEDICAL INC	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N019416 001</u>	Jan 17, 1986

CALCIUM GLUCONATE

SOLUTION; INTRAVENOUS

CALCIUM GLUCONATE

<u>AP</u>	B BRAUN MEDICAL INC	<u>10GM/100ML (100MG/ML)</u>	<u>A216541 001</u>	Aug 21, 2023
<u>AP</u>	+! FRESENIUS KABI USA	<u>1GM/10ML (100MG/ML)</u>	<u>N208418 001</u>	Jun 15, 2017
<u>AP</u>	+!	<u>5GM/50ML (100MG/ML)</u>	<u>N208418 002</u>	Jun 15, 2017
<u>AP</u>	+!	<u>10GM/100ML (100MG/ML)</u>	<u>N208418 003</u>	Jun 15, 2017
<u>AP</u>	NIVAGEN PHARMS INC	<u>1GM/10ML (100MG/ML)</u>	<u>A213071 001</u>	Oct 14, 2022
<u>AP</u>		<u>5GM/50ML (100MG/ML)</u>	<u>A213071 002</u>	Oct 19, 2023
<u>AP</u>	SOMERSET	<u>5GM/50ML (100MG/ML)</u>	<u>A217689 001</u>	Oct 19, 2023
<u>AP</u>		<u>10GM/100ML (100MG/ML)</u>	<u>A217689 002</u>	Oct 19, 2023

CALCIUM GLUCONATE IN SODIUM CHLORIDE

<u>AP</u>	AMNEAL	<u>1GM/50ML (20MG/ML)</u>	<u>A217174 001</u>	Sep 05, 2023
<u>AP</u>		<u>2GM/100ML (20MG/ML)</u>	<u>A217174 002</u>	Sep 05, 2023
<u>AP</u>	+! FRESENIUS KABI USA	<u>1GM/50ML (20MG/ML)</u>	<u>N208418 004</u>	Jun 17, 2021
<u>AP</u>	+!	<u>2GM/100ML (20MG/ML)</u>	<u>N208418 005</u>	Jun 17, 2021
	+! HQ SPCLT PHARMA	1GM/50ML (20MG/ML)	N210906 001	Oct 29, 2018
	+!	1GM/100ML (10MG/ML)	N210906 003	Jun 04, 2021
	+!	2GM/100ML (20MG/ML)	N210906 002	Oct 29, 2018

CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE

SOLUTION; ORAL

XYWAV

	+! JAZZ	0.234GM/ML; 0.096GM/ML; 0.13GM/ML; 0.04GM/ML	N212690 001	Jul 21, 2020
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PRESCRIPTION DRUG PRODUCT LIST

CANAGLIFLOZIN

TABLET; ORAL

INVOKANA

+	JANSSEN PHARMS	100MG	N204042	001	Mar 29, 2013
+	!	300MG	N204042	002	Mar 29, 2013

CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

INVOKAMET

+	JANSSEN PHARMS	50MG; 500MG	N204353	001	Aug 08, 2014
+		50MG; 1GM	N204353	002	Aug 08, 2014
+		150MG; 500MG	N204353	003	Aug 08, 2014
+	!	150MG; 1GM	N204353	004	Aug 08, 2014

TABLET, EXTENDED RELEASE; ORAL

INVOKAMET XR

+	JANSSEN PHARMS	50MG; 500MG	N205879	001	Sep 20, 2016
+		50MG; 1GM	N205879	002	Sep 20, 2016
+		150MG; 500MG	N205879	003	Sep 20, 2016
+	!	150MG; 1GM	N205879	004	Sep 20, 2016

CANDESARTAN CILEXETIL

TABLET; ORAL

ATACAND

<u>AB</u>	+	ANI PHARMS	<u>4MG</u>	<u>N020838</u>	<u>001</u>	Jun 04, 1998
<u>AB</u>	+		<u>8MG</u>	<u>N020838</u>	<u>002</u>	Jun 04, 1998
<u>AB</u>	+		<u>16MG</u>	<u>N020838</u>	<u>003</u>	Jun 04, 1998
<u>AB</u>	+	!	<u>32MG</u>	<u>N020838</u>	<u>004</u>	Jun 04, 1998

CANDESARTAN CILEXETIL

<u>AB</u>		ALEMBIC	<u>4MG</u>	<u>A210302</u>	<u>001</u>	Dec 04, 2018
<u>AB</u>			<u>8MG</u>	<u>A210302</u>	<u>002</u>	Dec 04, 2018
<u>AB</u>			<u>16MG</u>	<u>A210302</u>	<u>003</u>	Dec 04, 2018
<u>AB</u>			<u>32MG</u>	<u>A209119</u>	<u>001</u>	Jun 20, 2017
<u>AB</u>		MACLEODS PHARMS LTD	<u>4MG</u>	<u>A203813</u>	<u>001</u>	Dec 05, 2016
<u>AB</u>			<u>8MG</u>	<u>A203813</u>	<u>002</u>	Dec 05, 2016
<u>AB</u>			<u>16MG</u>	<u>A203813</u>	<u>003</u>	Dec 05, 2016
<u>AB</u>			<u>32MG</u>	<u>A203813</u>	<u>004</u>	Dec 05, 2016
<u>AB</u>		MYLAN	<u>4MG</u>	<u>A078702</u>	<u>001</u>	May 03, 2013
<u>AB</u>			<u>8MG</u>	<u>A078702</u>	<u>002</u>	May 03, 2013
<u>AB</u>			<u>16MG</u>	<u>A078702</u>	<u>003</u>	May 03, 2013
<u>AB</u>			<u>32MG</u>	<u>A078702</u>	<u>004</u>	May 03, 2013
<u>AB</u>		PRINSTON INC	<u>8MG</u>	<u>A206233</u>	<u>001</u>	Aug 21, 2023
<u>AB</u>			<u>16MG</u>	<u>A206233</u>	<u>002</u>	Aug 21, 2023
<u>AB</u>			<u>32MG</u>	<u>A206233</u>	<u>003</u>	Aug 21, 2023
<u>AB</u>		ZYDUS LIFESCIENCES	<u>4MG</u>	<u>A091390</u>	<u>001</u>	Aug 23, 2017
<u>AB</u>			<u>8MG</u>	<u>A091390</u>	<u>002</u>	Aug 23, 2017
<u>AB</u>			<u>16MG</u>	<u>A091390</u>	<u>003</u>	Aug 23, 2017
<u>AB</u>			<u>32MG</u>	<u>A091390</u>	<u>004</u>	Aug 23, 2017

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ATACAND HCT

<u>AB</u>	+	ANI PHARMS	<u>16MG; 12.5MG</u>	<u>N021093</u>	<u>001</u>	Sep 05, 2000
<u>AB</u>	+		<u>32MG; 12.5MG</u>	<u>N021093</u>	<u>002</u>	Sep 05, 2000
<u>AB</u>	+	!	<u>32MG; 25MG</u>	<u>N021093</u>	<u>003</u>	May 16, 2008

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>		DR REDDYS LABS LTD	<u>16MG; 12.5MG</u>	<u>A202965</u>	<u>001</u>	Jun 03, 2013
<u>AB</u>			<u>32MG; 12.5MG</u>	<u>A202965</u>	<u>002</u>	Jun 03, 2013
<u>AB</u>			<u>32MG; 25MG</u>	<u>A202965</u>	<u>003</u>	Jun 03, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>16MG; 12.5MG</u>	<u>A204100</u>	<u>001</u>	Feb 27, 2015
<u>AB</u>			<u>32MG; 12.5MG</u>	<u>A204100</u>	<u>002</u>	Feb 27, 2015
<u>AB</u>			<u>32MG; 25MG</u>	<u>A204100</u>	<u>003</u>	Feb 27, 2015
<u>AB</u>		MYLAN	<u>16MG; 12.5MG</u>	<u>A090704</u>	<u>001</u>	Dec 04, 2012
<u>AB</u>			<u>32MG; 12.5MG</u>	<u>A090704</u>	<u>002</u>	Dec 04, 2012
<u>AB</u>			<u>32MG; 25MG</u>	<u>A090704</u>	<u>003</u>	Dec 04, 2012
<u>AB</u>		PRINSTON INC	<u>16MG; 12.5MG</u>	<u>A207455</u>	<u>001</u>	Apr 11, 2018
<u>AB</u>			<u>32MG; 12.5MG</u>	<u>A207455</u>	<u>002</u>	Apr 11, 2018
<u>AB</u>			<u>32MG; 25MG</u>	<u>A207455</u>	<u>003</u>	Apr 11, 2018
<u>AB</u>		ZYDUS LIFESCIENCES	<u>16MG; 12.5MG</u>	<u>A203466</u>	<u>001</u>	Nov 27, 2017
<u>AB</u>			<u>32MG; 12.5MG</u>	<u>A203466</u>	<u>002</u>	Nov 27, 2017
<u>AB</u>			<u>32MG; 25MG</u>	<u>A203466</u>	<u>003</u>	Nov 27, 2017

PRESCRIPTION DRUG PRODUCT LIST

CANGRELOR

POWDER; INTRAVENOUS

KENGREAL

+! CHIESI

50MG/VIAL

N204958 001 Jun 22, 2015

CANNABIDIOL

SOLUTION; ORAL

EPIDIOLEX

+! JAZZ PHARMS RES

100MG/ML

N210365 001 Sep 28, 2018

CANTHARIDIN

SOLUTION; TOPICAL

YCANTH

+! VERRICA PHARMS

0.7%

N212905 001 Jul 21, 2023

CAPECITABINE

TABLET; ORAL

CAPECITABINE

<u>AB</u>	ACCORD HLTHCARE	<u>150MG</u>	<u>A202593 001</u>	Apr 23, 2015
<u>AB</u>		<u>500MG</u>	<u>A202593 002</u>	Apr 23, 2015
<u>AB</u>	ALKEM LABS LTD	<u>150MG</u>	<u>A207652 001</u>	Nov 24, 2017
<u>AB</u>		<u>500MG</u>	<u>A207652 002</u>	Nov 24, 2017
<u>AB</u>	DR REDDYS	<u>150MG</u>	<u>A204345 001</u>	Dec 04, 2020
<u>AB</u>		<u>500MG</u>	<u>A204345 002</u>	Dec 04, 2020
<u>AB</u>	EUGIA PHARMA	<u>150MG</u>	<u>A210604 001</u>	Apr 17, 2018
<u>AB</u>		<u>500MG</u>	<u>A210604 002</u>	Apr 17, 2018
<u>AB</u>	HIKMA	<u>150MG</u>	<u>A200483 001</u>	Jul 14, 2016
<u>AB</u>		<u>500MG</u>	<u>A200483 002</u>	Jul 14, 2016
<u>AB</u>	MSN	<u>150MG</u>	<u>A209365 001</u>	Jul 02, 2018
<u>AB</u>		<u>500MG</u>	<u>A209365 002</u>	Jul 02, 2018
<u>AB</u>	RELIANCE LIFE	<u>150MG</u>	<u>A211724 001</u>	Apr 27, 2020
<u>AB</u>		<u>500MG</u>	<u>A211724 002</u>	Apr 27, 2020
<u>AB</u>	RISING	<u>150MG</u>	<u>A090943 001</u>	Aug 08, 2014
<u>AB</u>		<u>500MG</u>	<u>A090943 002</u>	Aug 08, 2014
<u>AB</u>	SHILPA	<u>150MG</u>	<u>A207456 001</u>	Dec 12, 2016
<u>AB</u>		<u>500MG</u>	<u>A207456 002</u>	Dec 12, 2016
<u>AB</u>	SUN PHARM	<u>150MG</u>	<u>A204668 001</u>	Jun 21, 2019
<u>AB</u>		<u>500MG</u>	<u>A204668 002</u>	Jun 21, 2019
<u>AB</u>	TEVA PHARMS USA	<u>150MG</u>	<u>A091649 001</u>	Sep 16, 2013
<u>AB</u>		<u>500MG</u>	<u>A091649 002</u>	Sep 16, 2013
<u>AB</u>	TEYRO LABS	<u>150MG</u>	<u>A217237 001</u>	Oct 23, 2023
<u>AB</u>		<u>500MG</u>	<u>A217237 002</u>	Oct 23, 2023
	<u>XELODA</u>			
<u>AB</u>	+ CHEPLAPHARM	<u>150MG</u>	<u>N020896 001</u>	Apr 30, 1998
<u>AB</u>	+!	<u>500MG</u>	<u>N020896 002</u>	Apr 30, 1998

CAPIVASERTIB

TABLET; ORAL

TRUQAP

+ ASTRAZENECA

160MG

N218197 001 Nov 16, 2023

+!

200MG

N218197 002 Nov 16, 2023

CAPMATINIB HYDROCHLORIDE

TABLET; ORAL

TABRECTA

+ NOVARTIS PHARM

EQ 150MG BASE

N213591 001 May 06, 2020

+!

EQ 200MG BASE

N213591 002 May 06, 2020

CAPSAICIN

PATCH; TOPICAL

QUTENZA

+! AVERITAS

8%

N022395 001 Nov 16, 2009

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

<u>AB</u>	AJANTA PHARMA LTD	<u>12.5MG</u>	<u>A212809 001</u>	Dec 13, 2019
<u>AB</u>		<u>25MG</u>	<u>A212809 002</u>	Dec 13, 2019
<u>AB</u>		<u>50MG</u>	<u>A212809 003</u>	Dec 13, 2019
<u>AB</u>		<u>100MG</u>	<u>A212809 004</u>	Dec 13, 2019
<u>AB</u>	CHANGZHOU PHARM	<u>12.5MG</u>	<u>A214442 001</u>	Jan 27, 2023
<u>AB</u>		<u>25MG</u>	<u>A214442 002</u>	Jan 27, 2023
<u>AB</u>		<u>50MG</u>	<u>A214442 003</u>	Jan 27, 2023
<u>AB</u>		<u>100MG</u>	<u>A214442 004</u>	Jan 27, 2023
<u>AB</u>	COREPHARMA	<u>12.5MG</u>	<u>A074737 001</u>	Oct 28, 1998
<u>AB</u>		<u>25MG</u>	<u>A074737 002</u>	Oct 28, 1998

PRESCRIPTION DRUG PRODUCT LIST

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

<u>AB</u>		<u>50MG</u>	<u>A074737 003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>A074737 004</u>	Oct 28, 1998
<u>AB</u>	HIKMA INTL PHARMS	<u>12.5MG</u>	<u>A074505 001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>A074505 002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>A074505 003</u>	Feb 13, 1996
<u>AB</u>	!	<u>100MG</u>	<u>A074505 004</u>	Feb 13, 1996
<u>AB</u>	KENTON	<u>12.5MG</u>	<u>A074677 004</u>	May 30, 1997
<u>AB</u>		<u>25MG</u>	<u>A074677 002</u>	May 30, 1997
<u>AB</u>		<u>50MG</u>	<u>A074677 001</u>	May 30, 1997
<u>AB</u>		<u>100MG</u>	<u>A074677 003</u>	May 30, 1997
<u>AB</u>	PRINSTON INC	<u>12.5MG</u>	<u>A074477 001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>A074477 002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>A074477 003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>A074477 004</u>	Feb 13, 1996
<u>AB</u>	WOCKHARDT	<u>12.5MG</u>	<u>A074532 001</u>	Mar 28, 1997
<u>AB</u>		<u>25MG</u>	<u>A074532 002</u>	Mar 28, 1997
<u>AB</u>		<u>50MG</u>	<u>A074532 003</u>	Mar 28, 1997
<u>AB</u>		<u>100MG</u>	<u>A074532 004</u>	Mar 28, 1997

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

	RISING	25MG; 15MG	A074896 001	Dec 29, 1997
!		25MG; 25MG	A074896 002	Dec 29, 1997
!		50MG; 15MG	A074896 004	Dec 29, 1997
		50MG; 25MG	A074896 003	Dec 29, 1997

CARBACHOL

SOLUTION; INTRAOCULAR

MIOSTAT

+	ALCON	0.01%	N016968 001	
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CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A078986 001</u>	Nov 25, 2011
<u>AB</u>		<u>200MG</u>	<u>A078986 002</u>	Nov 25, 2011
<u>AB</u>		<u>300MG</u>	<u>A078986 003</u>	Nov 25, 2011
<u>AB</u>	NOSTRUM LABS INC	<u>100MG</u>	<u>A076697 001</u>	May 20, 2011
<u>AB</u>		<u>200MG</u>	<u>A076697 002</u>	May 20, 2011
<u>AB</u>		<u>300MG</u>	<u>A076697 003</u>	May 20, 2011
<u>AB</u>	TARO	<u>100MG</u>	<u>A201106 001</u>	Jun 21, 2013
<u>AB</u>		<u>200MG</u>	<u>A201106 002</u>	Jun 21, 2013
<u>AB</u>		<u>300MG</u>	<u>A201106 003</u>	Jun 21, 2013
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>A078592 001</u>	Sep 20, 2012
<u>AB</u>		<u>200MG</u>	<u>A078592 002</u>	Sep 20, 2012
<u>AB</u>		<u>300MG</u>	<u>A078592 003</u>	Sep 20, 2012

CARBATROL

<u>AB</u>	+	TAKEDA PHARMS USA	<u>100MG</u>	<u>N020712 003</u>	Sep 30, 1997
<u>AB</u>	+		<u>200MG</u>	<u>N020712 001</u>	Sep 30, 1997
<u>AB</u>	+	!	<u>300MG</u>	<u>N020712 002</u>	Sep 30, 1997
		EQUETRO			
	+	VALIDUS PHARMS	100MG	N021710 001	Dec 10, 2004
	+		200MG	N021710 002	Dec 10, 2004
	+		300MG	N021710 003	Dec 10, 2004

SUSPENSION; ORAL

CARBAMAZEPINE

<u>AB</u>	CHARTWELL RX	<u>100MG/5ML</u>	<u>A075714 001</u>	Jun 05, 2002
<u>AB</u>	NOVITIUM PHARMA	<u>100MG/5ML</u>	<u>A214277 001</u>	Oct 06, 2022

TEGRETOL

<u>AB</u>	+	NOVARTIS	<u>100MG/5ML</u>	<u>N018927 001</u>	Dec 18, 1987
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TERIL

<u>AB</u>	TARO	<u>100MG/5ML</u>	<u>A076729 001</u>	Sep 20, 2004
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TABLET; ORAL

CARBAMAZEPINE

<u>AB</u>	APOTEX INC	<u>200MG</u>	<u>A075948 001</u>	Feb 27, 2002
<u>AB</u>	TARO	<u>200MG</u>	<u>A074649 001</u>	Oct 03, 1996
<u>AB</u>	TORRENT PHARMS	<u>100MG</u>	<u>A077272 001</u>	Dec 07, 2005
<u>AB</u>		<u>200MG</u>	<u>A077272 002</u>	Dec 07, 2005
<u>AB</u>	UMEDICA	<u>100MG</u>	<u>A207798 001</u>	Apr 15, 2020
<u>AB</u>		<u>200MG</u>	<u>A207798 002</u>	Apr 15, 2020

PRESCRIPTION DRUG PRODUCT LIST

CARBAMAZEPINE

TABLET;ORAL

CARBAMAZEPINE

AB	UNICHEM	200MG	A213284 001	Aug 22, 2022
	EPITOL			
AB	TEVA	200MG	A070541 001	Sep 17, 1986
	TEGRETOL			
AB	+! NOVARTIS	200MG	N016608 001	
	CARBAMAZEPINE			
	TORRENT PHARMS	300MG	A077272 003	Dec 07, 2005
		400MG	A077272 004	Dec 07, 2005

TABLET, CHEWABLE;ORAL

CARBAMAZEPINE

AB	TARO PHARM INDS	100MG	A075687 001	Oct 24, 2000
AB	TORRENT PHARMS	100MG	A075712 001	Jul 05, 2001
	EPITOL			
AB	TEVA	100MG	A073524 001	Jul 29, 1992
	CARBAMAZEPINE			
	! TARO PHARM INDS	200MG	A075687 002	Jul 29, 2002

TABLET, EXTENDED RELEASE;ORAL

CARBAMAZEPINE

AB	AMNEAL PHARMS	100MG	A212704 001	Sep 22, 2023
AB		200MG	A212704 002	Sep 22, 2023
AB		400MG	A212704 003	Sep 22, 2023
AB	ANBISON LAB	100MG	A212948 001	Sep 30, 2021
AB		200MG	A212948 002	Sep 30, 2021
AB		400MG	A212948 003	Sep 30, 2021
AB	CSPC OUYI	100MG	A213311 001	Apr 13, 2021
AB		200MG	A213311 002	Apr 13, 2021
AB		400MG	A213311 003	Apr 13, 2021
AB	RICONPHARMA LLC	100MG	A216404 001	Dec 02, 2022
AB		200MG	A216404 002	Dec 02, 2022
AB		400MG	A216404 003	Dec 02, 2022
AB	SCIECURE PHARMA INC	100MG	A216235 001	Mar 02, 2023
AB		200MG	A216235 002	Mar 02, 2023
AB		400MG	A216235 003	Mar 02, 2023
AB	TARO	100MG	A078115 001	Mar 31, 2009
AB		200MG	A078115 002	Mar 31, 2009
AB		400MG	A078115 003	Mar 31, 2009
AB	UMEDICA	100MG	A216594 001	Aug 18, 2022
AB		200MG	A216594 002	Aug 18, 2022
AB		400MG	A216594 003	Aug 18, 2022
AB	UNIQUE PHARM	100MG	A211623 001	Apr 24, 2020
AB		200MG	A211623 002	Apr 24, 2020
AB		400MG	A211623 003	Apr 24, 2020
AB	ZHEJIANG JIUZHOU	100MG	A215591 001	Mar 31, 2022
AB		200MG	A215591 002	Mar 31, 2022
AB		400MG	A215591 003	Mar 31, 2022
AB	ZYDUS PHARMS	100MG	A205571 001	Feb 07, 2019
AB		200MG	A205571 002	Feb 07, 2019
AB		400MG	A205571 003	Feb 07, 2019
	TEGRETOL-XR			
AB	+ NOVARTIS	100MG	N020234 001	Mar 25, 1996
AB	+	200MG	N020234 002	Mar 25, 1996
AB	+!	400MG	N020234 003	Mar 25, 1996

CARBIDOPA

TABLET;ORAL

CARBIDOPA

AB	ALVOGEN	25MG	A204291 001	Jan 08, 2016
AB	AUROBINDO PHARMA	25MG	A211055 001	Oct 21, 2019
AB	BEXIMCO PHARMS USA	25MG	A217961 001	Dec 11, 2023
AB	EDENBRIDGE PHARMS	25MG	A205304 001	Feb 17, 2016
AB	NOVEL LABS INC	25MG	A204763 001	Oct 20, 2017
AB	ZYDUS PHARMS	25MG	A209910 001	May 07, 2018
	LODOSYN			
AB	+! ATON	25MG	N017830 001	

PRESCRIPTION DRUG PRODUCT LIST

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

<u>AB</u>	RISING	<u>12.5MG;200MG;50MG</u>	<u>A213212 001</u>	Jan 25, 2022
<u>AB</u>		<u>18.75MG;200MG;75MG</u>	<u>A213212 002</u>	Jan 25, 2022
<u>AB</u>		<u>25MG;200MG;100MG</u>	<u>A213212 003</u>	Jan 25, 2022
<u>AB</u>		<u>31.25MG;200MG;125MG</u>	<u>A213212 004</u>	Jan 25, 2022
<u>AB</u>		<u>37.5MG;200MG;150MG</u>	<u>A213212 005</u>	Jan 25, 2022
<u>AB</u>		<u>50MG;200MG;200MG</u>	<u>A213212 006</u>	Jan 25, 2022
<u>AB</u>	SUN PHARM	<u>25MG;200MG;100MG</u>	<u>A079085 001</u>	May 10, 2012
<u>AB</u>		<u>37.5MG;200MG;150MG</u>	<u>A079085 002</u>	May 10, 2012
	<u>STALEVO 100</u>			
<u>AB</u>	+ ORION PHARMA	<u>25MG;200MG;100MG</u>	<u>N021485 002</u>	Jun 11, 2003
	<u>STALEVO 125</u>			
<u>AB</u>	+ ORION PHARMA	<u>31.25MG;200MG;125MG</u>	<u>N021485 006</u>	Aug 29, 2008
	<u>STALEVO 150</u>			
<u>AB</u>	+ ORION PHARMA	<u>37.5MG;200MG;150MG</u>	<u>N021485 003</u>	Jun 11, 2003
	<u>STALEVO 200</u>			
<u>AB</u>	+! ORION PHARMA	<u>50MG;200MG;200MG</u>	<u>N021485 004</u>	Aug 02, 2007
	<u>STALEVO 50</u>			
<u>AB</u>	+! ORION PHARMA	<u>12.5MG;200MG;50MG</u>	<u>N021485 001</u>	Jun 11, 2003
	<u>STALEVO 75</u>			
<u>AB</u>	+ ORION PHARMA	<u>18.75MG;200MG;75MG</u>	<u>N021485 005</u>	Aug 29, 2008

CARBIDOPA; LEVODOPA

CAPSULE, EXTENDED RELEASE; ORAL

RYTARY

	+ IMPAX LABS INC	23.75MG;95MG	N203312 001	Jan 07, 2015
		36.25MG;145MG	N203312 002	Jan 07, 2015
		48.75MG;195MG	N203312 003	Jan 07, 2015
	+!	61.25MG;245MG	N203312 004	Jan 07, 2015

SUSPENSION; ENTERAL

DUOPA

	+! ABBVIE	4.63MG/ML;20MG/ML	N203952 001	Jan 09, 2015
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TABLET; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG;100MG</u>	<u>A074260 001</u>	Sep 03, 1993
<u>AB</u>		<u>25MG;100MG</u>	<u>A074260 002</u>	Sep 03, 1993
<u>AB</u>	!	<u>25MG;250MG</u>	<u>A074260 003</u>	Sep 03, 1993
<u>AB</u>	APOTEX INC	<u>10MG;100MG</u>	<u>A077120 001</u>	Jun 02, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A077120 002</u>	Jun 02, 2008
<u>AB</u>		<u>25MG;250MG</u>	<u>A077120 003</u>	Jun 02, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG;100MG</u>	<u>A216537 001</u>	Nov 28, 2022
<u>AB</u>		<u>25MG;100MG</u>	<u>A216537 002</u>	Nov 28, 2022
<u>AB</u>		<u>25MG;250MG</u>	<u>A216537 003</u>	Nov 28, 2022
<u>AB</u>	DR REDDYS LABS SA	<u>10MG;100MG</u>	<u>A073618 001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;100MG</u>	<u>A073589 001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;250MG</u>	<u>A073607 001</u>	Aug 28, 1992
<u>AB</u>	MYLAN	<u>10MG;100MG</u>	<u>A090324 001</u>	Sep 28, 2009
<u>AB</u>		<u>25MG;100MG</u>	<u>A090324 002</u>	Sep 28, 2009
<u>AB</u>		<u>25MG;250MG</u>	<u>A090324 003</u>	Sep 28, 2009
<u>AB</u>	RUBICON	<u>10MG;100MG</u>	<u>A216505 001</u>	Sep 21, 2022
<u>AB</u>		<u>25MG;100MG</u>	<u>A216505 002</u>	Sep 21, 2022
<u>AB</u>		<u>25MG;250MG</u>	<u>A216505 003</u>	Sep 21, 2022
<u>AB</u>	SCIEGEN PHARMS INC	<u>10MG;100MG</u>	<u>A214092 001</u>	May 07, 2021
<u>AB</u>		<u>25MG;100MG</u>	<u>A214092 002</u>	May 07, 2021
<u>AB</u>		<u>25MG;250MG</u>	<u>A214092 003</u>	May 07, 2021
<u>AB</u>	SUN PHARM INDS	<u>10MG;100MG</u>	<u>A078536 001</u>	Oct 28, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A078536 002</u>	Oct 28, 2008
<u>AB</u>		<u>25MG;250MG</u>	<u>A078536 003</u>	Oct 28, 2008

SINEMET

<u>AB</u>	+ ORGANON	<u>10MG;100MG</u>	<u>N017555 001</u>	
<u>AB</u>		<u>25MG;100MG</u>	<u>N017555 003</u>	
<u>AB</u>		<u>25MG;250MG</u>	<u>N017555 002</u>	
	DHIVY			
	+! AVION PHARMS	25MG;100MG	N214869 001	Nov 12, 2021

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	ACCORD HLTHCARE	<u>25MG;100MG</u>	<u>A202323 001</u>	Feb 08, 2013
<u>AB</u>		<u>50MG;200MG</u>	<u>A202323 002</u>	Feb 08, 2013
<u>AB</u>	ALEMBIC	<u>25MG;100MG</u>	<u>A210341 001</u>	Jun 05, 2019
<u>AB</u>		<u>50MG;200MG</u>	<u>A210341 002</u>	Jun 05, 2019

PRESCRIPTION DRUG PRODUCT LIST

CARBIDOPA; LEVODOPA

TABLET, EXTENDED RELEASE;ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	APOTEX	<u>25MG;100MG</u>	<u>A076212</u>	<u>001</u>	Jun 16, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>A076212</u>	<u>002</u>	Jun 16, 2004
<u>AB</u>	IMPAX LABS	<u>25MG;100MG</u>	<u>A076521</u>	<u>001</u>	May 14, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>A076521</u>	<u>002</u>	May 14, 2004
<u>AB</u>	MYLAN	<u>25MG;100MG</u>	<u>A075091</u>	<u>002</u>	Apr 21, 2000
<u>AB</u>		<u>50MG;200MG</u>	<u>A075091</u>	<u>001</u>	Sep 30, 1999
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG;100MG</u>	<u>A214091</u>	<u>001</u>	Oct 05, 2021
<u>AB</u>	!	<u>50MG;200MG</u>	<u>A214091</u>	<u>002</u>	Oct 05, 2021
<u>AB</u>	SUN PHARM INDS	<u>25MG;100MG</u>	<u>A077828</u>	<u>001</u>	Aug 23, 2007
<u>AB</u>		<u>50MG;200MG</u>	<u>A077828</u>	<u>002</u>	Aug 23, 2007

TABLET, ORALLY DISINTEGRATING;ORAL

CARBIDOPA AND LEVODOPA

SUN PHARM

10MG;100MG

A078690 001 Jul 31, 2009

25MG;100MG

A078690 002 Jul 31, 2009

! 25MG;250MG

A078690 003 Jul 31, 2009

CARBINOXAMINE MALEATE

SOLUTION;ORAL

CARBINOXAMINE MALEATE

! GENUS

4MG/5ML

A040458 001 Apr 25, 2003

SUSPENSION, EXTENDED RELEASE;ORAL

KARBINAL ER

+! AYTU

4MG/5ML

N022556 001 Mar 28, 2013

TABLET;ORAL

CARBINOXAMINE MALEATE

<u>AA</u>	!	GENUS	<u>4MG</u>	<u>A040442</u>	<u>001</u>	Mar 19, 2003
<u>AA</u>		INVAGEN PHARMS	<u>4MG</u>	<u>A090435</u>	<u>001</u>	Apr 15, 2010
<u>AA</u>		MISSION PHARMACAL	<u>4MG</u>	<u>A090756</u>	<u>001</u>	May 27, 2011
	!	MIKART	6MG	A207484	001	May 31, 2016

CARBOPLATIN

INJECTABLE;INTRAVENOUS

CARBOPLATIN

<u>AP</u>		ACCORD HLTHCARE	<u>50MG/5ML (10MG/ML)</u>	<u>A206775</u>	<u>001</u>	Feb 09, 2017
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A206775</u>	<u>002</u>	Feb 09, 2017
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A206775</u>	<u>003</u>	Feb 09, 2017
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A206775</u>	<u>004</u>	Feb 09, 2017
<u>AP</u>		EPIC PHARMA LLC	<u>50MG/5ML (10MG/ML)</u>	<u>A090475</u>	<u>001</u>	Jul 29, 2009
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A090475</u>	<u>002</u>	Jul 29, 2009
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A090475</u>	<u>003</u>	Jul 29, 2009
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A091268</u>	<u>002</u>	Jul 28, 2010
<u>AP</u>		EUGIA PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A205487</u>	<u>001</u>	Mar 28, 2016
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A205487</u>	<u>002</u>	Mar 28, 2016
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A205487</u>	<u>003</u>	Mar 28, 2016
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A205487</u>	<u>004</u>	Aug 03, 2020
<u>AP</u>		FRESENIUS KABI USA	<u>450MG/45ML (10MG/ML)</u>	<u>A077247</u>	<u>003</u>	Oct 21, 2004
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A077266</u>	<u>003</u>	Feb 15, 2006
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A077266</u>	<u>004</u>	Feb 15, 2006
<u>AP</u>		GLAND PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A207324</u>	<u>001</u>	Feb 15, 2017
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A207324</u>	<u>002</u>	Feb 15, 2017
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A207324</u>	<u>003</u>	Feb 15, 2017
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A207324</u>	<u>004</u>	Feb 15, 2017
<u>AP</u>		HIKMA	<u>50MG/5ML (10MG/ML)</u>	<u>A077244</u>	<u>001</u>	Oct 15, 2004
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A077244</u>	<u>002</u>	Oct 15, 2004
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A077244</u>	<u>003</u>	Oct 15, 2004
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A077244</u>	<u>004</u>	Jan 20, 2006
<u>AP</u>	!	HOSPIRA	<u>50MG/5ML (10MG/ML)</u>	<u>A076517</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>	!		<u>150MG/15ML (10MG/ML)</u>	<u>A076517</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>	!		<u>450MG/45ML (10MG/ML)</u>	<u>A076517</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>	!		<u>600MG/60ML (10MG/ML)</u>	<u>A077059</u>	<u>001</u>	Nov 23, 2004
<u>AP</u>		NOVAST LABS	<u>50MG/5ML (10MG/ML)</u>	<u>A208487</u>	<u>001</u>	Apr 26, 2017
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A208487</u>	<u>002</u>	Apr 26, 2017
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A208487</u>	<u>003</u>	Apr 26, 2017
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A208487</u>	<u>004</u>	Apr 26, 2017
<u>AP</u>		PHARMACHEMIE BV	<u>50MG/5ML (10MG/ML)</u>	<u>A077269</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A077269</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>			<u>450MG/45ML (10MG/ML)</u>	<u>A077269</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>			<u>600MG/60ML (10MG/ML)</u>	<u>A077269</u>	<u>004</u>	Dec 28, 2007
<u>AP</u>		SANDOZ	<u>50MG/5ML (10MG/ML)</u>	<u>A078280</u>	<u>001</u>	May 08, 2008
<u>AP</u>			<u>150MG/15ML (10MG/ML)</u>	<u>A078280</u>	<u>002</u>	May 08, 2008

PRESCRIPTION DRUG PRODUCT LIST

CARBOPLATIN

INJECTABLE; INTRAVENOUS

CARBOPLATIN

<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A078280 003</u>	May 08, 2008
<u>AP</u>	SUN PHARM	<u>50MG/5ML (10MG/ML)</u>	<u>A077926 001</u>	Sep 19, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077926 002</u>	Sep 19, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077926 003</u>	Sep 19, 2008
<u>AP</u>	TEYRO LABS	<u>50MG/5ML (10MG/ML)</u>	<u>A077861 001</u>	Jan 18, 2007
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077861 002</u>	Jan 18, 2007
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077861 003</u>	Jan 18, 2007
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077861 004</u>	Jan 18, 2007
	! ACCORD HLTHCARE	1GM/100ML (10MG/ML)	A206775 005	Apr 06, 2020

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

CARBOPROST TROMETHAMINE

<u>AP</u>	ALEMBIC	<u>EQ 0.25MG BASE/ML</u>	<u>A217198 001</u>	Jun 15, 2023
<u>AP</u>	AMNEAL	<u>EQ 0.25MG BASE/ML</u>	<u>A215337 001</u>	Jan 27, 2022
<u>AP</u>	CAPLIN	<u>EQ 0.25MG BASE/ML</u>	<u>A216882 001</u>	Feb 13, 2023
<u>AP</u>	DR REDDYS	<u>EQ 0.25MG BASE/ML</u>	<u>A211941 001</u>	Jul 02, 2019
<u>AP</u>	EUGIA PHARMA	<u>EQ 0.25MG BASE/ML</u>	<u>A216939 001</u>	May 25, 2023
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 0.25MG BASE/ML</u>	<u>A217657 001</u>	Aug 07, 2023
<u>AP</u>	LONG GROVE PHARMS	<u>EQ 0.25MG BASE/ML</u>	<u>A214499 001</u>	Aug 09, 2022
<u>AP</u>	SOLA PHARMS	<u>EQ 0.25MG BASE/ML</u>	<u>A216824 001</u>	May 19, 2023
<u>AP</u>	STERISCIENCE	<u>EQ 0.25MG BASE/ML</u>	<u>A216897 001</u>	Jul 25, 2023
<u>AP</u>	SUNNY	<u>EQ 0.25MG BASE/ML</u>	<u>A213118 001</u>	Mar 25, 2021

HEMABATE

<u>AP</u>	+! PFIZER	<u>EQ 0.25MG BASE/ML</u>	<u>N017989 001</u>	
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CARFILZOMIB

POWDER; INTRAVENOUS

CARFILZOMIB

<u>AP</u>	DR REDDYS	<u>60MG/VIAL</u>	<u>A209422 001</u>	Sep 09, 2019
<u>AP</u>	+! ONYX PHARMS AMGEN	<u>60MG/VIAL</u>	<u>N202714 001</u>	Jul 20, 2012
	+	10MG/VIAL	N202714 003	Jun 07, 2018
	+	30MG/VIAL	N202714 002	Jun 03, 2016

CARGLUMIC ACID

TABLET, FOR SUSPENSION; ORAL

CARBAGLU

<u>AB</u>	+! RECORDATI RARE	<u>200MG</u>	<u>N022562 001</u>	Mar 18, 2010
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CARGLUMIC ACID

<u>AB</u>	NAVINTA LLC	<u>200MG</u>	<u>A213395 001</u>	Jun 22, 2022
<u>AB</u>	NOVITIUM PHARMA	<u>200MG</u>	<u>A213729 001</u>	Oct 13, 2021

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL

VRAYLAR

	+! ABBVIE	EQ 1.5MG BASE	N204370 001	Sep 17, 2015
	+	EQ 3MG BASE	N204370 002	Sep 17, 2015
	+	EQ 4.5MG BASE	N204370 003	Sep 17, 2015
	+	EQ 6MG BASE	N204370 004	Sep 17, 2015

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

<u>AA</u>	ACCELRX LABS	<u>350MG</u>	<u>A040576 001</u>	Jun 07, 2005
<u>AA</u>	AUROBINDO PHARMA	<u>350MG</u>	<u>A040792 001</u>	Aug 06, 2009
<u>AA</u>	CHARTWELL RX	<u>350MG</u>	<u>A040245 001</u>	Sep 08, 1997
<u>AA</u>	MLV	<u>350MG</u>	<u>A211789 001</u>	Oct 20, 2021
<u>AA</u>	NATCO	<u>350MG</u>	<u>A090988 001</u>	Oct 28, 2014
<u>AA</u>	NOSTRUM LABS INC	<u>350MG</u>	<u>A207237 002</u>	Sep 21, 2020
<u>AA</u>	NOVAST LABS	<u>350MG</u>	<u>A040823 001</u>	Oct 22, 2008
<u>AA</u>	ORIENT PHARMA CO LTD	<u>350MG</u>	<u>A205085 001</u>	Oct 28, 2014
<u>AA</u>	OXFORD PHARMS	<u>350MG</u>	<u>A040188 001</u>	Mar 07, 1997
<u>AA</u>	SCIEGEN PHARMS INC	<u>350MG</u>	<u>A203374 001</u>	Jan 27, 2014
<u>AA</u>	WANBANG BIOPHARMS	<u>350MG</u>	<u>A081025 001</u>	Apr 13, 1989
<u>AA</u>	WATSON LABS	<u>350MG</u>	<u>A087499 001</u>	Apr 20, 1982
<u>AA</u>	WILSHIRE PHARMS INC	<u>350MG</u>	<u>A205126 002</u>	Jul 08, 2015

SOMA

<u>AA</u>	+ MYLAN SPECIALITY LP	<u>350MG</u>	<u>N011792 001</u>	
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CARISOPRODOL

<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>	<u>A040792 002</u>	Nov 08, 2016
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PRESCRIPTION DRUG PRODUCT LIST

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

AB	NOSTRUM LABS INC	250MG	<u>A207237</u>	<u>001</u>	May 11, 2017
AB	WILSHIRE PHARMS INC	250MG	<u>A205126</u>	<u>001</u>	Jul 08, 2015

SOMA

AB	+ ! MYLAN SPECIALITY LP	250MG	<u>N011792</u>	<u>004</u>	Sep 13, 2007
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CARMUSTINE

IMPLANT; INTRACRANIAL

GLIADEL

+ !	AZURITY	7.7MG	N020637	001	Sep 23, 1996
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INJECTABLE; INJECTION

BICNU

AP	+ ! AVET LIFESCIENCES	100MG/VIAL	<u>N017422</u>	<u>001</u>	
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CARMUSTINE

AP	ALEMBIC	100MG/VIAL	<u>A215730</u>	<u>001</u>	Oct 20, 2023
AP	AMNEAL	100MG/VIAL	<u>A211229</u>	<u>001</u>	Oct 16, 2018
AP	DR REDDYS	100MG/VIAL	<u>A213207</u>	<u>001</u>	Oct 22, 2020
AP	HENGRUI PHARMA	100MG/VIAL	<u>A211202</u>	<u>001</u>	Mar 12, 2021
AP	INGENUS PHARMS LLC	100MG/VIAL	<u>A211011</u>	<u>001</u>	Jun 21, 2022
AP	MEITHEAL	100MG/VIAL	<u>A213460</u>	<u>001</u>	Aug 02, 2021
AP	MSN	100MG/VIAL	<u>A214814</u>	<u>001</u>	May 11, 2023
AP	NAVINTA LLC	100MG/VIAL	<u>A210179</u>	<u>001</u>	Sep 11, 2018
AP	PENN LIFE	100MG/VIAL	<u>A209278</u>	<u>001</u>	Apr 02, 2019

POWDER; INTRAVENOUS

CARMUSTINE

+ !	ACCORD HLTHCARE	50MG/VIAL	N215000	001	May 16, 2022
+ !		300MG/VIAL	N215000	002	May 16, 2022

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

!	SANDOZ	1%	A075476	001	Jan 03, 2000
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CARVEDILOL

TABLET; ORAL

CARVEDILOL

AB	AUROBINDO PHARMA	3.125MG	<u>A078332</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A078332</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A078332</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A078332</u>	<u>004</u>	Sep 05, 2007
AB	BEXIMCO USA	3.125MG	<u>A078384</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A078384</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A078384</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A078384</u>	<u>004</u>	Sep 05, 2007
AB	CHARTWELL MOLECULAR	3.125MG	<u>A077474</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A077474</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A077474</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A077474</u>	<u>004</u>	Sep 05, 2007
AB	DR REDDYS LABS LTD	3.125MG	<u>A076649</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A076649</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A076649</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A076649</u>	<u>004</u>	Sep 05, 2007
AB	GLENMARK GENERICS	3.125MG	<u>A078251</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A078251</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A078251</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A078251</u>	<u>004</u>	Sep 05, 2007
AB	LUPIN	3.125MG	<u>A078217</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A078217</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A078217</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A078217</u>	<u>004</u>	Sep 05, 2007
AB	MYLAN	3.125MG	<u>A077316</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A077316</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A077316</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A077316</u>	<u>004</u>	Sep 05, 2007
AB	RUBICON	3.125MG	<u>A078165</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A078165</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A078165</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A078165</u>	<u>004</u>	Sep 05, 2007
AB	SANDOZ	3.125MG	<u>A078227</u>	<u>001</u>	Sep 05, 2007
AB		6.25MG	<u>A078227</u>	<u>002</u>	Sep 05, 2007
AB		12.5MG	<u>A078227</u>	<u>003</u>	Sep 05, 2007
AB		25MG	<u>A078227</u>	<u>004</u>	Sep 05, 2007

PRESCRIPTION DRUG PRODUCT LIST

CARVEDILOL

TABLET; ORAL

CARVEDILOL

<u>AB</u>	SUN PHARM INDS LTD	<u>3.125MG</u>	<u>A076989 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076989 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076989 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076989 004</u>	Sep 05, 2007
<u>AB</u>	TARO	<u>3.125MG</u>	<u>A077780 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077780 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077780 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077780 004</u>	Sep 05, 2007
<u>AB</u>	TEVA	<u>3.125MG</u>	<u>A076373 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076373 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076373 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076373 004</u>	Sep 05, 2007
<u>AB</u>	ZYDUS PHARMS USA INC	<u>3.125MG</u>	<u>A077614 004</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077614 001</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077614 002</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077614 003</u>	Sep 05, 2007
<u>COREG</u>				
<u>AB</u>	+ WOODWARD	<u>3.125MG</u>	<u>N020297 004</u>	May 29, 1997
<u>AB</u>	+	<u>6.25MG</u>	<u>N020297 003</u>	Sep 14, 1995
<u>AB</u>	+!	<u>12.5MG</u>	<u>N020297 002</u>	Sep 14, 1995
<u>AB</u>	+	<u>25MG</u>	<u>N020297 001</u>	Sep 14, 1995

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

CARVEDILOL PHOSPHATE

<u>AB</u>	IMPAX LABS INC	<u>10MG</u>	<u>A204717 001</u>	May 07, 2018
<u>AB</u>		<u>20MG</u>	<u>A204717 002</u>	May 07, 2018
<u>AB</u>		<u>40MG</u>	<u>A204717 003</u>	May 07, 2018
<u>AB</u>		<u>80MG</u>	<u>A204717 004</u>	May 07, 2018
<u>AB</u>	SUN PHARM INDUSTRIES	<u>10MG</u>	<u>A090132 001</u>	Oct 25, 2017
<u>AB</u>		<u>20MG</u>	<u>A090132 002</u>	Oct 25, 2017
<u>AB</u>		<u>40MG</u>	<u>A090132 003</u>	Oct 25, 2017
<u>AB</u>		<u>80MG</u>	<u>A090132 004</u>	Oct 25, 2017
<u>COREG CR</u>				
<u>AB</u>	+ WOODWARD	<u>10MG</u>	<u>N022012 001</u>	Oct 20, 2006
<u>AB</u>	+	<u>20MG</u>	<u>N022012 002</u>	Oct 20, 2006
<u>AB</u>	+!	<u>40MG</u>	<u>N022012 003</u>	Oct 20, 2006
<u>AB</u>	+	<u>80MG</u>	<u>N022012 004</u>	Oct 20, 2006

CASIMERSEN

SOLUTION; INTRAVENOUS

AMONDYS 45

+! SAREPTA THERAPS INC 100MG/2ML (50MG/ML) N213026 001 Feb 25, 2021

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CANCIDAS

<u>AP</u>	+! MERCK	<u>50MG/VIAL</u>	<u>N021227 001</u>	Jan 26, 2001
<u>AP</u>	+!	<u>70MG/VIAL</u>	<u>N021227 002</u>	Jan 26, 2001

CASPOFUNGIN ACETATE

<u>AP</u>	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>N206110 001</u>	Dec 30, 2016
<u>AP</u>		<u>70MG/VIAL</u>	<u>N206110 002</u>	Dec 30, 2016
<u>AP</u>	GLAND	<u>50MG/VIAL</u>	<u>A207092 001</u>	Sep 29, 2017
<u>AP</u>		<u>70MG/VIAL</u>	<u>A207092 002</u>	Sep 29, 2017
<u>AP</u>	HENGRUI PHARMA	<u>50MG/VIAL</u>	<u>A200833 001</u>	Jun 28, 2018
<u>AP</u>		<u>70MG/VIAL</u>	<u>A200833 002</u>	Jun 28, 2018
<u>AP</u>	UBI	<u>50MG/VIAL</u>	<u>A211263 001</u>	Oct 01, 2021
<u>AP</u>		<u>70MG/VIAL</u>	<u>A211263 002</u>	Oct 01, 2021

CEDAZURIDINE; DECITABINE

TABLET; ORAL

INQOVI

+! OTSUKA 100MG; 35MG N212576 001 Jul 07, 2020

PRESCRIPTION DRUG PRODUCT LIST

CEFACLOR

CAPSULE; ORAL

CEFACLOR

YUNG SHIN PHARM	EQ 250MG BASE	A065146 001	Jan 22, 2004
!	EQ 500MG BASE	A065146 002	Jan 22, 2004

FOR SUSPENSION; ORAL

CEFACLOR

YUNG SHIN PHARM	EQ 125MG BASE/5ML	A065412 001	Feb 17, 2012
	EQ 187MG BASE/5ML	A065412 002	Feb 17, 2012
	EQ 250MG BASE/5ML	A065412 003	Feb 17, 2012
!	EQ 375MG BASE/5ML	A065412 004	Feb 17, 2012

TABLET, EXTENDED RELEASE; ORAL

CEFACLOR

TEVA	EQ 375MG BASE	A065058 001	Sep 04, 2002
!	EQ 500MG BASE	A065058 002	Sep 04, 2002

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A065352 001</u>	Jan 25, 2007
<u>AB</u>	LUPIN	<u>EQ 500MG BASE</u>	<u>A065392 001</u>	May 29, 2007
<u>AB</u>	! TEVA PHARMS	<u>EQ 500MG BASE</u>	<u>A065282 001</u>	Jan 20, 2006

FOR SUSPENSION; ORAL

CEFADROXIL

<u>AB</u>	AUROBINDO	<u>EQ 250MG BASE/5ML</u>	<u>A065349 001</u>	Apr 25, 2013
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>A065349 002</u>	Apr 25, 2013
<u>AB</u>	LUPIN	<u>EQ 250MG BASE/5ML</u>	<u>A065396 001</u>	Feb 21, 2008
<u>AB</u>	!	<u>EQ 500MG BASE/5ML</u>	<u>A065396 002</u>	Feb 21, 2008

TABLET; ORAL

CEFADROXIL

! TEVA PHARMS	EQ 1GM BASE	A062774 001	Apr 08, 1987
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CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065303 001</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065303 002</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065306 001</u>	Oct 22, 2008
<u>AP</u>	! HIKMA FARMACEUTICA	<u>EQ 500MG BASE/VIAL</u>	<u>A065047 001</u>	Sep 18, 2001
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A065047 002</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065143 001</u>	Oct 18, 2004
<u>AP</u>	QILU	<u>EQ 1GM BASE/VIAL</u>	<u>A203661 001</u>	Dec 28, 2015
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A209217 001</u>	Oct 17, 2018
<u>AP</u>	SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A062831 001</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062831 002</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065345 001</u>	May 09, 2007
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A062831 003</u>	Sep 25, 1992

CEFAZOLIN AND DEXTROSE

+! B BRAUN	EQ 1GM BASE/VIAL	N050779 002	Jul 27, 2000
+	EQ 2GM BASE/VIAL	N050779 003	Jan 13, 2012

CEFAZOLIN SODIUM

! ACS DOBFAR	EQ 20GM BASE/VIAL	A065306 002	Aug 18, 2014
QILU	EQ 2GM BASE/VIAL	A203661 002	Mar 11, 2022
! SAMSON MEDCL	EQ 100GM BASE/VIAL	A065141 001	Nov 29, 2006
!	EQ 300GM BASE/VIAL	A065141 002	Nov 29, 2006

POWDER; INTRAVENOUS

CEFAZOLIN SODIUM

+! HIKMA	EQ 2GM BASE/VIAL	N216109 001	Oct 07, 2022
+!	EQ 3GM BASE/VIAL	N216109 002	Oct 07, 2022
+! HQ SPCLT PHARMA	EQ 2GM BASE/VIAL	N211413 001	May 08, 2023

SOLUTION; INTRAVENOUS

CEFAZOLIN IN PLASTIC CONTAINER

+! BAXTER HLTHCARE CORP	EQ 1GM BASE/50ML (EQ 20MG BASE/ML)	N207131 002	Feb 01, 2021
+!	EQ 2GM BASE/100ML (EQ 20MG BASE/ML)	N207131 001	Aug 07, 2015

CEFDINIR

CAPSULE; ORAL

CEFDINIR

<u>AB</u>	ALKEM LABS LTD	<u>300MG</u>	<u>A210220 001</u>	Feb 19, 2021
<u>AB</u>	ANDA REPOSITORY	<u>300MG</u>	<u>A065418 001</u>	Jul 18, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>300MG</u>	<u>A065434 001</u>	Jan 07, 2008
<u>AB</u>	LUPIN	<u>300MG</u>	<u>A065264 001</u>	May 19, 2006
<u>AB</u>	! SANDOZ	<u>300MG</u>	<u>A065330 001</u>	Apr 06, 2007

PRESCRIPTION DRUG PRODUCT LIST

CEFDINIR

CAPSULE; ORAL

CEFDINIR

AB	TEVA PHARMS	300MG	A065368 001	May 09, 2007
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FOR SUSPENSION; ORAL

CEFDINIR

AB	ALKEM LABS LTD	125MG/5ML	A210534 001	Feb 19, 2021
AB		250MG/5ML	A210534 002	Feb 19, 2021
AB	ANDA REPOSITORY	125MG/5ML	A065429 001	Jul 18, 2007
AB		250MG/5ML	A065429 002	Jul 18, 2007
AB	AUROBINDO PHARMA	125MG/5ML	A065473 001	Dec 14, 2007
AB	!	250MG/5ML	A065473 002	Dec 14, 2007
AB	LUPIN	125MG/5ML	A065259 001	May 31, 2006
AB		250MG/5ML	A065259 002	May 07, 2007
AB	SANDOZ	125MG/5ML	A065337 001	Apr 06, 2007
AB		250MG/5ML	A065337 002	Apr 06, 2007
AB	TEVA PHARMS	125MG/5ML	A065332 001	May 04, 2007
AB		250MG/5ML	A065332 002	May 04, 2007

CEFEPIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFEPIME HYDROCHLORIDE

AP	ACS DOBFAR	EQ 1GM BASE/VIAL	A065441 001	Mar 20, 2008
AP		EQ 2GM BASE/VIAL	A065441 002	Mar 20, 2008
AP	ASTRAL	EQ 1GM BASE/VIAL	A212721 001	Jul 21, 2020
AP		EQ 2GM BASE/VIAL	A212721 002	Jul 21, 2020
AP	! QILU	EQ 1GM BASE/VIAL	A203704 002	Feb 01, 2016
AP	!	EQ 2GM BASE/VIAL	A203704 003	Feb 01, 2016
AP	SAGENT PHARMS INC	EQ 1GM BASE/VIAL	A091048 001	Jan 04, 2017
AP		EQ 2GM BASE/VIAL	A091048 002	Jan 04, 2017

CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER

	B BRAUN	EQ 1GM BASE/VIAL	N050821 001	May 06, 2010
		EQ 2GM BASE/VIAL	N050821 002	May 06, 2010

CEFEPIME HYDROCHLORIDE

!	QILU	EQ 500MG BASE/VIAL	A203704 001	Feb 01, 2016
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CEFEPIME IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	EQ 1GM BASE/50ML (EQ 20MG BASE/ML)	N050817 001	Aug 05, 2008
+	!	EQ 2GM BASE/100ML (EQ 20MG BASE/ML)	N050817 002	Aug 05, 2008

POWDER; INTRAVENOUS

CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER

	SAMSON MEDCL	EQ 100GM BASE	A209408 001	Aug 21, 2018
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CEFIDEROCOL SULFATE TOSYLATE

POWDER; INTRAVENOUS

FETROJA

+	SHIONOGI INC	EQ 1GM BASE/VIAL	N209445 001	Nov 14, 2019
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CEFIXIME

CAPSULE; ORAL

CEFIXIME

AB	ALKEM LABS LTD	400MG	A210574 001	Oct 09, 2018
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SUPRAX

AB	! LUPIN LTD	400MG	N203195 001	Jun 01, 2012
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FOR SUSPENSION; ORAL

CEFIXIME

AB	ALKEM LABS LTD	100MG/5ML	A211775 001	Feb 19, 2021
AB		200MG/5ML	A211775 002	Feb 19, 2021
AB	AUROBINDO PHARMA LTD	100MG/5ML	A204835 001	Apr 14, 2015
AB		200MG/5ML	A204835 002	Apr 14, 2015
AB	BELCHER	100MG/5ML	A206938 001	Feb 06, 2017
AB		200MG/5ML	A206938 002	Feb 06, 2017
AB		500MG/5ML	A206939 001	Feb 06, 2017

SUPRAX

AB	! LUPIN LTD	500MG/5ML	N202091 001	Feb 20, 2013
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AB	! LUPIN PHARMS	200MG/5ML	A065355 001	Apr 10, 2007
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TABLET, CHEWABLE; ORAL

SUPRAX

	LUPIN LTD	100MG	A065380 001	Oct 25, 2010
		150MG	A065380 002	Oct 25, 2010
	!	200MG	A065380 003	Oct 25, 2010

PRESCRIPTION DRUG PRODUCT LIST

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTETAN

!	FRESENIUS KABI USA	EQ 1GM BASE/VIAL	A065374	001	Aug 09, 2007
!		EQ 2GM BASE/VIAL	A065374	002	Aug 09, 2007

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

<u>AP</u>	!	ACS DOBFAR	<u>EQ 1GM BASE/VIAL</u>	<u>A065414</u>	<u>001</u>	Jun 12, 2009
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065414</u>	<u>002</u>	Jun 12, 2009
<u>AP</u>	!		<u>EQ 10GM BASE/VIAL</u>	<u>A065415</u>	<u>001</u>	May 19, 2010
<u>AP</u>		HIKMA	<u>EQ 1GM BASE/VIAL</u>	<u>A065051</u>	<u>001</u>	Sep 11, 2000
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065051</u>	<u>002</u>	Sep 11, 2000
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065050</u>	<u>001</u>	Sep 11, 2000
<u>AP</u>		HIKMA FARMACEUTICA	<u>EQ 1GM BASE/VIAL</u>	<u>A065238</u>	<u>001</u>	Mar 12, 2010
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065238</u>	<u>002</u>	Mar 12, 2010
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065239</u>	<u>001</u>	Mar 02, 2010

CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	+	B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>N065214</u>	<u>001</u>	Mar 10, 2006
<u>AP</u>	+		<u>EQ 2GM BASE/VIAL</u>	<u>N065214</u>	<u>002</u>	Mar 10, 2006

POWDER; INTRAVENOUS

CEFOXITIN IN PLASTIC CONTAINER

		SAMSON MEDCL	EQ 100GM BASE	A200938	001	Nov 16, 2015
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CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

CEFPODOXIME PROXETIL

		AUROBINDO PHARMA LTD	EQ 50MG BASE/5ML	A065409	001	Jun 08, 2007
	!		EQ 100MG BASE/5ML	A065409	002	Jun 08, 2007

TABLET; ORAL

CEFPODOXIME PROXETIL

<u>AB</u>		ALKEM LABS LTD	<u>EQ 100MG BASE</u>	<u>A210568</u>	<u>001</u>	May 18, 2022
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A210568</u>	<u>002</u>	May 18, 2022
<u>AB</u>		ANDA REPOSITORY	<u>EQ 100MG BASE</u>	<u>A065388</u>	<u>001</u>	Nov 14, 2007
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A065388</u>	<u>002</u>	Nov 14, 2007
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 100MG BASE</u>	<u>A065370</u>	<u>001</u>	Jun 11, 2007
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A065370</u>	<u>002</u>	Jun 11, 2007
<u>AB</u>		SANDOZ	<u>EQ 100MG BASE</u>	<u>A065462</u>	<u>001</u>	May 28, 2008
<u>AB</u>	!		<u>EQ 200MG BASE</u>	<u>A065462</u>	<u>002</u>	May 28, 2008

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

<u>AB</u>		APOTEX INC	<u>125MG/5ML</u>	<u>A065351</u>	<u>001</u>	Feb 29, 2012
<u>AB</u>			<u>250MG/5ML</u>	<u>A065351</u>	<u>002</u>	Feb 29, 2012
<u>AB</u>		AUROBINDO PHARMA	<u>125MG/5ML</u>	<u>A065381</u>	<u>001</u>	Jan 30, 2007
<u>AB</u>			<u>250MG/5ML</u>	<u>A065381</u>	<u>002</u>	Jan 30, 2007
<u>AB</u>		LUPIN	<u>125MG/5ML</u>	<u>A065261</u>	<u>001</u>	Dec 19, 2005
<u>AB</u>	!		<u>250MG/5ML</u>	<u>A065261</u>	<u>002</u>	Dec 19, 2005

TABLET; ORAL

CEFPROZIL

<u>AB</u>		APOTEX INC	<u>250MG</u>	<u>A065327</u>	<u>001</u>	Mar 26, 2008
<u>AB</u>			<u>500MG</u>	<u>A065327</u>	<u>002</u>	Mar 26, 2008
<u>AB</u>		AUROBINDO PHARMA	<u>250MG</u>	<u>A065340</u>	<u>001</u>	May 24, 2007
<u>AB</u>			<u>500MG</u>	<u>A065340</u>	<u>002</u>	May 24, 2007
<u>AB</u>		CHARTWELL RX	<u>250MG</u>	<u>A065235</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>			<u>500MG</u>	<u>A065235</u>	<u>002</u>	Nov 14, 2005
<u>AB</u>		LUPIN	<u>250MG</u>	<u>A065276</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>	!		<u>500MG</u>	<u>A065276</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>		TEVA	<u>250MG</u>	<u>A065208</u>	<u>001</u>	Dec 06, 2005
<u>AB</u>			<u>500MG</u>	<u>A065208</u>	<u>002</u>	Dec 06, 2005

CEFTAROLINE FOSAMIL

POWDER; INTRAVENOUS

TEFLARO

+	!	ABBVIE	400MG/VIAL	N200327	001	Oct 29, 2010
+	!		600MG/VIAL	N200327	002	Oct 29, 2010

PRESCRIPTION DRUG PRODUCT LIST

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

<u>AP</u>	!	ACS DOBFAR	<u>1GM/VIAL</u>	<u>A062640 002</u>	Nov 20, 1985
<u>AP</u>	!		<u>2GM/VIAL</u>	<u>A062640 003</u>	Nov 20, 1985
<u>AP</u>	!		<u>6GM/VIAL</u>	<u>A062640 004</u>	Feb 03, 1992

TAZICEF

<u>AP</u>		HOSPIRA	<u>1GM/VIAL</u>	<u>A062662 002</u>	Mar 06, 1986
<u>AP</u>			<u>1GM/VIAL</u>	<u>A064032 001</u>	Oct 31, 1993
<u>AP</u>			<u>2GM/VIAL</u>	<u>A062662 003</u>	Mar 06, 1986
<u>AP</u>			<u>2GM/VIAL</u>	<u>A064032 002</u>	Oct 31, 1993
<u>AP</u>			<u>6GM/VIAL</u>	<u>A062662 004</u>	Mar 06, 1986

CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM

POWDER; INTRAVENOUS

ZERBAXA

+	!	CUBIST PHARMS LLC	EQ 1GM BASE/VIAL;EQ 0.5GM BASE/VIAL	N206829 001	Dec 19, 2014
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CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE

<u>AP</u>		ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065329 001</u>	Jul 24, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065329 002</u>	Jul 24, 2008
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065329 003</u>	Jul 24, 2008
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065328 001</u>	Jul 24, 2008
<u>AP</u>		QILU	<u>EQ 10GM BASE/VIAL</u>	<u>A209218 001</u>	Oct 17, 2018
<u>AP</u>	!	SANDOZ	<u>EQ 10GM BASE/VIAL</u>	<u>A065168 001</u>	May 17, 2005
<u>AP</u>	!	SANDOZ INC	<u>EQ 1GM BASE/VIAL</u>	<u>A065204 001</u>	May 03, 2005
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065204 002</u>	May 03, 2005
<u>AP</u>		WOCKHARDT	<u>EQ 1GM BASE/VIAL</u>	<u>A065180 001</u>	May 12, 2006

CEFTRIAZONE AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	+	!	B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>N050796 001</u>	Apr 20, 2005
<u>AP</u>	+	!		<u>EQ 2GM BASE/VIAL</u>	<u>N050796 002</u>	Apr 20, 2005

CEFTRIAZONE SODIUM

<u>AP</u>		ANDA REPOSITORY	<u>EQ 10GM BASE/VIAL</u>	<u>A091117 001</u>	Jan 20, 2017
<u>AP</u>		HIKMA	<u>EQ 10GM BASE/VIAL</u>	<u>A090701 001</u>	Oct 04, 2017

CEFTRIAZONE

		SAMSON MEDCL	EQ 100GM BASE/VIAL	A090057 001	Apr 25, 2014
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CEFTRIAZONE IN PLASTIC CONTAINER

!		BAXTER HLTHCARE	EQ 20MG BASE/ML	A065224 001	Aug 23, 2005
!			EQ 40MG BASE/ML	A065224 002	Aug 23, 2005

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAZONE

<u>AP</u>		AKORN	<u>EQ 1GM BASE/VIAL</u>	<u>A065305 003</u>	Jan 11, 2008
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065305 004</u>	Jan 11, 2008
<u>AP</u>			<u>EQ 250MG BASE/VIAL</u>	<u>A065305 001</u>	Jan 11, 2008
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065305 002</u>	Jan 11, 2008
<u>AP</u>		ASTRAL	<u>EQ 250MG BASE/VIAL</u>	<u>A091049 001</u>	Jun 11, 2018
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A091049 002</u>	Jun 11, 2018
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A091049 003</u>	Jun 11, 2018
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A091049 004</u>	Jun 11, 2018
<u>AP</u>		HIKMA FARMACEUTICA	<u>EQ 250MG BASE/VIAL</u>	<u>A065342 001</u>	Jan 10, 2008
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065342 002</u>	Jan 10, 2008
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A065342 003</u>	Jan 10, 2008
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065342 004</u>	Jan 10, 2008
<u>AP</u>		QILU	<u>EQ 250MG BASE/VIAL</u>	<u>A203702 001</u>	Jun 29, 2016
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A203702 002</u>	Jun 29, 2016
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A203702 003</u>	Jun 29, 2016
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A203702 004</u>	Jun 29, 2016
<u>AP</u>	!	SANDOZ	<u>EQ 250MG BASE/VIAL</u>	<u>A065169 001</u>	May 09, 2005
<u>AP</u>	!		<u>EQ 500MG BASE/VIAL</u>	<u>A065169 002</u>	May 09, 2005
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A065169 003</u>	May 09, 2005
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065169 004</u>	May 09, 2005
<u>AP</u>		WOCKHARDT	<u>EQ 250MG BASE/VIAL</u>	<u>A065391 001</u>	Apr 12, 2007
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A065391 002</u>	Apr 12, 2007
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065391 003</u>	Apr 12, 2007

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

<u>AB</u>		ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A065496 001</u>	Jun 07, 2010
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A065496 002</u>	Jun 07, 2010
<u>AB</u>		ANDA REPOSITORY	<u>EQ 125MG BASE</u>	<u>A065359 001</u>	Feb 15, 2008
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>A065359 002</u>	Feb 15, 2008

PRESCRIPTION DRUG PRODUCT LIST

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065359 003</u>	Feb 15, 2008
<u>AB</u>	APOTEX	<u>EQ 250MG BASE</u>	<u>A065069 001</u>	Oct 02, 2002
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065069 002</u>	Oct 02, 2002
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 125MG BASE</u>	<u>A065308 001</u>	Mar 29, 2006
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065308 002</u>	Mar 29, 2006
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065308 003</u>	Mar 29, 2006
<u>AB</u>	CHARTWELL RX	<u>EQ 125MG BASE</u>	<u>A065166 001</u>	Jul 29, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065166 002</u>	Jul 29, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065166 003</u>	Jul 29, 2005
<u>AB</u>	LUPIN	<u>EQ 250MG BASE</u>	<u>A065135 001</u>	Jul 25, 2003
<u>AB</u>	!	<u>EQ 500MG BASE</u>	<u>A065135 002</u>	Jul 25, 2003

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME SODIUM

<u>AP</u>	ACS DOBFAR SPA	<u>EQ 1.5GM BASE/VIAL</u>	<u>A064125 002</u>	May 30, 1997
<u>AP</u>	!	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065048 002</u>	Jan 09, 2004

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

<u>AB</u>	ACS DOBFAR SPA	<u>EQ 750MG BASE/VIAL</u>	<u>A064125 001</u>	May 30, 1997
<u>AB</u>	!	<u>EQ 750MG BASE/VIAL</u>	<u>A065048 001</u>	Jan 09, 2004

CELECOXIB

CAPSULE; ORAL

CELEBREX

<u>AB</u>	+	UPJOHN	<u>50MG</u>	<u>N020998 004</u>	Dec 15, 2006
<u>AB</u>	+		<u>100MG</u>	<u>N020998 001</u>	Dec 31, 1998
<u>AB</u>	+		<u>200MG</u>	<u>N020998 002</u>	Dec 31, 1998
<u>AB</u>	+	!	<u>400MG</u>	<u>N020998 003</u>	Aug 29, 2002

CELECOXIB

<u>AB</u>		ALEMBIC	<u>50MG</u>	<u>A204519 001</u>	Aug 21, 2015
<u>AB</u>			<u>100MG</u>	<u>A204519 002</u>	Aug 21, 2015
<u>AB</u>			<u>200MG</u>	<u>A204519 003</u>	Aug 21, 2015
<u>AB</u>			<u>400MG</u>	<u>A204519 004</u>	Aug 21, 2015
<u>AB</u>		APOTEX	<u>50MG</u>	<u>A204197 001</u>	Jun 02, 2015
<u>AB</u>			<u>100MG</u>	<u>A204197 002</u>	Jun 02, 2015
<u>AB</u>			<u>200MG</u>	<u>A204197 003</u>	Jun 02, 2015
<u>AB</u>		AUROBINDO PHARMA	<u>50MG</u>	<u>A206827 001</u>	Feb 01, 2016
<u>AB</u>			<u>100MG</u>	<u>A206827 002</u>	Feb 01, 2016
<u>AB</u>			<u>200MG</u>	<u>A206827 003</u>	Feb 01, 2016
<u>AB</u>			<u>400MG</u>	<u>A206827 004</u>	Feb 01, 2016
<u>AB</u>		CADILA PHARMS LTD	<u>50MG</u>	<u>A208701 001</u>	Nov 14, 2019
<u>AB</u>			<u>100MG</u>	<u>A208701 002</u>	Nov 14, 2019
<u>AB</u>			<u>200MG</u>	<u>A208701 003</u>	Nov 14, 2019
<u>AB</u>			<u>400MG</u>	<u>A208701 004</u>	Nov 14, 2019
<u>AB</u>		CIPLA	<u>50MG</u>	<u>A207446 001</u>	Sep 23, 2015
<u>AB</u>			<u>100MG</u>	<u>A207446 002</u>	Sep 23, 2015
<u>AB</u>			<u>200MG</u>	<u>A207446 003</u>	Sep 23, 2015
<u>AB</u>			<u>400MG</u>	<u>A207446 004</u>	Sep 23, 2015
<u>AB</u>		CSPC OUYI	<u>50MG</u>	<u>A210071 001</u>	Jan 23, 2018
<u>AB</u>			<u>100MG</u>	<u>A210071 002</u>	Jan 23, 2018
<u>AB</u>			<u>200MG</u>	<u>A210071 003</u>	Jan 23, 2018
<u>AB</u>		LUPIN LTD	<u>50MG</u>	<u>A202240 001</u>	Oct 29, 2014
<u>AB</u>			<u>100MG</u>	<u>A202240 002</u>	Jun 09, 2015
<u>AB</u>			<u>200MG</u>	<u>A202240 003</u>	Jun 09, 2015
<u>AB</u>			<u>400MG</u>	<u>A202240 004</u>	Jun 09, 2015
<u>AB</u>		MACLEODS PHARMS LTD	<u>50MG</u>	<u>A204590 001</u>	Mar 16, 2016
<u>AB</u>			<u>100MG</u>	<u>A204590 002</u>	Mar 16, 2016
<u>AB</u>			<u>200MG</u>	<u>A204590 003</u>	Mar 16, 2016
<u>AB</u>			<u>400MG</u>	<u>A204590 004</u>	Mar 16, 2016
<u>AB</u>		MICRO LABS	<u>50MG</u>	<u>A204776 001</u>	Apr 30, 2018
<u>AB</u>			<u>100MG</u>	<u>A204776 002</u>	Apr 30, 2018
<u>AB</u>			<u>200MG</u>	<u>A204776 003</u>	Apr 30, 2018
<u>AB</u>			<u>400MG</u>	<u>A204776 004</u>	Apr 30, 2018
<u>AB</u>		NANJING	<u>200MG</u>	<u>A213598 001</u>	May 13, 2020
<u>AB</u>		QINGDAO BAHEAL PHARM	<u>50MG</u>	<u>A208856 001</u>	Aug 07, 2019
<u>AB</u>			<u>100MG</u>	<u>A208856 002</u>	Aug 07, 2019
<u>AB</u>			<u>200MG</u>	<u>A208856 003</u>	Aug 07, 2019
<u>AB</u>			<u>400MG</u>	<u>A208856 004</u>	Aug 07, 2019
<u>AB</u>		SCIEGEN PHARMS INC	<u>50MG</u>	<u>A205129 001</u>	Dec 03, 2020

PRESCRIPTION DRUG PRODUCT LIST

CELECOXIB

CAPSULE; ORAL

CELECOXIB

<u>AB</u>		<u>100MG</u>	<u>A205129 002</u>	Dec 03, 2020
<u>AB</u>		<u>200MG</u>	<u>A205129 003</u>	Dec 03, 2020
<u>AB</u>		<u>400MG</u>	<u>A205129 004</u>	Dec 03, 2020
<u>AB</u>	TEVA	<u>50MG</u>	<u>A076898 001</u>	May 30, 2014
<u>AB</u>		<u>100MG</u>	<u>A076898 002</u>	May 30, 2014
<u>AB</u>		<u>200MG</u>	<u>A076898 003</u>	May 30, 2014
<u>AB</u>		<u>400MG</u>	<u>A076898 004</u>	May 30, 2014
<u>AB</u>	TIANJIN TIANYAO	<u>50MG</u>	<u>A207872 001</u>	Feb 25, 2020
<u>AB</u>		<u>100MG</u>	<u>A207872 002</u>	Feb 25, 2020
<u>AB</u>		<u>200MG</u>	<u>A207872 003</u>	Feb 25, 2020
<u>AB</u>		<u>400MG</u>	<u>A207872 004</u>	Feb 25, 2020
<u>AB</u>	TORRENT	<u>50MG</u>	<u>A207677 001</u>	Dec 23, 2015
<u>AB</u>		<u>100MG</u>	<u>A207677 002</u>	Dec 23, 2015
<u>AB</u>		<u>200MG</u>	<u>A207677 003</u>	Dec 23, 2015
<u>AB</u>		<u>400MG</u>	<u>A207677 004</u>	Dec 23, 2015
<u>AB</u>	UMEDICA	<u>50MG</u>	<u>A210628 001</u>	Nov 27, 2019
<u>AB</u>		<u>100MG</u>	<u>A210628 002</u>	Nov 27, 2019
<u>AB</u>		<u>200MG</u>	<u>A210628 003</u>	Nov 27, 2019
<u>AB</u>		<u>400MG</u>	<u>A210628 004</u>	Nov 27, 2019
<u>AB</u>	WATSON LABS INC	<u>50MG</u>	<u>A200562 001</u>	Feb 11, 2015
<u>AB</u>		<u>100MG</u>	<u>A200562 002</u>	Feb 11, 2015
<u>AB</u>		<u>200MG</u>	<u>A200562 003</u>	Feb 11, 2015
<u>AB</u>		<u>400MG</u>	<u>A200562 004</u>	Feb 11, 2015
<u>AB</u>	YILING	<u>50MG</u>	<u>A211412 001</u>	Mar 06, 2020
<u>AB</u>		<u>100MG</u>	<u>A211412 002</u>	Mar 06, 2020
<u>AB</u>		<u>200MG</u>	<u>A211412 003</u>	Mar 06, 2020
<u>AB</u>		<u>400MG</u>	<u>A211412 004</u>	Mar 06, 2020
BX	AMNEAL PHARMS	50MG	A208833 001	May 31, 2018
BX		100MG	A208833 002	May 31, 2018
BX		200MG	A208833 003	May 31, 2018
BX		400MG	A208833 004	May 31, 2018

SOLUTION; ORAL

ELYXYB

+! SCILEX HLDG 25MG/ML N212157 001 May 05, 2020

CELECOXIB; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

SEGLENTIS

+! KOWA PHARMS 56MG; 44MG N213426 001 Oct 15, 2021

CENOBAAMATE

TABLET; ORAL

XCOPRI

+!	SK LIFE	12.5MG	N212839 001	Mar 10, 2020
+		25MG	N212839 002	Mar 10, 2020
+		50MG	N212839 003	Mar 10, 2020
+		100MG	N212839 004	Mar 10, 2020
+		150MG	N212839 005	Mar 10, 2020
+		200MG	N212839 006	Mar 10, 2020

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

<u>AB</u>	ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A090836 001</u>	Dec 20, 2010
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A090836 002</u>	Dec 20, 2010
<u>AB</u>	!	<u>EQ 750MG BASE</u>	<u>A090836 004</u>	Mar 29, 2013
<u>AB</u>	ANDA REPOSITORY	<u>EQ 250MG BASE</u>	<u>A065248 001</u>	Jun 28, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065248 002</u>	Jun 28, 2005
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A065253 001</u>	Nov 16, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065253 002</u>	Nov 16, 2005
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A065253 003</u>	Dec 06, 2023
<u>AB</u>	BELCHER PHARMS	<u>EQ 250MG BASE</u>	<u>A062713 001</u>	Jul 15, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062713 002</u>	Jul 15, 1988
<u>AB</u>	CHARTWELL RX	<u>EQ 250MG BASE</u>	<u>A065152 001</u>	Feb 24, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065152 002</u>	Feb 24, 2005
<u>AB</u>	LUPIN	<u>EQ 250MG BASE</u>	<u>A065229 001</u>	Nov 25, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065229 002</u>	Nov 25, 2005
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A062702 001</u>	Feb 13, 1987
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062702 002</u>	Feb 13, 1987
	ALKEM LABS LTD	EQ 333MG BASE	A090836 003	Mar 29, 2013

PRESCRIPTION DRUG PRODUCT LIST

CEPHALEXIN

FOR SUSPENSION;ORAL

CEPHALEXIN

<u>AB</u>	ALKEM LABS LTD	<u>EQ 125MG BASE/5ML</u>	<u>A210221 001</u>	Mar 26, 2019
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A210221 002</u>	Mar 26, 2019
<u>AB</u>	LUPIN	<u>EQ 125MG BASE/5ML</u>	<u>A065234 001</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065234 002</u>	Aug 17, 2005
<u>AB</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>A062703 001</u>	Feb 13, 1987
<u>AB</u>	!	<u>EQ 250MG BASE/5ML</u>	<u>A062703 002</u>	Feb 13, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 125MG BASE/5ML</u>	<u>A065336 001</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065336 002</u>	Jul 25, 2007

TABLET;ORAL

CEPHALEXIN

TEVA

EQ 250MG BASE

A063023 001 Jan 12, 1989

!

EQ 500MG BASE

A063024 001 Jan 12, 1989

CERITINIB

TABLET;ORAL

ZYKADIA

+! NOVARTIS

150MG

N211225 001 Mar 18, 2019

CETIRIZINE HYDROCHLORIDE

SOLUTION;INTRAVENOUS

QUZYTIR

+! JDP

10MG/ML (10MG/ML)

N211415 001 Oct 04, 2019

SOLUTION/DROPS;OPHTHALMIC

ZERVIAE

+! HARROW EYE

EQ 0.24% BASE

N208694 001 May 30, 2017

SYRUP;ORAL

CETIRIZINE HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>5MG/5ML</u>	<u>A090766 001</u>	Oct 07, 2009
<u>AA</u>	BAJAJ	<u>5MG/5ML</u>	<u>A090191 001</u>	Nov 12, 2009
<u>AA</u>	BIONPHARMA	<u>5MG/5ML</u>	<u>A078488 001</u>	Oct 06, 2008
<u>AA</u>	CHARTWELL MOLECULAR	<u>5MG/5ML</u>	<u>A078876 001</u>	May 11, 2012
<u>AA</u>	! PADAGIS US	<u>5MG/5ML</u>	<u>A078398 001</u>	Jun 17, 2008
<u>AA</u>	TARO	<u>5MG/5ML</u>	<u>A076601 001</u>	Jun 20, 2008
<u>AA</u>	TEVA PHARMS	<u>5MG/5ML</u>	<u>A077279 001</u>	May 27, 2008

CETRORELIX ACETATE

POWDER;SUBCUTANEOUS

CETRORELIX ACETATE

<u>AP</u>	TEVA PHARMS INC	<u>EQ 0.25MG BASE/VIAL</u>	<u>A215737 001</u>	Aug 12, 2022
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CETROTIDE

<u>AP</u>	+! EMD SERONO INC	<u>EQ 0.25MG BASE/VIAL</u>	<u>N021197 001</u>	Aug 11, 2000
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CEVIMELINE HYDROCHLORIDE

CAPSULE;ORAL

CEVIMELINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA	<u>30MG</u>	<u>A215056 001</u>	Apr 18, 2023
<u>AB</u>	BIONPHARMA	<u>30MG</u>	<u>A218290 001</u>	Nov 08, 2023
<u>AB</u>	HIKMA	<u>30MG</u>	<u>A091591 001</u>	Jul 08, 2013
<u>AB</u>	NOVEL LABS INC	<u>30MG</u>	<u>A204746 001</u>	Dec 30, 2016
<u>AB</u>	RISING	<u>30MG</u>	<u>A203775 001</u>	Jun 04, 2014
<u>AB</u>	RUBICON	<u>30MG</u>	<u>A216682 001</u>	Apr 06, 2023

EVOXAC

<u>AB</u>	+! COSETTE	<u>30MG</u>	<u>N020989 002</u>	Jan 11, 2000
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CHENODIOL

TABLET;ORAL

CHENODIOL

! LGM PHARMA

250MG

A091019 001 Oct 22, 2009

CHLORAMBUCIL

TABLET;ORAL

LEUKERAN

+! WAYLIS THERAP

2MG

N010669 002

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE;INJECTION

CHLORAMPHENICOL SODIUM SUCCINATE

! FRESENIUS KABI USA EQ 1GM BASE/VIAL

A062365 001 Aug 25, 1982

PRESCRIPTION DRUG PRODUCT LIST

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

<u>AB</u>	BARR	<u>5MG</u>	<u>A084768</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>A083116</u>	<u>001</u>	
<u>AB</u>	!	<u>25MG</u>	<u>A084769</u>	<u>001</u>	
<u>AB</u>	CHARTWELL RX	<u>5MG</u>	<u>A084678</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>A084041</u>	<u>001</u>	
<u>AB</u>		<u>25MG</u>	<u>A084679</u>	<u>002</u>	
<u>LIBRIUM</u>					
<u>AB</u>	+	BAUSCH	<u>25MG</u>	<u>A085475</u>	<u>001</u>
<u>AB</u>	+	VALEANT PHARM INTL	<u>5MG</u>	<u>A085461</u>	<u>001</u>
<u>AB</u>	+		<u>10MG</u>	<u>A085472</u>	<u>001</u>

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE

<u>AB</u>	ALEMBIC	<u>5MG;2.5MG</u>	<u>A216969</u>	<u>001</u>	Sep 15, 2023
<u>AB</u>	ALKEM LABS LTD	<u>5MG;2.5MG</u>	<u>A214065</u>	<u>001</u>	Apr 26, 2021
<u>AB</u>	AMNEAL	<u>5MG;2.5MG</u>	<u>A215555</u>	<u>001</u>	Oct 25, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG;2.5MG</u>	<u>A216419</u>	<u>001</u>	Sep 14, 2023
<u>AB</u>	CHARTWELL RX	<u>5MG;2.5MG</u>	<u>A213530</u>	<u>001</u>	Oct 20, 2020
<u>AB</u>	COREPHARMA	<u>5MG;2.5MG</u>	<u>A215453</u>	<u>001</u>	Jul 29, 2022
<u>AB</u>	DR REDDYS	<u>5MG;2.5MG</u>	<u>A214698</u>	<u>001</u>	May 10, 2021
<u>AB</u>	MICRO LABS	<u>5MG;2.5MG</u>	<u>A215835</u>	<u>001</u>	Jul 19, 2022
<u>AB</u>	MISEMER	<u>5MG;2.5MG</u>	<u>A210579</u>	<u>001</u>	Jul 29, 2020
<u>AB</u>	NUVO PHARMS INC	<u>5MG;2.5MG</u>	<u>A211421</u>	<u>001</u>	Jul 07, 2020
<u>LIBRAX</u>					
<u>AB</u>	+	BAUSCH	<u>5MG;2.5MG</u>	<u>N012750</u>	<u>001</u>

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

<u>AT</u>	BAJAJ	<u>0.12%</u>	<u>A075561</u>	<u>001</u>	Nov 14, 2000
<u>AT</u>	CHARTWELL RX	<u>0.12%</u>	<u>A075006</u>	<u>001</u>	Mar 03, 2004
<u>AT</u>	LYNE	<u>0.12%</u>	<u>A074291</u>	<u>001</u>	Dec 28, 1995
<u>AT</u>	PHARM ASSOC	<u>0.12%</u>	<u>A074522</u>	<u>001</u>	Dec 15, 1995
<u>AT</u>	XTTRIUM	<u>0.12%</u>	<u>A077789</u>	<u>001</u>	Jun 18, 2009
<u>PERIDEX</u>					
<u>AT</u>	+	3M	<u>0.12%</u>	<u>N019028</u>	<u>001</u> Aug 13, 1986
<u>PERIOGARD</u>					
<u>AT</u>	COLGATE-PALMOLIVE CO	<u>0.12%</u>	<u>A203212</u>	<u>001</u>	Jan 28, 2016

CHLOROPROCAINE HYDROCHLORIDE

GEL; OPHTHALMIC

IHEEZO

+! HARROW EYE 3%

N216227 001 Sep 27, 2022

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

<u>AP</u>	HIKMA	<u>2%</u>	<u>A040273</u>	<u>001</u>	Sep 09, 1998
<u>AP</u>		<u>3%</u>	<u>A040273</u>	<u>002</u>	Sep 09, 1998
<u>NESACAINE</u>					
<u>AP</u>	+	FRESENIUS KABI USA	<u>2%</u>	<u>N009435</u>	<u>002</u>
<u>NESACAINE-MPF</u>					
<u>AP</u>	+	FRESENIUS KABI USA	<u>2%</u>	<u>N009435</u>	<u>006</u> May 02, 1996
<u>AP</u>	+		<u>3%</u>	<u>N009435</u>	<u>007</u> May 02, 1996
<u>NESACAINE</u>					
	+	FRESENIUS KABI USA	1%	N009435	001

CHLOROQUINE PHOSPHATE

TABLET; ORAL

CHLOROQUINE PHOSPHATE

<u>AA</u>	IMPAX LABS	<u>250MG</u>	<u>A080880</u>	<u>001</u>	
<u>AA</u>		<u>500MG</u>	<u>A040516</u>	<u>001</u>	Aug 29, 2003
<u>AA</u>	IPCA LABS LTD	<u>250MG</u>	<u>A090610</u>	<u>001</u>	Dec 03, 2009
<u>AA</u>		<u>500MG</u>	<u>A090249</u>	<u>001</u>	Dec 03, 2009
<u>AA</u>	!	NATCO PHARMA LTD	<u>250MG</u>	<u>A091621</u>	<u>001</u> Jan 21, 2011
<u>AA</u>	!		<u>500MG</u>	<u>A090612</u>	<u>001</u> Jan 21, 2011
<u>AA</u>	SUVEN PHARMS	<u>500MG</u>	<u>A214756</u>	<u>001</u>	Sep 03, 2021

PRESCRIPTION DRUG PRODUCT LIST

CHLOROTHIAZIDE

SUSPENSION; ORAL

DIURIL

+! SALIX PHARMS 250MG/5ML N011870 001

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

CHLOROTHIAZIDE SODIUM

AP	AM REGENT	<u>EQ 500MG BASE/VIAL</u>	<u>A202561 001</u>	Apr 22, 2013
AP	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A090896 001</u>	Oct 16, 2009
AP	RK PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A202493 001</u>	Jun 18, 2014
AP	! SAGENT PHARMS INC	<u>EQ 500MG BASE/VIAL</u>	<u>A202462 001</u>	May 29, 2015
AP	SUN PHARM	<u>EQ 500MG BASE/VIAL</u>	<u>A091546 001</u>	Jul 26, 2011

CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE

TABLET, EXTENDED RELEASE; ORAL

TUXARIN ER

MAINPOINTE 8MG; 54.3MG N206323 001 Jun 22, 2015

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

! PADAGIS US 4MG/5ML; 5MG/5ML; 60MG/5ML A204627 001 Apr 29, 2014

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

! TRIS PHARMA INC EQ 8MG MALEATE/5ML; EQ 10MG BITARTRATE/5ML A091632 001 Oct 01, 2010

CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

! GENUS 30MG/ML A214542 001 Jun 02, 2021
! 100MG/ML A214542 002 Jun 02, 2021

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

AP	EUGIA PHARMA	<u>25MG/ML</u>	<u>A211816 001</u>	Jul 07, 2020
AP	+! WEST-WARD PHARMS	<u>25MG/ML</u>	<u>A083329 001</u>	

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

AB	ALEMBIC	<u>10MG</u>	<u>A217350 001</u>	Jul 18, 2023
AB		<u>25MG</u>	<u>A217350 002</u>	Jul 18, 2023
AB		<u>50MG</u>	<u>A217350 003</u>	Jul 18, 2023
AB		<u>100MG</u>	<u>A217350 004</u>	Jul 18, 2023
AB		<u>200MG</u>	<u>A217350 005</u>	Jul 18, 2023
AB	AMNEAL PHARMS CO	<u>10MG</u>	<u>A209755 001</u>	Sep 10, 2018
AB		<u>25MG</u>	<u>A209755 002</u>	Sep 10, 2018
AB		<u>50MG</u>	<u>A209755 003</u>	Sep 10, 2018
AB		<u>100MG</u>	<u>A209755 004</u>	Sep 10, 2018
AB		<u>200MG</u>	<u>A209755 005</u>	Sep 10, 2018
AB	APPCO	<u>10MG</u>	<u>A213590 001</u>	Aug 31, 2020
AB		<u>25MG</u>	<u>A213590 002</u>	Aug 31, 2020
AB		<u>50MG</u>	<u>A213590 003</u>	Aug 31, 2020
AB		<u>100MG</u>	<u>A213590 004</u>	Aug 31, 2020
AB		<u>200MG</u>	<u>A213590 005</u>	Aug 31, 2020
AB	CHARTWELL RX	<u>10MG</u>	<u>A212630 001</u>	Nov 29, 2021
AB		<u>25MG</u>	<u>A212630 002</u>	Nov 29, 2021
AB		<u>50MG</u>	<u>A212630 003</u>	Nov 29, 2021
AB		<u>100MG</u>	<u>A212630 004</u>	Nov 29, 2021
AB		<u>200MG</u>	<u>A212630 005</u>	Nov 29, 2021
AB	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A212144 001</u>	Mar 23, 2021
AB		<u>25MG</u>	<u>A212144 002</u>	Mar 23, 2021
AB		<u>50MG</u>	<u>A212144 003</u>	Mar 23, 2021
AB		<u>100MG</u>	<u>A212144 004</u>	Mar 23, 2021
AB		<u>200MG</u>	<u>A212144 005</u>	Mar 23, 2021
AB	LANNETT CO INC	<u>10MG</u>	<u>A212996 001</u>	Jan 22, 2021
AB		<u>25MG</u>	<u>A212996 002</u>	Jan 22, 2021
AB		<u>50MG</u>	<u>A212996 003</u>	Jan 22, 2021
AB		<u>100MG</u>	<u>A212996 004</u>	Jan 22, 2021
AB		<u>200MG</u>	<u>A212996 005</u>	Jan 22, 2021
AB	LUPIN	<u>10MG</u>	<u>A213327 001</u>	Jul 13, 2023
AB		<u>25MG</u>	<u>A213327 002</u>	Jul 13, 2023
AB		<u>50MG</u>	<u>A213327 003</u>	Jul 13, 2023

PRESCRIPTION DRUG PRODUCT LIST

CHLORPROMAZINE HYDROCHLORIDE

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

<u>AB</u>		<u>100MG</u>	<u>A213327 004</u>	Jul 13, 2023
<u>AB</u>		<u>200MG</u>	<u>A213327 005</u>	Jul 13, 2023
<u>AB</u>	MSN	<u>25MG</u>	<u>A214827 001</u>	Jan 27, 2022
<u>AB</u>		<u>50MG</u>	<u>A214827 002</u>	Jan 27, 2022
<u>AB</u>		<u>100MG</u>	<u>A214827 003</u>	Jan 27, 2022
<u>AB</u>		<u>200MG</u>	<u>A214827 004</u>	Jan 27, 2022
<u>AB</u>	SUN PHARM	<u>10MG</u>	<u>A214256 001</u>	Oct 26, 2020
<u>AB</u>		<u>25MG</u>	<u>A214256 002</u>	Oct 26, 2020
<u>AB</u>		<u>50MG</u>	<u>A214256 003</u>	Oct 26, 2020
<u>AB</u>		<u>100MG</u>	<u>A214256 004</u>	Oct 26, 2020
<u>AB</u>		<u>200MG</u>	<u>A214256 005</u>	Oct 26, 2020
<u>AB</u>	TEVA PHARMS	<u>10MG</u>	<u>A215659 001</u>	Oct 25, 2021
<u>AB</u>		<u>25MG</u>	<u>A215659 002</u>	Oct 25, 2021
<u>AB</u>		<u>50MG</u>	<u>A215659 003</u>	Oct 25, 2021
<u>AB</u>		<u>100MG</u>	<u>A215659 004</u>	Oct 25, 2021
<u>AB</u>		<u>200MG</u>	<u>A215659 005</u>	Oct 25, 2021
<u>AB</u>	+ UPSHER SMITH LABS	<u>10MG</u>	<u>A083386 001</u>	
<u>AB</u>	+!	<u>25MG</u>	<u>A084112 001</u>	
<u>AB</u>	+	<u>50MG</u>	<u>A084113 001</u>	
<u>AB</u>	+!	<u>100MG</u>	<u>A084114 001</u>	
<u>AB</u>	+	<u>200MG</u>	<u>A084115 001</u>	
<u>AB</u>	ZYDUS	<u>10MG</u>	<u>A213368 001</u>	Jan 17, 2020
<u>AB</u>		<u>25MG</u>	<u>A213368 002</u>	Jan 17, 2020
<u>AB</u>		<u>50MG</u>	<u>A213368 003</u>	Jan 17, 2020
<u>AB</u>		<u>100MG</u>	<u>A213368 004</u>	Jan 17, 2020
<u>AB</u>		<u>200MG</u>	<u>A213368 005</u>	Jan 17, 2020

CHLORTHALIDONE

TABLET;ORAL

CHLORTHALIDONE

<u>AB</u>	AJANTA PHARMA LTD	<u>25MG</u>	<u>A214129 001</u>	Nov 27, 2020
<u>AB</u>		<u>50MG</u>	<u>A214129 002</u>	Nov 27, 2020
<u>AB</u>	ALEMBIC	<u>25MG</u>	<u>A216262 001</u>	Aug 26, 2022
<u>AB</u>		<u>50MG</u>	<u>A216262 002</u>	Aug 26, 2022
<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A213412 001</u>	Feb 11, 2020
<u>AB</u>		<u>50MG</u>	<u>A213412 002</u>	Feb 11, 2020
<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A207204 001</u>	Jul 01, 2019
<u>AB</u>		<u>50MG</u>	<u>A207204 002</u>	Jul 01, 2019
<u>AB</u>	APPCO	<u>25MG</u>	<u>A210742 001</u>	Oct 12, 2018
<u>AB</u>		<u>50MG</u>	<u>A210742 002</u>	Oct 12, 2018
<u>AB</u>	CHARTWELL RX	<u>25MG</u>	<u>A211063 001</u>	Feb 26, 2019
<u>AB</u>		<u>50MG</u>	<u>A211063 002</u>	Feb 26, 2019
<u>AB</u>	INVENTIA	<u>25MG</u>	<u>A211320 001</u>	Feb 09, 2022
<u>AB</u>		<u>50MG</u>	<u>A211320 002</u>	Feb 09, 2022
<u>AB</u>	MANKIND PHARMA	<u>25MG</u>	<u>A215587 001</u>	Aug 04, 2022
<u>AB</u>		<u>50MG</u>	<u>A215587 002</u>	Aug 04, 2022
<u>AB</u>	+ MYLAN	<u>25MG</u>	<u>A086831 002</u>	
<u>AB</u>	+!	<u>50MG</u>	<u>A086831 001</u>	
<u>AB</u>	NOVAST LABS	<u>25MG</u>	<u>A206904 001</u>	Mar 30, 2017
<u>AB</u>		<u>50MG</u>	<u>A206904 002</u>	Mar 30, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A089286 002</u>	Jul 21, 1986
<u>AB</u>		<u>50MG</u>	<u>A089286 001</u>	Jul 21, 1986
<u>AB</u>	TAGI	<u>25MG</u>	<u>A212878 001</u>	Feb 24, 2022
<u>AB</u>		<u>50MG</u>	<u>A212875 001</u>	Feb 24, 2022
<u>AB</u>	UMEDICA	<u>25MG</u>	<u>A207222 001</u>	May 24, 2018
<u>AB</u>		<u>50MG</u>	<u>A207222 002</u>	May 24, 2018
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A211627 001</u>	Aug 06, 2019
<u>AB</u>		<u>50MG</u>	<u>A211627 002</u>	Aug 06, 2019
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A207813 001</u>	May 10, 2019
<u>AB</u>		<u>50MG</u>	<u>A207813 002</u>	May 10, 2019
<u>THALITONE</u>				
BX	+! CASPER PHARMA LLC	25MG	N019574 002	Feb 12, 1992
	+	15MG	N019574 001	Dec 20, 1988

PRESCRIPTION DRUG PRODUCT LIST

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A089853 001</u>	May 04, 1988
<u>AA</u>	BELCHER	<u>250MG</u>	<u>A215540 001</u>	Jan 24, 2023
<u>AA</u>	COREPHARMA	<u>250MG</u>	<u>A214702 001</u>	Apr 10, 2023
<u>AA</u>		<u>500MG</u>	<u>A214941 002</u>	Oct 20, 2022
<u>AA</u>	DR REDDYS LABS SA	<u>500MG</u>	<u>A211849 002</u>	Jul 21, 2020
<u>AA</u>	GRAVITI PHARMS	<u>250MG</u>	<u>A216925 001</u>	Aug 09, 2023
<u>AA</u>	! MIKART	<u>250MG</u>	<u>A207483 001</u>	Jun 24, 2016
<u>AA</u>	NOVITIUM PHARMA	<u>500MG</u>	<u>A212254 001</u>	Sep 12, 2019
<u>AA</u>	RISE PHARMA	<u>250MG</u>	<u>A215158 001</u>	Jul 29, 2021
<u>AA</u>	! WATSON LABS	<u>500MG</u>	<u>A089859 001</u>	May 04, 1988
<u>AB</u>	APTAPHARMA INC	<u>375MG</u>	<u>A212053 001</u>	Sep 14, 2020
<u>AB</u>		<u>750MG</u>	<u>A212053 002</u>	Sep 14, 2020
<u>AB</u>	AUROBINDO PHARMA LTD	<u>375MG</u>	<u>A089853 002</u>	Aug 17, 2023
<u>AB</u>		<u>750MG</u>	<u>A089853 003</u>	Aug 17, 2023
<u>AB</u>	COREPHARMA	<u>375MG</u>	<u>A214941 001</u>	Oct 20, 2022
<u>AB</u>		<u>750MG</u>	<u>A214941 003</u>	Oct 20, 2022
<u>AB</u>	DR REDDYS LABS SA	<u>375MG</u>	<u>A211849 001</u>	Jul 21, 2020
<u>AB</u>		<u>750MG</u>	<u>A211849 003</u>	Jul 21, 2020
<u>AB</u>	! MIKART	<u>375MG</u>	<u>A040861 001</u>	Jun 01, 2010
<u>AB</u>	!	<u>750MG</u>	<u>A040861 002</u>	Jun 01, 2010
<u>AB</u>	NOVITIUM PHARMA	<u>375MG</u>	<u>A212253 001</u>	Nov 27, 2019
<u>AB</u>		<u>750MG</u>	<u>A212253 002</u>	Nov 27, 2019
<u>AB</u>	PAR PHARM INC	<u>375MG</u>	<u>A212743 002</u>	Apr 29, 2021
<u>AB</u>		<u>750MG</u>	<u>A212743 001</u>	Nov 02, 2020
<u>AB</u>	RISING	<u>375MG</u>	<u>A213126 001</u>	Apr 05, 2022
<u>AB</u>		<u>500MG</u>	<u>A213126 002</u>	Apr 05, 2022
<u>AB</u>		<u>750MG</u>	<u>A213126 003</u>	Apr 05, 2022
<u>AB</u>	TEVA PHARMS USA INC	<u>375MG</u>	<u>A212898 001</u>	Jun 17, 2020
<u>AB</u>		<u>750MG</u>	<u>A212898 002</u>	Jun 17, 2020
<u>AB</u>	UPSHER SMITH LABS	<u>375MG</u>	<u>A212047 001</u>	Jan 27, 2023
<u>AB</u>		<u>750MG</u>	<u>A212047 002</u>	Jan 27, 2023

CHOLESTYRAMINE

POWDER; ORAL

CHOLESTYRAMINE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211119 001</u>	Apr 06, 2020
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211119 002</u>	Apr 06, 2020
<u>AB</u>	ALKEM LABS LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211856 001</u>	Oct 19, 2021
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211856 002</u>	Oct 19, 2021
<u>AB</u>	! EPIC PHARMA LLC	<u>EQ 4GM RESIN/PACKET</u>	<u>A074557 001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074557 002</u>	Aug 15, 1996
<u>AB</u>	PAR PHARM	<u>EQ 4GM RESIN/PACKET</u>	<u>A077204 001</u>	Aug 26, 2005
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077204 002</u>	Aug 26, 2005
<u>AB</u>	TAGI	<u>EQ 4GM RESIN/PACKET</u>	<u>A209597 001</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A209597 002</u>	Mar 09, 2021
<u>AB</u>	ZYDUS PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A202901 001</u>	Jul 02, 2018

CHOLESTYRAMINE LIGHT

<u>AB</u>	ALKEM LABS LTD	<u>EQ 4GM RESIN/PACKET</u>	<u>A211799 001</u>	Oct 19, 2021
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A211799 002</u>	Oct 19, 2021
<u>AB</u>	! EPIC PHARMA LLC	<u>EQ 4GM RESIN/PACKET</u>	<u>A074558 001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074558 002</u>	Aug 15, 1996
<u>AB</u>	PAR PHARM	<u>EQ 4GM RESIN/PACKET</u>	<u>A077203 001</u>	Aug 26, 2005
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077203 002</u>	Aug 26, 2005
<u>AB</u>	TAGI	<u>EQ 4GM RESIN/PACKET</u>	<u>A209599 001</u>	Nov 12, 2020
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A209599 002</u>	Nov 12, 2020
<u>AB</u>	ZYDUS PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A202902 001</u>	Apr 25, 2017

LOCHOLEST

<u>AB</u>	CHARTWELL RX	<u>EQ 4GM RESIN/PACKET</u>	<u>A074561 001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074561 002</u>	Aug 15, 1996

LOCHOLEST LIGHT

<u>AB</u>	CHARTWELL RX	<u>EQ 4GM RESIN/PACKET</u>	<u>A074562 001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074562 002</u>	Aug 15, 1996

PREVALITE

<u>AB</u>	UPSHER SMITH LABS	<u>EQ 4GM RESIN/PACKET</u>	<u>A073263 001</u>	Feb 22, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A073263 002</u>	Oct 30, 1997

PRESCRIPTION DRUG PRODUCT LIST

CHOLIC ACID

CAPSULE; ORAL

CHOLBAM

+ MIRUM

50MG

N205750 001 Mar 17, 2015

+!

250MG

N205750 002 Mar 17, 2015

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11**AP** DECATUR**4-33.1mCi/ML****A206319 001** Nov 13, 2015**AP** +! MCPRF**4-33.1mCi/ML****N203155 001** Sep 12, 2012CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID**AB** ACTAVIS ELIZABETH**EQ 45MG FENOFIBRIC ACID****A200920 001** Oct 07, 2015**AB** **EQ 135MG FENOFIBRIC ACID****A200920 002** Oct 07, 2015**AB** ALEMBIC **EQ 45MG FENOFIBRIC ACID****A208705 001** May 12, 2017**AB** **EQ 135MG FENOFIBRIC ACID****A208705 002** May 12, 2017**AB** ANCHEN PHARMS **EQ 45MG FENOFIBRIC ACID****A201573 002** Jul 18, 2013**AB** **EQ 135MG FENOFIBRIC ACID****A201573 001** Jul 18, 2013**AB** AUROBINDO PHARMA**EQ 45MG FENOFIBRIC ACID****A212598 001** Jul 25, 2019

LTD

AB **EQ 135MG FENOFIBRIC ACID****A212598 002** Jul 25, 2019**AB** GRAVITI PHARMS **EQ 45MG FENOFIBRIC ACID****A211626 001** Jul 18, 2019**AB** **EQ 135MG FENOFIBRIC ACID****A211626 002** Jul 18, 2019**AB** IMPAX LABS INC **EQ 45MG FENOFIBRIC ACID****A200264 001** Sep 07, 2016**AB** **EQ 135MG FENOFIBRIC ACID****A200264 002** Sep 07, 2016**AB** LUPIN LTD **EQ 45MG FENOFIBRIC ACID****A200750 001** Dec 04, 2013**AB** **EQ 135MG FENOFIBRIC ACID****A200750 002** Dec 04, 2013**AB** MICRO LABS **EQ 45MG FENOFIBRIC ACID****A213450 001** Jun 16, 2020**AB** **EQ 135MG FENOFIBRIC ACID****A213450 002** Jun 16, 2020**AB** SCINOPHARM TAIWAN **EQ 45MG FENOFIBRIC ACID****A210469 001** Jul 05, 2019**AB** **EQ 135MG FENOFIBRIC ACID****A210469 002** Jul 05, 2019**AB** YICHANG HUMANWELL **EQ 45MG FENOFIBRIC ACID****A212562 001** Dec 23, 2020**AB** **EQ 135MG FENOFIBRIC ACID****A212562 002** Dec 23, 2020TRILIPIX**AB** + ABBVIE**EQ 45MG FENOFIBRIC ACID****N022224 001** Dec 15, 2008**AB** +!**EQ 135MG FENOFIBRIC ACID****N022224 002** Dec 15, 2008CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA

EQ 0.004MG CHROMIUM/ML

N018961 001 Jun 26, 1986

CICLESONIDE

AEROSOL, METERED; INHALATION

ALVESCO

+! COVIS

0.08MG/INH

N021658 002 Jan 10, 2008

+!

0.16MG/INH

N021658 003 Jan 10, 2008

AEROSOL, METERED; NASAL

ZETONNA

+! COVIS

0.037MG/INH

N202129 001 Jan 20, 2012

SPRAY, METERED; NASAL

OMNARIS

+! COVIS

0.05MG/SPRAY

N022004 001 Oct 20, 2006

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX**AB** COSETTE**0.77%****A078463 001** Dec 20, 2010**AB** FOUGERA PHARMS**0.77%****A076435 001** Dec 29, 2004**AB** GLENMARK PHARMS**0.77%****A090273 001** Nov 10, 2009**AB** PADAGIS ISRAEL**0.77%****A077364 001** Mar 03, 2006**AB** TARO**0.77%****A076790 001** Apr 12, 2005LOPROX**AB** +! MEDIMETRIKS PHARMS**0.77%****N018748 001** Dec 30, 1982

GEL; TOPICAL

CICLOPIROX**AB** FOUGERA PHARMS**0.77%****A077896 001** Jun 10, 2008**AB** ! GLENMARK GENERICS**0.77%****A091595 001** Feb 29, 2012**AB** PADAGIS US**0.77%****A078266 001** Jan 07, 2009

SHAMPOO; TOPICAL

CICLOPIROX**AT** ACTAVIS MID**1%****A090490 001** Nov 24, 2009

PRESCRIPTION DRUG PRODUCT LIST

CICLOPIROX

SHAMPOO; TOPICAL

CICLOPIROX

ATLANTIC

<u>AT</u>	FOUGERA PHARMS	<u>1%</u>	<u>A090146</u>	<u>001</u>	May 25, 2010
<u>AT</u>	PADAGIS US	<u>1%</u>	<u>A078594</u>	<u>001</u>	Feb 16, 2010
<u>AT</u>	TARO	<u>1%</u>	<u>A090269</u>	<u>001</u>	Feb 23, 2011

LOPROX

<u>AT</u>	+! BAUSCH	<u>1%</u>	<u>N021159</u>	<u>001</u>	Feb 28, 2003
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SOLUTION; TOPICAL

CICLOPIROX

<u>AT</u>	ACELLA	<u>8%</u>	<u>A078172</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	CHARTWELL RX	<u>8%</u>	<u>A078046</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	COSETTE	<u>8%</u>	<u>A078233</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	ENCUBE	<u>8%</u>	<u>A077687</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	HIKMA	<u>8%</u>	<u>A078270</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	! PADAGIS US	<u>8%</u>	<u>A077623</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	RISING	<u>8%</u>	<u>A078124</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	TARO PHARM INDS	<u>8%</u>	<u>A078144</u>	<u>001</u>	Sep 18, 2007

SUSPENSION; TOPICAL

CICLOPIROX

<u>AB</u>	FOUGERA PHARMS	<u>0.77%</u>	<u>A076422</u>	<u>001</u>	Aug 06, 2004
<u>AB</u>	PADAGIS ISRAEL	<u>0.77%</u>	<u>A077676</u>	<u>001</u>	Dec 15, 2006
<u>AB</u>	TARO	<u>0.77%</u>	<u>A077092</u>	<u>001</u>	Aug 10, 2005

LOPROX

<u>AB</u>	+! MEDIMETRIKS PHARMS	<u>0.77%</u>	<u>N019824</u>	<u>001</u>	Dec 30, 1988
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CIDOFIVIR

INJECTABLE; INJECTION

CIDOFIVIR

<u>AP</u>	AVET LIFESCIENCES	<u>EQ 75MG BASE/ML</u>	<u>A202501</u>	<u>001</u>	Jul 26, 2012
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>EQ 75MG BASE/ML</u>	<u>A201276</u>	<u>001</u>	Jun 27, 2012

CILASTATIN SODIUM; IMIPENEM

POWDER; INTRAVENOUS

IMIPENEM AND CILASTATIN

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	<u>A090577</u>	<u>002</u>	Dec 21, 2011
<u>AP</u>	HQ SPCLT PHARMA	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	<u>A207594</u>	<u>001</u>	Dec 12, 2019

PRIMAXIN

<u>AP</u>	+! MERCK	<u>EQ 500MG BASE/VIAL; 500MG/VIAL</u>	<u>N050587</u>	<u>002</u>	Nov 26, 1985
	IMIPENEM AND CILASTATIN				
	! ACS DOBFAR	<u>EQ 250MG BASE/VIAL; 250MG/VIAL</u>	<u>A090577</u>	<u>001</u>	Dec 21, 2011

CILASTATIN SODIUM; IMIPENEM; RELEBACTAM

POWDER; INTRAVENOUS

RECARBRIO

	+! MSD MERCK CO	<u>EQ 500MG</u> <u>BASE/VIAL; 500MG/VIAL; 250MG/VIAL</u>	<u>N212819</u>	<u>001</u>	Jul 16, 2019
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CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

<u>AB</u>	APOTEX INC	<u>50MG</u>	<u>A077030</u>	<u>001</u>	Dec 10, 2004
<u>AB</u>		<u>100MG</u>	<u>A077030</u>	<u>002</u>	Dec 10, 2004
<u>AB</u>	CHARTWELL RX	<u>50MG</u>	<u>A077831</u>	<u>002</u>	Sep 24, 2012
<u>AB</u>		<u>100MG</u>	<u>A077831</u>	<u>001</u>	Sep 24, 2012
<u>AB</u>	EPIC PHARMA LLC	<u>50MG</u>	<u>A077022</u>	<u>002</u>	Mar 11, 2005
<u>AB</u>		<u>100MG</u>	<u>A077022</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>	SLATE RUN PHARMA	<u>50MG</u>	<u>A077208</u>	<u>002</u>	Mar 29, 2006
<u>AB</u>		<u>100MG</u>	<u>A077208</u>	<u>001</u>	Mar 29, 2006
<u>AB</u>	! TEVA	<u>50MG</u>	<u>A077027</u>	<u>001</u>	Nov 24, 2004
<u>AB</u>	!	<u>100MG</u>	<u>A077027</u>	<u>002</u>	Nov 24, 2004

CIMETIDINE

TABLET; ORAL

CIMETIDINE

<u>AB</u>	MYLAN	<u>200MG</u>	<u>A074246</u>	<u>001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>A074246</u>	<u>002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>A074246</u>	<u>003</u>	May 17, 1994
<u>AB</u>	!	<u>800MG</u>	<u>A074246</u>	<u>004</u>	May 17, 1994
<u>AB</u>	TEVA	<u>200MG</u>	<u>A074151</u>	<u>001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>A074151</u>	<u>002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>A074151</u>	<u>003</u>	May 17, 1994
<u>AB</u>		<u>800MG</u>	<u>A074463</u>	<u>001</u>	May 17, 1994

PRESCRIPTION DRUG PRODUCT LIST

CIMETIDINE HYDROCHLORIDE

SOLUTION;ORAL

CIMETIDINE HYDROCHLORIDE

<u>AA</u>	!	PAI HOLDINGS PHARM	<u>EQ 300MG BASE/5ML</u>	<u>A074664</u>	<u>001</u>	Oct 28, 1997
<u>AA</u>		PHARM ASSOC	<u>EQ 300MG BASE/5ML</u>	<u>A074553</u>	<u>001</u>	Jan 27, 1997

CINACALCET HYDROCHLORIDE

TABLET;ORAL

CINACALCET HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 30MG BASE</u>	<u>A211892</u>	<u>001</u>	May 15, 2020
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A211892</u>	<u>002</u>	May 15, 2020
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A211892</u>	<u>003</u>	May 15, 2020
<u>AB</u>		ALKEM LABS LTD	<u>EQ 30MG BASE</u>	<u>A210570</u>	<u>001</u>	May 17, 2019
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A210570</u>	<u>002</u>	May 17, 2019
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A210570</u>	<u>003</u>	May 17, 2019
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 30MG BASE</u>	<u>A206125</u>	<u>001</u>	Mar 08, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A206125</u>	<u>002</u>	Mar 08, 2018
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A206125</u>	<u>003</u>	Mar 08, 2018
<u>AB</u>		CHARTWELL RX	<u>EQ 30MG BASE</u>	<u>A213325</u>	<u>001</u>	May 18, 2020
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A213325</u>	<u>002</u>	May 18, 2020
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A213325</u>	<u>003</u>	May 18, 2020
<u>AB</u>		CIPLA	<u>EQ 30MG BASE</u>	<u>A208915</u>	<u>001</u>	Mar 08, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A208915</u>	<u>002</u>	Mar 08, 2018
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A208915</u>	<u>003</u>	Mar 08, 2018
<u>AB</u>		DR REDDYS	<u>EQ 30MG BASE</u>	<u>A208368</u>	<u>001</u>	Sep 18, 2020
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A208368</u>	<u>002</u>	Sep 18, 2020
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A208368</u>	<u>003</u>	Sep 18, 2020
<u>AB</u>		HETERO LABS LTD V	<u>EQ 30MG BASE</u>	<u>A209403</u>	<u>001</u>	Oct 07, 2020
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A209403</u>	<u>002</u>	Oct 07, 2020
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A209403</u>	<u>003</u>	Oct 07, 2020
<u>AB</u>		SLATE RUN PHARMA	<u>EQ 30MG BASE</u>	<u>A210207</u>	<u>001</u>	Aug 01, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A210207</u>	<u>002</u>	Aug 01, 2018
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A210207</u>	<u>003</u>	Aug 01, 2018
<u>AB</u>		STRIDES PHARMA	<u>EQ 30MG BASE</u>	<u>A209226</u>	<u>001</u>	Apr 30, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A209226</u>	<u>002</u>	Apr 30, 2018
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A209226</u>	<u>003</u>	Apr 30, 2018
<u>AB</u>		SUN PHARM	<u>EQ 30MG BASE</u>	<u>A207008</u>	<u>001</u>	Oct 11, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A207008</u>	<u>002</u>	Oct 11, 2018
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A207008</u>	<u>003</u>	Oct 11, 2018
<u>AB</u>		WATSON LABS TEVA	<u>EQ 30MG BASE</u>	<u>A204377</u>	<u>001</u>	Dec 27, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A204377</u>	<u>002</u>	Dec 27, 2018
<u>AB</u>			<u>EQ 90MG BASE</u>	<u>A204377</u>	<u>003</u>	Dec 27, 2018

SENSIPAR

<u>AB</u>	+	AMGEN	<u>EQ 30MG BASE</u>	<u>N021688</u>	<u>001</u>	Mar 08, 2004
<u>AB</u>	+		<u>EQ 60MG BASE</u>	<u>N021688</u>	<u>002</u>	Mar 08, 2004
<u>AB</u>	+	!	<u>EQ 90MG BASE</u>	<u>N021688</u>	<u>003</u>	Mar 08, 2004

CIPROFLOXACIN

FOR SUSPENSION;ORAL

CIPRO

<u>AB</u>	+	BAYER HLTHCARE	<u>250MG/5ML</u>	<u>N020780</u>	<u>001</u>	Sep 26, 1997
<u>AB</u>	+	!	<u>500MG/5ML</u>	<u>N020780</u>	<u>002</u>	Sep 26, 1997

CIPROFLOXACIN

<u>AB</u>		CHARTWELL	<u>250MG/5ML</u>	<u>A200563</u>	<u>001</u>	Mar 05, 2014
<u>AB</u>			<u>500MG/5ML</u>	<u>A200563</u>	<u>002</u>	Mar 05, 2014

INJECTABLE;INJECTION

CIPROFLOXACIN

<u>AP</u>	!	BAXTER HLTHCARE CORP	<u>200MG/20ML (10MG/ML)</u>	<u>A078062</u>	<u>001</u>	Apr 29, 2008
<u>AP</u>	!		<u>400MG/40ML (10MG/ML)</u>	<u>A078062</u>	<u>002</u>	Apr 29, 2008
<u>AP</u>		HIKMA FARMACEUTICA	<u>200MG/20ML (10MG/ML)</u>	<u>A076717</u>	<u>001</u>	Dec 22, 2009
<u>AP</u>			<u>400MG/40ML (10MG/ML)</u>	<u>A076717</u>	<u>002</u>	Dec 22, 2009

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>		BAXTER HLTHCARE CORP	<u>200MG/100ML</u>	<u>A078024</u>	<u>001</u>	Mar 18, 2008
<u>AP</u>			<u>400MG/200ML</u>	<u>A078024</u>	<u>002</u>	Mar 18, 2008
<u>AP</u>		HIKMA FARMACEUTICA	<u>400MG/200ML</u>	<u>A078431</u>	<u>001</u>	Nov 18, 2009
<u>AP</u>	!	HOSPIRA	<u>200MG/100ML</u>	<u>A077753</u>	<u>001</u>	Mar 18, 2008
<u>AP</u>	!		<u>400MG/200ML</u>	<u>A077753</u>	<u>002</u>	Mar 18, 2008
<u>AP</u>		INFORLIFE	<u>200MG/100ML</u>	<u>A078252</u>	<u>001</u>	Mar 18, 2008
<u>AP</u>			<u>400MG/200ML</u>	<u>A078252</u>	<u>002</u>	Mar 18, 2008

PRESCRIPTION DRUG PRODUCT LIST

CIPROFLOXACIN HYDROCHLORIDE

OINTMENT;OPHTHALMIC

CILOXAN

+! SANDOZ EQ 0.3% BASE N020369 001 Mar 30, 1998

SOLUTION/DROPS;OPHTHALMIC

CILOXAN**AT** +! SANDOZ **EQ 0.3% BASE** **N019992 001** Dec 31, 1990CIPROFLOXACIN HYDROCHLORIDE**AT** ALTAIRE PHARMS INC **EQ 0.3% BASE** **A204613 001** May 03, 2018**AT** FDC LTD **EQ 0.3% BASE** **A077568 001** Jun 30, 2008**AT** RISING **EQ 0.3% BASE** **A077689 001** Dec 13, 2006**AT** RUBICON **EQ 0.3% BASE** **A075928 001** Jun 09, 2004**AT** WATSON LABS INC **EQ 0.3% BASE** **A076673 001** Jan 21, 2005

SOLUTION/DROPS;OTIC

CETRAKAL

+! WRASER PHARMS EQ 0.2% BASE N021918 001 May 01, 2009

TABLET;ORAL

CIPRO**AB** + BAYER HLTHCARE **EQ 250MG BASE** **N019537 002** Oct 22, 1987**AB** +! **EQ 500MG BASE** **N019537 003** Oct 22, 1987CIPROFLOXACIN HYDROCHLORIDE**AB** AMNEAL **EQ 250MG BASE** **A075939 002** Jun 09, 2004**AB** **EQ 500MG BASE** **A075939 003** Jun 09, 2004**AB** **EQ 750MG BASE** **A075939 004** Jun 09, 2004**AB** AUROBINDO PHARMA **EQ 250MG BASE** **A077859 001** Apr 26, 2007**AB** **EQ 500MG BASE** **A077859 002** Apr 26, 2007**AB** **EQ 750MG BASE** **A077859 003** Apr 26, 2007**AB** CARLSBAD **EQ 250MG BASE** **A076126 002** Jun 09, 2004**AB** **EQ 500MG BASE** **A076126 003** Jun 09, 2004**AB** **EQ 750MG BASE** **A076126 004** Jun 09, 2004**AB** CHARTWELL **EQ 250MG BASE** **A076896 001** Nov 04, 2004**AB** **EQ 500MG BASE** **A076896 002** Nov 04, 2004**AB** **EQ 750MG BASE** **A076896 003** Nov 04, 2004**AB** DR REDDYS LABS LTD **EQ 250MG BASE** **A075593 003** Jun 09, 2004**AB** **EQ 500MG BASE** **A075593 004** Jun 09, 2004**AB** **EQ 750MG BASE** **A075593 001** Jun 09, 2004**AB** HIKMA **EQ 250MG BASE** **A076558 002** Jun 09, 2004**AB** **EQ 500MG BASE** **A076558 003** Jun 09, 2004**AB** **EQ 750MG BASE** **A076558 004** Jun 09, 2004**AB** IVAX SUB TEVA **EQ 250MG BASE** **A076089 002** Jun 09, 2004

PHARMS

AB **EQ 500MG BASE** **A076089 003** Jun 09, 2004**AB** **EQ 750MG BASE** **A076089 004** Jun 09, 2004**AB** RISING **EQ 500MG BASE** **A075817 003** Jun 09, 2004**AB** UNIQUE **EQ 250MG BASE** **A076639 001** Sep 10, 2004**AB** **EQ 500MG BASE** **A076639 002** Sep 10, 2004**AB** **EQ 750MG BASE** **A076639 003** Sep 10, 2004**AB** WATSON LABS **EQ 250MG BASE** **A076794 002** Jun 09, 2004**AB** **EQ 500MG BASE** **A076794 003** Jun 09, 2004**AB** **EQ 750MG BASE** **A076794 004** Jun 09, 2004**AB** YILING **EQ 250MG BASE** **A208921 001** Jun 22, 2018**AB** **EQ 500MG BASE** **A208921 002** Jun 22, 2018CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE

SOLUTION/DROPS;OTIC

OTOVEL

+! LABORATORIOS SALVAT EQ 0.3% BASE;0.025% N208251 001 Apr 29, 2016

CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE

SUSPENSION/DROPS;OTIC

CIPRO HC

+! SANDOZ EQ 0.2% BASE;1% N020805 001 Feb 10, 1998

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CIPROFLOXACIN EXTENDED RELEASE**AB** ! ANCHEN PHARMS **425.2MG;EQ 574.9MG BASE** **A078166 001** Nov 27, 2007**AB** DR REDDYS LABS LTD **425.2MG;EQ 574.9MG BASE** **A077701 001** Mar 26, 2007

! ANCHEN PHARMS 212.6MG;EQ 287.5MG BASE A078166 002 Nov 27, 2007

PRESCRIPTION DRUG PRODUCT LIST

CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS; OTIC

CIPRODEX

<u>AB</u>	<u>+</u> !	SANDOZ	<u>0.3%;0.1%</u>	<u>N021537</u>	<u>001</u>	Jul 18, 2003
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CIPROFLOXACIN AND DEXAMETHASONE

<u>AB</u>		DR REDDYS	<u>0.3%;0.1%</u>	<u>A205548</u>	<u>001</u>	Aug 10, 2020
<u>AB</u>		SENTISS	<u>0.3%;0.1%</u>	<u>A215768</u>	<u>001</u>	Jun 09, 2023
<u>AB</u>		SUN PHARM	<u>0.3%;0.1%</u>	<u>A210470</u>	<u>001</u>	Aug 30, 2022

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

<u>AP</u>		CAPLIN	<u>EQ 2MG BASE/ML</u>	<u>A217725</u>	<u>001</u>	Jun 09, 2023
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A217725</u>	<u>002</u>	Jun 09, 2023
<u>AP</u>		EUGIA PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A209144</u>	<u>001</u>	May 08, 2020
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A203183</u>	<u>001</u>	Feb 26, 2015
<u>AP</u>		HENGRUI PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A209334</u>	<u>001</u>	Aug 30, 2017
<u>AP</u>		HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A203078</u>	<u>001</u>	Jun 28, 2022
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A203079</u>	<u>001</u>	Apr 19, 2023
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A203078</u>	<u>002</u>	Dec 18, 2023
<u>AP</u>		HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A203236</u>	<u>001</u>	Mar 30, 2018
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A203238</u>	<u>001</u>	Mar 30, 2018
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A203236</u>	<u>002</u>	Mar 30, 2018
<u>AP</u>		MEITHEAL	<u>EQ 2MG BASE/ML</u>	<u>A211668</u>	<u>001</u>	Apr 25, 2019
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A211669</u>	<u>001</u>	Apr 25, 2019
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A211668</u>	<u>002</u>	Apr 25, 2019
<u>AP</u>		PIRAMAL CRITICAL	<u>EQ 2MG BASE/ML</u>	<u>A215516</u>	<u>001</u>	May 24, 2022
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A215517</u>	<u>001</u>	Dec 14, 2022
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A215516</u>	<u>002</u>	May 24, 2022
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 2MG BASE/ML</u>	<u>A201836</u>	<u>001</u>	Nov 06, 2020
<u>AP</u>			<u>EQ 2MG BASE/ML</u>	<u>A201851</u>	<u>001</u>	Nov 06, 2020
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A201836</u>	<u>002</u>	Nov 06, 2020
<u>AP</u>		SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A200159</u>	<u>001</u>	Feb 03, 2012
<u>AP</u>		SOMERSET	<u>EQ 2MG BASE/ML</u>	<u>A209132</u>	<u>001</u>	Apr 24, 2019
<u>AP</u>		ZYDUS PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A212171</u>	<u>001</u>	Nov 04, 2019
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A212171</u>	<u>002</u>	Nov 04, 2019

CISATRACURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>		EUGIA PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A209665</u>	<u>001</u>	Oct 27, 2020
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A203182</u>	<u>001</u>	Feb 26, 2015
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A203182</u>	<u>002</u>	Feb 26, 2015
<u>AP</u>		HENGRUI PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A204960</u>	<u>001</u>	Jan 27, 2017
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A204960</u>	<u>002</u>	Sep 19, 2017
<u>AP</u>		SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A200154</u>	<u>001</u>	Feb 03, 2012
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A200154</u>	<u>002</u>	Feb 03, 2012
<u>AP</u>		SOMERSET THERAPS LLC	<u>EQ 2MG BASE/ML</u>	<u>A206791</u>	<u>001</u>	Feb 20, 2019
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A206791</u>	<u>002</u>	Feb 20, 2019

NIMBEX

<u>AP</u>	<u>+</u> !	ABBVIE	<u>EQ 2MG BASE/ML</u>	<u>N020551</u>	<u>001</u>	Dec 15, 1995
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NIMBEX PRESERVATIVE FREE

<u>AP</u>	<u>+</u> !	ABBVIE	<u>EQ 2MG BASE/ML</u>	<u>N020551</u>	<u>003</u>	Dec 15, 1995
<u>AP</u>	<u>+</u> !		<u>EQ 10MG BASE/ML</u>	<u>N020551</u>	<u>002</u>	Dec 15, 1995

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

<u>AP</u>		ACCORD HLTHCARE	<u>1MG/ML</u>	<u>A206774</u>	<u>001</u>	Aug 18, 2015
<u>AP</u>	!	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A074735</u>	<u>001</u>	Jul 16, 1999
<u>AP</u>		GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A207323</u>	<u>001</u>	Mar 17, 2017
<u>AP</u>		HIKMA	<u>1MG/ML</u>	<u>A075036</u>	<u>001</u>	Nov 07, 2000
<u>AP</u>	+	HQ SPCLT PHARMA	<u>1MG/ML</u>	<u>N018057</u>	<u>004</u>	Nov 08, 1988
<u>AP</u>		PHARMACHEMIE BV	<u>1MG/ML</u>	<u>A074656</u>	<u>001</u>	May 16, 2000
	<u>+</u> !	HQ SPCLT PHARMA	50MG/VIAL	N018057	002	

CITALOPRAM HYDROBROMIDE

CAPSULE; ORAL

CITALOPRAM HYDROBROMIDE

+! ALMATICA

EQ 30MG BASE

N215428 001 Jan 31, 2022

SOLUTION; ORAL

CITALOPRAM HYDROBROMIDE

<u>AA</u>		AUROBINDO PHARMA	<u>EQ 10MG BASE/5ML</u>	<u>A077812</u>	<u>001</u>	Aug 28, 2006
<u>AA</u>		CHARTWELL MOLECULAR	<u>EQ 10MG BASE/5ML</u>	<u>A077629</u>	<u>001</u>	Jun 14, 2006
<u>AA</u>		HETERO LABS LTD III	<u>EQ 10MG BASE/5ML</u>	<u>A201450</u>	<u>001</u>	Dec 15, 2015
<u>AA</u>	!	HIKMA	<u>EQ 10MG BASE/5ML</u>	<u>A077043</u>	<u>001</u>	Dec 13, 2004

PRESCRIPTION DRUG PRODUCT LIST

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CELEXA

AB	+	ABBVIE	EQ 10MG BASE	N020822 001	Apr 27, 2000
AB	+		EQ 20MG BASE	N020822 002	Jul 17, 1998
AB	+		EQ 40MG BASE	N020822 003	Jul 17, 1998

CITALOPRAM HYDROBROMIDE

AB		AMNEAL PHARMS NY	EQ 10MG BASE	A077289 001	Nov 30, 2006
AB			EQ 20MG BASE	A077289 002	Nov 30, 2006
AB			EQ 40MG BASE	A077289 003	Nov 30, 2006
AB		APOTEX INC	EQ 10MG BASE	A077046 001	Nov 24, 2004
AB		AUROBINDO	EQ 10MG BASE	A077031 001	Oct 28, 2004
AB			EQ 20MG BASE	A077031 002	Oct 28, 2004
AB			EQ 40MG BASE	A077031 003	Oct 28, 2004
AB		CHARTWELL MOLECULAR	EQ 10MG BASE	A077044 001	Nov 05, 2004
AB			EQ 20MG BASE	A077044 002	Nov 05, 2004
AB			EQ 40MG BASE	A077044 003	Nov 05, 2004
AB		COSETTE	EQ 10MG BASE	A077048 001	Nov 16, 2004
AB			EQ 20MG BASE	A077048 002	Nov 16, 2004
AB			EQ 40MG BASE	A077048 003	Nov 16, 2004
AB		DR REDDYS LABS LTD	EQ 10MG BASE	A077038 001	Oct 28, 2004
AB			EQ 20MG BASE	A077038 002	Oct 28, 2004
AB			EQ 40MG BASE	A077038 003	Oct 28, 2004
AB		EPIC PHARMA	EQ 10MG BASE	A077045 003	Apr 29, 2005
AB			EQ 20MG BASE	A077045 002	Apr 29, 2005
AB			EQ 40MG BASE	A077045 001	Apr 29, 2005
AB		GLENMARK GENERICS	EQ 10MG BASE	A077654 001	Feb 27, 2009
AB			EQ 20MG BASE	A077654 002	Feb 27, 2009
AB			EQ 40MG BASE	A077654 003	Feb 27, 2009
AB		INVAGEN PHARMS	EQ 10MG BASE	A077534 001	Oct 03, 2006
AB			EQ 20MG BASE	A077534 002	Oct 03, 2006
AB			EQ 40MG BASE	A077534 003	Oct 03, 2006
AB		MYLAN	EQ 10MG BASE	A077042 001	Nov 05, 2004
AB			EQ 20MG BASE	A077042 002	Nov 05, 2004
AB			EQ 40MG BASE	A077042 003	Nov 05, 2004
AB		TORPHARM	EQ 20MG BASE	A077046 002	Nov 24, 2004
AB			EQ 40MG BASE	A077046 003	Nov 24, 2004
AB		TORRENT PHARMS	EQ 10MG BASE	A078216 001	Mar 27, 2007
AB			EQ 20MG BASE	A078216 002	Mar 27, 2007
AB			EQ 40MG BASE	A078216 003	Mar 27, 2007

CITRIC ACID; GLUCONOLACTONE; MAGNESIUM CARBONATE

SOLUTION; IRRIGATION

RENACIDIN

+	UNITED GUARDIAN	6.602GM/100ML;198MG/100ML;3.177GM/100ML	N019481 001	Oct 02, 1990
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CITRIC ACID; LACTIC ACID; POTASSIUM BITARTRATE

GEL; VAGINAL

PHEXXI

+	EVOFEM INC	1%;1.8%;0.4%	N208352 001	May 22, 2020
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CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

FOR SOLUTION; ORAL

SODIUM PICOSULFATE, MAGNESIUM OXIDE AND ANHYDROUS CITRIC ACID

!	HERETO LABS LTD V	12GM/PACKET;3.5GM/PACKET;10MG/PACKET	A212789 001	Jul 18, 2022
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SOLUTION; ORAL

CLENPIQ

+	FERRING PHARMS INC	12GM/BOT;3.5GM/BOT;10MG/BOT	N209589 001	Nov 28, 2017
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CITRIC ACID; UREA C-13

FOR SOLUTION, TABLET, FOR SOLUTION; ORAL

IDKIT:HP

+	MERIDIAN BIOSCIENCE	4GM;75MG	N021314 001	Dec 17, 2002
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CLADRIBINE

INJECTABLE; INJECTION

CLADRIBINE

AP	!	FRESENIUS KABI USA	1MG/ML	A076571 001	Apr 22, 2004
AP		HIKMA	1MG/ML	A075405 001	Feb 28, 2000
AP		HISUN PHARM HANGZHOU	1MG/ML	A210856 001	Nov 25, 2019

TABLET; ORAL

MAVENCLAD

+	EMD SERONO INC	10MG	N022561 001	Mar 29, 2019
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PRESCRIPTION DRUG PRODUCT LIST

CLARITHROMYCIN

FOR SUSPENSION;ORAL

CLARITHROMYCIN

SANDOZ

125MG/5ML

A065283 002 Sep 04, 2007

!

250MG/5ML

A065283 003 Sep 04, 2007

TABLET;ORAL

CLARITHROMYCINAB ALEMBIC250MGA210459 001 Jan 31, 2022AB500MGA210459 002 Jan 31, 2022AB ! AUROBINDO250MGA065489 001 Jul 25, 2012AB !500MGA065489 002 Jul 25, 2012AB CHARTWELL250MGA065384 001 Aug 20, 2007AB500MGA065384 002 Aug 20, 2007AB HEC PHARM250MGA203584 001 Sep 28, 2015AB500MGA203584 002 Sep 28, 2015AB SANDOZ250MGA065144 001 Oct 18, 2005AB500MGA065136 001 Aug 25, 2005AB STRIDES PHARMA250MGA202710 001 Jun 10, 2013AB500MGA202710 002 Jun 10, 2013

TABLET, EXTENDED RELEASE;ORAL

CLARITHROMYCINAB ACTAVIS LABS FL INC500MGA065145 001 Jun 24, 2004AB ! DR REDDYS LABS SA500MGA065154 001 May 18, 2005AB NOSTRUM LABS INC500MGA203243 001 Feb 29, 2016AB SUNSHINE500MGA208987 001 Jul 09, 2018CLASCOTERONE

CREAM;TOPICAL

WINLEVI

+! SUN PHARM

1%

N213433 001 Aug 26, 2020

CLEMASTINE FUMARATE

SYRUP;ORAL

CLEMASTINE FUMARATE

! GENUS

EQ 0.5MG BASE/5ML

A073399 001 Jun 30, 1994

TABLET;ORAL

CLEMASTINE FUMARATE

! TEVA

2.68MG

A073283 001 Jan 31, 1992

CLEVIDIPINE

EMULSION;INTRAVENOUS

CLEVIPREX

+! CHIESI

25MG/50ML (0.5MG/ML)

N022156 001 Aug 01, 2008

+!

50MG/100ML (0.5MG/ML)

N022156 002 Aug 01, 2008

CLINDAMYCIN HYDROCHLORIDE

CAPSULE;ORAL

CLEOCIN HYDROCHLORIDEAB + PFIZEREQ 75MG BASEN050162 001AB +EQ 150MG BASEN050162 002AB +!EQ 300MG BASEN050162 003 Apr 14, 1988CLINDAMYCIN HYDROCHLORIDEAB AUROBINDO PHARMAEQ 150MG BASEA065442 001 Aug 26, 2009ABEQ 300MG BASEA065442 002 Aug 26, 2009AB CHARTWELL MOLECULAREQ 75MG BASEA065243 002 Aug 12, 2005AB COSETTEEQ 150MG BASEA063029 001 Sep 20, 1989ABEQ 300MG BASEA063029 002 Aug 05, 2005AB EPIC PHARMA LLCEQ 150MG BASEA065194 001 Mar 22, 2004ABEQ 300MG BASEA065194 002 Mar 22, 2004AB GLENMARK PHARMS LTDEQ 75MG BASEA216957 001 Mar 10, 2023ABEQ 150MG BASEA216957 002 Mar 10, 2023ABEQ 300MG BASEA216957 003 Mar 10, 2023AB MICRO LABSEQ 75MG BASEA207402 001 Nov 05, 2018ABEQ 150MG BASEA207402 002 Nov 05, 2018ABEQ 300MG BASEA207402 003 Nov 05, 2018AB SUN PHARM INDS LTDEQ 150MG BASEA065061 001 Feb 02, 2001ABEQ 300MG BASEA065061 002 Feb 02, 2001AB ZYDUS PHARMS USAEQ 75MG BASEA065217 001 Jan 31, 2005ABEQ 150MG BASEA065217 002 Jan 31, 2005ABEQ 300MG BASEA065217 003 Jan 31, 2005

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION;ORAL

CLEOCIN

AA	!	PFIZER	<u>EQ 75MG BASE/5ML</u>	<u>A062644</u>	<u>001</u>	Apr 07, 1986
<u>CLINDAMYCIN PALMITATE HYDROCHLORIDE</u>						
AA		AMNEAL PHARMS	<u>EQ 75MG BASE/5ML</u>	<u>A203513</u>	<u>001</u>	Mar 13, 2014
AA		AUROBINDO PHARMA LTD	<u>EQ 75MG BASE/5ML</u>	<u>A202409</u>	<u>001</u>	Apr 30, 2013
AA		CHARTWELL RX	<u>EQ 75MG BASE/5ML</u>	<u>A206958</u>	<u>001</u>	May 05, 2017
AA		HERITAGE PHARMS INC	<u>EQ 75MG BASE/5ML</u>	<u>A207047</u>	<u>001</u>	May 11, 2018
AA		LYNE	<u>EQ 75MG BASE/5ML</u>	<u>A201821</u>	<u>001</u>	Aug 28, 2012
AA		PADAGIS US	<u>EQ 75MG BASE/5ML</u>	<u>A090902</u>	<u>001</u>	Jul 07, 2010

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM;TOPICAL

CLINDAMYCIN PHOSPHATE

AT		GLENMARK PHARMS LTD	<u>1%</u>	<u>A210778</u>	<u>001</u>	Sep 20, 2021
AT	!	PADAGIS ISRAEL	<u>1%</u>	<u>A090785</u>	<u>001</u>	Mar 31, 2010
AT		TARO	<u>1%</u>	<u>A210004</u>	<u>001</u>	Mar 11, 2020

CREAM;VAGINAL

CLEOCIN

AB	+	!	PFIZER	<u>EQ 2% BASE</u>	<u>N050680</u>	<u>002</u>	Mar 02, 1998
<u>CLINDAMYCIN PHOSPHATE</u>							
AB			FOUGERA PHARMS	<u>EQ 2% BASE</u>	<u>A065139</u>	<u>001</u>	Dec 27, 2004
CLINDESSE							
	+	!	PADAGIS US	<u>EQ 2% BASE</u>	<u>N050793</u>	<u>001</u>	Nov 30, 2004

GEL;TOPICAL

CLEOCIN T

AB1	+	!	PFIZER	<u>EQ 1% BASE</u>	<u>N050615</u>	<u>001</u>	Jan 07, 1987
<u>CLINDAMYCIN PHOSPHATE</u>							
AB1			AMNEAL	<u>EQ 1% BASE</u>	<u>A215219</u>	<u>001</u>	Nov 03, 2022
AB1			ENCUBE	<u>EQ 1% BASE</u>	<u>A212438</u>	<u>001</u>	Mar 11, 2021
AB1			FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A064160</u>	<u>001</u>	Jan 28, 2000
AB1			GLENMARK PHARMS LTD	<u>EQ 1% BASE</u>	<u>A214251</u>	<u>001</u>	Feb 10, 2021
AB1			PADAGIS ISRAEL	<u>EQ 1% BASE</u>	<u>A212104</u>	<u>001</u>	Dec 31, 2020
AB1			SOLARIS PHARMA CORP	<u>EQ 1% BASE</u>	<u>A211872</u>	<u>001</u>	Jul 29, 2020
AB1			TARO	<u>EQ 1% BASE</u>	<u>A214052</u>	<u>001</u>	Nov 10, 2020
AB1			ZYDUS LIFESCIENCES	<u>EQ 1% BASE</u>	<u>A216587</u>	<u>001</u>	Sep 19, 2023

CLINDAGEL

AB2	+	!	BAUSCH	<u>EQ 1% BASE</u>	<u>N050782</u>	<u>001</u>	Nov 27, 2000
<u>CLINDAMYCIN PHOSPHATE</u>							
AB2			AMNEAL	<u>EQ 1% BASE</u>	<u>A214668</u>	<u>001</u>	Aug 05, 2022
AB2			SOLARIS PHARMA CORP	<u>EQ 1% BASE</u>	<u>A212842</u>	<u>001</u>	Aug 13, 2021
GEL;VAGINAL							
XACIATO							
	+	!	DARE	<u>EQ 2% BASE</u>	<u>N215650</u>	<u>001</u>	Dec 07, 2021

INJECTABLE;INJECTION

CLEOCIN PHOSPHATE

AP			PFIZER	<u>EQ 150MG BASE/ML</u>	<u>A062803</u>	<u>001</u>	Oct 16, 1987
AP	+	!		<u>EQ 150MG BASE/ML</u>	<u>N050441</u>	<u>001</u>	
<u>CLINDAMYCIN PHOSPHATE</u>							
AP			ALMAJECT	<u>EQ 150MG BASE/ML</u>	<u>A062800</u>	<u>001</u>	Jul 24, 1987
AP				<u>EQ 150MG BASE/ML</u>	<u>A062943</u>	<u>001</u>	Sep 29, 1988
AP			FRESENIUS KABI USA	<u>EQ 150MG BASE/ML</u>	<u>A065346</u>	<u>001</u>	Mar 29, 2007
AP				<u>EQ 150MG BASE/ML</u>	<u>A065347</u>	<u>001</u>	May 09, 2007
AP			HIKMA	<u>EQ 150MG BASE/ML</u>	<u>A062889</u>	<u>001</u>	Apr 25, 1988
AP				<u>EQ 150MG BASE/ML</u>	<u>A065206</u>	<u>001</u>	Sep 24, 2004
AP			SAGENT PHARMS INC	<u>EQ 150MG BASE/ML</u>	<u>A090108</u>	<u>001</u>	Sep 30, 2011
AP				<u>EQ 150MG BASE/ML</u>	<u>A090109</u>	<u>001</u>	Sep 30, 2011
<u>CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER</u>							
AP			BAXTER HLTHCARE CORP	<u>EQ 6MG BASE/ML</u>	<u>A208084</u>	<u>001</u>	Jun 28, 2017
AP				<u>EQ 12MG BASE/ML</u>	<u>A208084</u>	<u>002</u>	Jun 28, 2017
AP				<u>EQ 18MG BASE/ML</u>	<u>A208084</u>	<u>003</u>	Jun 28, 2017
AP			RISING	<u>EQ 6MG BASE/ML</u>	<u>A203048</u>	<u>001</u>	Apr 04, 2013
AP				<u>EQ 12MG BASE/ML</u>	<u>A203048</u>	<u>002</u>	Apr 04, 2013
AP				<u>EQ 18MG BASE/ML</u>	<u>A203048</u>	<u>003</u>	Apr 04, 2013
AP	!		SANDOZ INC	<u>EQ 6MG BASE/ML</u>	<u>A201692</u>	<u>001</u>	May 31, 2012
AP	!			<u>EQ 12MG BASE/ML</u>	<u>A201692</u>	<u>002</u>	May 31, 2012
AP	!			<u>EQ 18MG BASE/ML</u>	<u>A201692</u>	<u>003</u>	May 31, 2012
<u>CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%</u>							
AP			HIKMA	<u>EQ 6MG BASE/ML</u>	<u>A214401</u>	<u>001</u>	Apr 05, 2023
AP				<u>EQ 12MG BASE/ML</u>	<u>A214401</u>	<u>002</u>	Apr 05, 2023

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

AP		<u>EQ 18MG BASE/ML</u>	<u>A214401 003</u>	Apr 05, 2023
	+!	ABRAXIS PHARM	EQ 900MG BASE/100ML	N050635 001 Dec 22, 1989

LOTION; TOPICAL

CLEOCIN T

AB	+!	PFIZER	<u>EQ 1% BASE</u>	<u>N050600 001</u>	May 31, 1989
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CLINDAMYCIN PHOSPHATE

AB		ENCUBE	<u>EQ 1% BASE</u>	<u>A215607 001</u>	Jan 18, 2022
AB		FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A065067 001</u>	Jan 31, 2002
AB		PADAGIS ISRAEL	<u>EQ 1% BASE</u>	<u>A214604 001</u>	Mar 08, 2021
AB		TARO	<u>EQ 1% BASE</u>	<u>A214526 001</u>	Jun 28, 2021

SOLUTION; INTRAVENOUS

CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE

	+!	BAXTER HLTHCARE	EQ 300MG BASE/50ML (EQ 6MG BASE/ML)	N208083 001	Apr 20, 2017
		CORP			
	+!		EQ 600MG BASE/50ML (EQ 12MG BASE/ML)	N208083 002	Apr 20, 2017
	+!		EQ 900MG BASE/50ML (EQ 18MG BASE/ML)	N208083 003	Apr 20, 2017

SOLUTION; TOPICAL

CLINDA-DERM

AT		PADAGIS US	<u>EQ 1% BASE</u>	<u>A063329 001</u>	Sep 30, 1992
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CLINDAMYCIN PHOSPHATE

AT		CADILA	<u>EQ 1% BASE</u>	<u>A208767 001</u>	Jul 16, 2018
AT		CHARTWELL RX	<u>EQ 1% BASE</u>	<u>A209846 001</u>	Feb 08, 2018
AT		ENCUBE ETHICALS	<u>EQ 1% BASE</u>	<u>A209914 001</u>	Jan 28, 2019
AT		FOUGERA PHARMS INC	<u>EQ 1% BASE</u>	<u>A064159 001</u>	Jun 05, 1997
AT	!	PADAGIS US	<u>EQ 1% BASE</u>	<u>A064050 001</u>	Nov 30, 1995
AT		TARO PHARM INDS	<u>EQ 1% BASE</u>	<u>A065184 001</u>	Mar 31, 2004

SUPPOSITORY; VAGINAL

CLEOCIN

	+!	PFIZER	100MG	N050767 001	Aug 13, 1999
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SWAB; TOPICAL

CLINDAMYCIN PHOSPHATE

AT		EPIC PHARMA LLC	<u>EQ 1% BASE</u>	<u>A065513 001</u>	Jun 17, 2010
AT	!	PADAGIS US	<u>EQ 1% BASE</u>	<u>A065049 001</u>	May 25, 2000

CLINDETS

AT		PADAGIS US	<u>EQ 1% BASE</u>	<u>A064136 001</u>	Sep 30, 1996
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CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE AND TRETINOIN

AB1		ACTAVIS MID	<u>1.2%;0.025%</u>	<u>A202564 001</u>	Jun 12, 2015
		ATLANTIC			

ZIANA

AB1	+!	BAUSCH	<u>1.2%;0.025%</u>	<u>N050802 001</u>	Nov 07, 2006
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CLINDAMYCIN PHOSPHATE AND TRETINOIN

AB2		ENCUBE	<u>1.2%;0.025%</u>	<u>A216943 001</u>	Sep 01, 2023
AB2		SOLARIS PHARMA CORP	<u>1.2%;0.025%</u>	<u>A212845 001</u>	Feb 10, 2022

VELTIN

AB2	+!	ALMIRALL	<u>1.2%;0.025%</u>	<u>N050803 001</u>	Jul 16, 2010
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CLOBAZAM

FILM; ORAL

SYMPAZAN

	+	OTTER PHARMS	5MG	N210833 001	Nov 01, 2018
	+		10MG	N210833 002	Nov 01, 2018
	+!		20MG	N210833 003	Nov 01, 2018

SUSPENSION; ORAL

CLOBAZAM

AB		ALKEM LABS LTD	<u>2.5MG/ML</u>	<u>A213039 001</u>	May 06, 2021
AB		AMNEAL	<u>2.5MG/ML</u>	<u>A210039 001</u>	Oct 22, 2018
AB		AUROBINDO PHARMA	<u>2.5MG/ML</u>	<u>A214404 001</u>	Mar 24, 2022
		LTD			
AB		BIONPHARMA	<u>2.5MG/ML</u>	<u>A208819 001</u>	Oct 22, 2018
AB		CHARTWELL MOLECULAR	<u>2.5MG/ML</u>	<u>A213110 001</u>	Apr 24, 2020
AB		HETERO LABS LTD III	<u>2.5MG/ML</u>	<u>A209796 001</u>	Feb 24, 2020
AB		LUPIN LTD	<u>2.5MG/ML</u>	<u>A210546 001</u>	Dec 28, 2018
AB		TARO	<u>2.5MG/ML</u>	<u>A210978 001</u>	Apr 15, 2019

ONFI

AB	+!	LUNDBECK PHARMS LLC	<u>2.5MG/ML</u>	<u>N203993 001</u>	Dec 14, 2012
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PRESCRIPTION DRUG PRODUCT LIST

CLOBAZAM

TABLET; ORAL

CLOBAZAM

<u>AB</u>	ALKEM LABS LTD	<u>10MG</u>	<u>A212714 001</u>	Sep 06, 2019
<u>AB</u>		<u>20MG</u>	<u>A212714 002</u>	Sep 06, 2019
<u>AB</u>	AMNEAL PHARMS CO	<u>10MG</u>	<u>A209718 001</u>	Oct 22, 2018
<u>AB</u>		<u>20MG</u>	<u>A209718 002</u>	Oct 22, 2018
<u>AB</u>	BIONPHARMA	<u>10MG</u>	<u>A208825 001</u>	Oct 22, 2018
<u>AB</u>		<u>20MG</u>	<u>A208825 002</u>	Oct 22, 2018
<u>AB</u>	BRECKENRIDGE	<u>10MG</u>	<u>A209308 001</u>	Oct 22, 2018
<u>AB</u>		<u>20MG</u>	<u>A209308 002</u>	Oct 22, 2018
<u>AB</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A209795 001</u>	Oct 22, 2018
<u>AB</u>		<u>20MG</u>	<u>A209795 002</u>	Oct 22, 2018
<u>AB</u>	LUPIN LTD	<u>10MG</u>	<u>A210545 001</u>	Dec 14, 2018
<u>AB</u>		<u>20MG</u>	<u>A210545 002</u>	Dec 14, 2018
<u>AB</u>	MICRO LABS	<u>10MG</u>	<u>A211711 001</u>	Jan 30, 2019
<u>AB</u>		<u>20MG</u>	<u>A211711 002</u>	Jan 30, 2019
<u>AB</u>	MSN	<u>10MG</u>	<u>A213404 001</u>	May 11, 2021
<u>AB</u>		<u>20MG</u>	<u>A213404 002</u>	May 11, 2021
<u>AB</u>	PIRAMAL HLTHCARE UK	<u>10MG</u>	<u>A209808 001</u>	Oct 22, 2018
<u>AB</u>		<u>20MG</u>	<u>A209808 002</u>	Oct 22, 2018
<u>AB</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A209687 001</u>	Oct 22, 2018
<u>AB</u>		<u>20MG</u>	<u>A209687 002</u>	Oct 22, 2018
<u>AB</u>	ZYDUS PHARMS	<u>10MG</u>	<u>A211449 001</u>	Oct 22, 2018
<u>AB</u>		<u>20MG</u>	<u>A211449 002</u>	Oct 22, 2018

ONFI

<u>AB</u>	+ LUNDBECK PHARMS LLC	<u>10MG</u>	<u>N202067 002</u>	Oct 21, 2011
<u>AB</u>	+!	<u>20MG</u>	<u>N202067 003</u>	Oct 21, 2011

CLOBAZAM

	ALKEM LABS LTD	5MG	A212714 003	Dec 21, 2020
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CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

<u>AB1</u>	ALEMBIC	<u>0.05%</u>	<u>A215838 001</u>	Apr 21, 2022
<u>AB1</u>	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A210809 001</u>	Feb 15, 2019
<u>AB1</u>	PADAGIS ISRAEL	<u>0.05%</u>	<u>A077763 001</u>	Mar 10, 2008
<u>AB1</u>	! TARO	<u>0.05%</u>	<u>A208779 001</u>	Oct 04, 2018
<u>AB2</u>	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A211450 001</u>	Sep 09, 2019
<u>AB2</u>	! PADAGIS ISRAEL	<u>0.05%</u>	<u>A201402 001</u>	Aug 14, 2012

CREAM; TOPICAL

CLOBETASOL PROPIONATE

<u>AB1</u>	ALEMBIC	<u>0.05%</u>	<u>A213291 001</u>	Jan 27, 2020
<u>AB1</u>	AMNEAL	<u>0.05%</u>	<u>A211256 001</u>	Dec 26, 2018
<u>AB1</u>	AUROBINDO PHARMA USA	<u>0.05%</u>	<u>A075338 001</u>	Feb 09, 2001
<u>AB1</u>	COSETTE	<u>0.05%</u>	<u>A074139 001</u>	Aug 03, 1994
<u>AB1</u>	! ENCUBE	<u>0.05%</u>	<u>A212982 001</u>	Aug 28, 2020
<u>AB1</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A074392 001</u>	Sep 30, 1996
<u>AB1</u>	GLENMARK PHARMS	<u>0.05%</u>	<u>A209095 001</u>	May 10, 2018
<u>AB1</u>	LUPIN LTD	<u>0.05%</u>	<u>A210208 001</u>	Jan 30, 2018
<u>AB1</u>	RISING	<u>0.05%</u>	<u>A211401 001</u>	Jan 11, 2019
<u>AB1</u>	TARO	<u>0.05%</u>	<u>A074249 001</u>	Jul 08, 1996
<u>AB1</u>	XIROMED	<u>0.05%</u>	<u>A210034 001</u>	Jun 15, 2018
<u>AB1</u>	ZYDUS PHARMS	<u>0.05%</u>	<u>A211074 001</u>	Oct 15, 2018

CORMAX

<u>AB1</u>	HIKMA	<u>0.05%</u>	<u>A074220 001</u>	May 16, 1997
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CLOBETASOL PROPIONATE (EMOLLIENT)

<u>AB2</u>	! FOUGERA PHARMS	<u>0.05%</u>	<u>A075430 001</u>	May 26, 1999
<u>AB2</u>	TARO	<u>0.05%</u>	<u>A075633 001</u>	May 17, 2000

IMPOYZ

	+! PRIMUS PHARMS	0.025%	N209483 001	Nov 28, 2017
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GEL; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	! FOUGERA PHARMS	<u>0.05%</u>	<u>A075368 001</u>	Feb 15, 2000
<u>AB</u>	PADAGIS US	<u>0.05%</u>	<u>A075027 001</u>	Oct 31, 1997
<u>AB</u>	TARO	<u>0.05%</u>	<u>A075279 001</u>	May 28, 1999

EMBELINE

<u>AB</u>	HIKMA	<u>0.05%</u>	<u>A076141 001</u>	Apr 12, 2002
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LOTION; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>0.05%</u>	<u>A078223 001</u>	Dec 04, 2008
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PRESCRIPTION DRUG PRODUCT LIST

CLOBETASOL PROPIONATE

LOTION; TOPICAL

CLOBETASOL PROPIONATE

AB	CADILA	<u>0.05%</u>	<u>A205249</u>	<u>001</u>	Sep 24, 2019
AB	LUPIN LTD	<u>0.05%</u>	<u>A209147</u>	<u>001</u>	Sep 22, 2017
AB	TARO	<u>0.05%</u>	<u>A200302</u>	<u>001</u>	Jul 02, 2012

CLOBEX

AB	+ ! GALDERMA LABS LP	<u>0.05%</u>	<u>N021535</u>	<u>001</u>	Jul 24, 2003
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OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

AB	ALEMBIC	<u>0.05%</u>	<u>A211800</u>	<u>001</u>	Mar 04, 2019
AB	AUROBINDO PHARMA USA	<u>0.05%</u>	<u>A075057</u>	<u>001</u>	Aug 12, 1998
AB	COSETTE	<u>0.05%</u>	<u>A074089</u>	<u>001</u>	Feb 16, 1994
AB	ENCUBE	<u>0.05%</u>	<u>A211295</u>	<u>001</u>	Nov 15, 2019
AB	! FOUGERA PHARMS	<u>0.05%</u>	<u>A074407</u>	<u>001</u>	Feb 23, 1996
AB	GLENMARK PHARMS	<u>0.05%</u>	<u>A208933</u>	<u>001</u>	Mar 20, 2017
AB	MACLEODS PHARMS LTD	<u>0.05%</u>	<u>A215990</u>	<u>001</u>	Dec 28, 2023
AB	NOVEL LABS INC	<u>0.05%</u>	<u>A208841</u>	<u>001</u>	May 04, 2018
AB	TARO	<u>0.05%</u>	<u>A074248</u>	<u>001</u>	Jul 12, 1996
AB	XIROMED	<u>0.05%</u>	<u>A209701</u>	<u>001</u>	Apr 17, 2018
AB	ZYDUS PHARMS	<u>0.05%</u>	<u>A210199</u>	<u>001</u>	Oct 27, 2017

EMBELINE

AB	HIKMA	<u>0.05%</u>	<u>A074221</u>	<u>001</u>	Mar 31, 1995
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SHAMPOO; TOPICAL

CLOBETASOL PROPIONATE

AB	ALEMBIC	<u>0.05%</u>	<u>A213290</u>	<u>001</u>	May 18, 2020
AB	AMNEAL	<u>0.05%</u>	<u>A214895</u>	<u>001</u>	Jun 14, 2021
AB	HIKMA	<u>0.05%</u>	<u>A209871</u>	<u>001</u>	Oct 27, 2017
AB	PADAGIS ISRAEL	<u>0.05%</u>	<u>A090974</u>	<u>001</u>	Aug 09, 2012
AB	TARO	<u>0.05%</u>	<u>A214867</u>	<u>001</u>	Mar 25, 2021

CLOBEX

AB	+ ! GALDERMA LABS	<u>0.05%</u>	<u>N021644</u>	<u>001</u>	Feb 05, 2004
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SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

AT	ALEMBIC	<u>0.05%</u>	<u>A212881</u>	<u>001</u>	Oct 21, 2019
AT	COSETTE	<u>0.05%</u>	<u>A074331</u>	<u>001</u>	Dec 15, 1995
AT	FOUGERA PHARMS	<u>0.05%</u>	<u>A075391</u>	<u>001</u>	Feb 08, 1999
AT	GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A210190</u>	<u>001</u>	Apr 18, 2018
AT	MACLEODS PHARMS LTD	<u>0.05%</u>	<u>A209361</u>	<u>001</u>	Oct 25, 2017
AT	NOVEL LABS INC	<u>0.05%</u>	<u>A206075</u>	<u>001</u>	Nov 23, 2015
AT	QUAGEN	<u>0.05%</u>	<u>A211240</u>	<u>001</u>	Feb 17, 2022
AT	SAPTALIS PHARMS	<u>0.05%</u>	<u>A211494</u>	<u>001</u>	Oct 02, 2019
AT	! TARO	<u>0.05%</u>	<u>A075224</u>	<u>001</u>	Nov 16, 1998
AT		<u>0.05%</u>	<u>A075363</u>	<u>001</u>	Dec 29, 2000

EMBELINE

AT	HIKMA	<u>0.05%</u>	<u>A074222</u>	<u>001</u>	Dec 06, 1995
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SPRAY; TOPICAL

CLOBETASOL PROPIONATE

AT	ALEMBIC	<u>0.05%</u>	<u>A211191</u>	<u>001</u>	Oct 02, 2019
AT	GLENMARK PHARMS	<u>0.05%</u>	<u>A209004</u>	<u>001</u>	Mar 26, 2018
AT	LUPIN LTD	<u>0.05%</u>	<u>A208125</u>	<u>001</u>	Mar 26, 2018
AT	PADAGIS US	<u>0.05%</u>	<u>A090898</u>	<u>001</u>	Jun 16, 2011
AT	TARO	<u>0.05%</u>	<u>A208842</u>	<u>001</u>	Mar 26, 2018
AT	ZYDUS PHARMS	<u>0.05%</u>	<u>A206378</u>	<u>001</u>	Feb 16, 2017

CLOBEX

AT	+ ! GALDERMA LABS LP	<u>0.05%</u>	<u>N021835</u>	<u>001</u>	Oct 27, 2005
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CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLOCORTOLONE PIVALATE

AB	TARO	<u>0.1%</u>	<u>A206370</u>	<u>001</u>	Apr 21, 2020
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CLODERM

AB	+ ! EPI HLTH	<u>0.1%</u>	<u>N017765</u>	<u>001</u>	
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CLOFARABINE

SOLUTION; INTRAVENOUS

CLOFARABINE

AP	ABON PHARMS LLC	<u>20MG/20ML (1MG/ML)</u>	<u>A204029</u>	<u>001</u>	May 09, 2017
AP	ACCORD HLTHCARE	<u>20MG/20ML (1MG/ML)</u>	<u>A212034</u>	<u>001</u>	Feb 22, 2019
AP	AMNEAL	<u>20MG/20ML (1MG/ML)</u>	<u>A208857</u>	<u>001</u>	Nov 06, 2017
AP	DR REDDYS	<u>20MG/20ML (1MG/ML)</u>	<u>A205375</u>	<u>001</u>	Nov 06, 2017
AP	GLAND PHARMA LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A207831</u>	<u>001</u>	Oct 31, 2018
AP	MEITHEAL	<u>20MG/20ML (1MG/ML)</u>	<u>A213461</u>	<u>001</u>	Oct 23, 2020

PRESCRIPTION DRUG PRODUCT LIST

CLOFARABINE

SOLUTION; INTRAVENOUS

CLOFARABINE

<u>AP</u>	MSN	<u>20MG/20ML (1MG/ML)</u>	<u>A209775 001</u>	Dec 06, 2017
<u>AP</u>	MYLAN LABS LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A208860 001</u>	Nov 06, 2017

CLOLAR

<u>AP</u>	+! GENZYME	<u>20MG/20ML (1MG/ML)</u>	<u>N021673 001</u>	Dec 28, 2004
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CLOMIPHENE CITRATE

TABLET; ORAL

CLOMIPHENE CITRATE

! COSETTE

50MG

A075528 001 Aug 30, 1999

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

ANAFRANIL

<u>AB</u>	+! SPECGX LLC	<u>25MG</u>	<u>N019906 001</u>	Dec 29, 1989
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<u>AB</u>	+	<u>50MG</u>	<u>N019906 002</u>	Dec 29, 1989
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<u>AB</u>	+	<u>75MG</u>	<u>N019906 003</u>	Dec 29, 1989
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CLOMIPRAMINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>25MG</u>	<u>A213897 001</u>	Oct 02, 2020
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<u>AB</u>		<u>50MG</u>	<u>A213897 002</u>	Oct 02, 2020
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<u>AB</u>		<u>75MG</u>	<u>A213897 003</u>	Oct 02, 2020
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<u>AB</u>	ALEMBIC	<u>25MG</u>	<u>A211822 001</u>	Aug 04, 2021
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<u>AB</u>		<u>50MG</u>	<u>A211822 002</u>	Aug 04, 2021
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<u>AB</u>		<u>75MG</u>	<u>A211822 003</u>	Aug 04, 2021
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<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A208632 001</u>	Oct 31, 2018
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<u>AB</u>		<u>50MG</u>	<u>A208632 002</u>	Oct 31, 2018
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<u>AB</u>		<u>75MG</u>	<u>A208632 003</u>	Oct 31, 2018
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<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A216440 001</u>	Mar 20, 2023
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<u>AB</u>		<u>50MG</u>	<u>A216440 002</u>	Mar 20, 2023
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<u>AB</u>		<u>75MG</u>	<u>A216440 003</u>	Mar 20, 2023
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<u>AB</u>	CHEMISTRY HLTH	<u>25MG</u>	<u>A211364 001</u>	Feb 07, 2020
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<u>AB</u>		<u>50MG</u>	<u>A211364 002</u>	Feb 07, 2020
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<u>AB</u>		<u>75MG</u>	<u>A211364 003</u>	Feb 07, 2020
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<u>AB</u>	JUBILANT CADISTA	<u>25MG</u>	<u>A212218 001</u>	Oct 21, 2019
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<u>AB</u>		<u>50MG</u>	<u>A212218 002</u>	Oct 21, 2019
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<u>AB</u>		<u>75MG</u>	<u>A212218 003</u>	Oct 21, 2019
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<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A209294 001</u>	Nov 21, 2018
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<u>AB</u>		<u>50MG</u>	<u>A209294 002</u>	Nov 21, 2018
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<u>AB</u>		<u>75MG</u>	<u>A209294 003</u>	Nov 21, 2018
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<u>AB</u>	MANKIND PHARMA	<u>25MG</u>	<u>A211767 001</u>	Apr 08, 2019
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<u>AB</u>		<u>50MG</u>	<u>A211767 002</u>	Apr 08, 2019
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<u>AB</u>		<u>75MG</u>	<u>A211767 003</u>	Apr 08, 2019
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<u>AB</u>	MICRO LABS	<u>25MG</u>	<u>A213219 001</u>	Jun 22, 2020
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<u>AB</u>		<u>50MG</u>	<u>A213219 002</u>	Jun 22, 2020
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<u>AB</u>		<u>75MG</u>	<u>A213219 003</u>	Jun 22, 2020
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<u>AB</u>	MYLAN	<u>25MG</u>	<u>A074947 001</u>	Apr 30, 1998
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<u>AB</u>		<u>50MG</u>	<u>A074947 002</u>	Apr 30, 1998
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<u>AB</u>		<u>75MG</u>	<u>A074947 003</u>	Apr 30, 1998
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<u>AB</u>	RK PHARMA	<u>25MG</u>	<u>A213221 001</u>	Jun 22, 2020
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<u>AB</u>		<u>50MG</u>	<u>A213221 002</u>	Jun 22, 2020
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<u>AB</u>		<u>75MG</u>	<u>A213221 003</u>	Jun 22, 2020
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<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A074364 001</u>	Mar 29, 1996
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<u>AB</u>		<u>25MG</u>	<u>A074953 001</u>	Jun 25, 1997
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<u>AB</u>		<u>50MG</u>	<u>A074364 002</u>	Mar 29, 1996
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<u>AB</u>		<u>50MG</u>	<u>A074953 002</u>	Jun 25, 1997
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<u>AB</u>		<u>75MG</u>	<u>A074364 003</u>	Mar 29, 1996
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<u>AB</u>		<u>75MG</u>	<u>A074953 003</u>	Jun 25, 1997
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<u>AB</u>	TARO	<u>25MG</u>	<u>A074694 001</u>	Dec 31, 1996
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<u>AB</u>		<u>50MG</u>	<u>A074694 002</u>	Dec 31, 1996
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<u>AB</u>		<u>75MG</u>	<u>A074694 003</u>	Dec 31, 1996
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<u>AB</u>	TULEX PHARMS INC	<u>25MG</u>	<u>A210653 001</u>	Apr 03, 2020
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<u>AB</u>		<u>50MG</u>	<u>A210653 002</u>	Apr 03, 2020
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<u>AB</u>		<u>75MG</u>	<u>A210653 003</u>	Apr 03, 2020
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<u>AB</u>	UNIQUE	<u>25MG</u>	<u>A212285 001</u>	Aug 07, 2020
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<u>AB</u>		<u>50MG</u>	<u>A212285 002</u>	Aug 07, 2020
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<u>AB</u>		<u>75MG</u>	<u>A212285 003</u>	Aug 07, 2020
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<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A208961 001</u>	Dec 27, 2017
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<u>AB</u>		<u>50MG</u>	<u>A208961 002</u>	Dec 27, 2017
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<u>AB</u>		<u>75MG</u>	<u>A208961 003</u>	Dec 27, 2017
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PRESCRIPTION DRUG PRODUCT LIST

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

<u>AB</u>	ACCORD HLTHCARE	<u>0.5MG</u>	<u>A077147 001</u>	May 02, 2005
<u>AB</u>		<u>1MG</u>	<u>A077147 002</u>	May 02, 2005
<u>AB</u>		<u>2MG</u>	<u>A077147 003</u>	May 02, 2005
<u>AB</u>	ACTAVIS ELIZABETH	<u>0.5MG</u>	<u>A074869 001</u>	Oct 31, 1996
<u>AB</u>		<u>1MG</u>	<u>A074869 002</u>	Oct 31, 1996
<u>AB</u>		<u>2MG</u>	<u>A074869 003</u>	Oct 31, 1996
<u>AB</u>	AUROBINDO PHARMA USA	<u>0.5MG</u>	<u>A075150 001</u>	Oct 05, 1998
<u>AB</u>		<u>1MG</u>	<u>A075150 002</u>	Oct 05, 1998
<u>AB</u>		<u>2MG</u>	<u>A075150 003</u>	Oct 05, 1998
<u>AB</u>	PRINSTON INC	<u>0.5MG</u>	<u>A077856 001</u>	Jun 28, 2006
<u>AB</u>		<u>1MG</u>	<u>A077856 002</u>	Jun 28, 2006
<u>AB</u>		<u>2MG</u>	<u>A077856 003</u>	Jun 28, 2006
<u>AB</u>	RUBICON	<u>0.5MG</u>	<u>A075468 001</u>	Oct 06, 2000
<u>AB</u>		<u>1MG</u>	<u>A075468 002</u>	Oct 06, 2000
<u>AB</u>		<u>2MG</u>	<u>A075468 003</u>	Oct 06, 2000
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A074979 001</u>	Aug 29, 1997
<u>AB</u>		<u>1MG</u>	<u>A074979 002</u>	Aug 29, 1997
<u>AB</u>		<u>2MG</u>	<u>A074979 003</u>	Aug 29, 1997
<u>AB</u>	TEVA	<u>0.5MG</u>	<u>A074569 001</u>	Sep 10, 1996
<u>AB</u>		<u>1MG</u>	<u>A074569 002</u>	Sep 10, 1996
<u>AB</u>		<u>2MG</u>	<u>A074569 003</u>	Sep 10, 1996

KLONOPIN

<u>AB</u>	+ CHEPLAPHARM	<u>0.5MG</u>	<u>N017533 001</u>	
<u>AB</u>	+!	<u>1MG</u>	<u>N017533 002</u>	
<u>AB</u>	+	<u>2MG</u>	<u>N017533 003</u>	

TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

<u>AB</u>	ALEMBIC	<u>0.125MG</u>	<u>A211033 001</u>	Jun 28, 2019
<u>AB</u>		<u>0.25MG</u>	<u>A211033 002</u>	Jun 28, 2019
<u>AB</u>		<u>0.5MG</u>	<u>A211033 003</u>	Jun 28, 2019
<u>AB</u>		<u>1MG</u>	<u>A211033 004</u>	Jun 28, 2019
<u>AB</u>		<u>2MG</u>	<u>A211033 005</u>	Jun 28, 2019
<u>AB</u>	BARR	<u>0.125MG</u>	<u>A077194 001</u>	Aug 10, 2005
<u>AB</u>		<u>0.25MG</u>	<u>A077194 002</u>	Aug 10, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077194 003</u>	Aug 10, 2005
<u>AB</u>		<u>1MG</u>	<u>A077194 004</u>	Aug 10, 2005
<u>AB</u>		<u>2MG</u>	<u>A077194 005</u>	Aug 10, 2005
<u>AB</u>	PAR PHARM	<u>0.125MG</u>	<u>A077171 001</u>	Aug 03, 2005
<u>AB</u>		<u>0.25MG</u>	<u>A077171 002</u>	Aug 03, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077171 003</u>	Aug 03, 2005
<u>AB</u>	!	<u>1MG</u>	<u>A077171 004</u>	Aug 03, 2005
<u>AB</u>		<u>2MG</u>	<u>A077171 005</u>	Aug 03, 2005
<u>AB</u>	SUN PHARM INDS INC	<u>0.125MG</u>	<u>A078654 001</u>	Aug 27, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A078654 002</u>	Aug 27, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A078654 003</u>	Aug 27, 2014
<u>AB</u>		<u>1MG</u>	<u>A078654 004</u>	Aug 27, 2014
<u>AB</u>		<u>2MG</u>	<u>A078654 005</u>	Aug 27, 2014

CLONIDINE

SYSTEM; TRANSDERMAL

CATAPRES-TTS-1

<u>AB</u>	+ LAVIPHARM	<u>0.1MG/24HR</u>	<u>N018891 001</u>	Oct 10, 1984
<u>AB</u>				
<u>AB</u>	+ LAVIPHARM	<u>0.2MG/24HR</u>	<u>N018891 002</u>	Oct 10, 1984
<u>AB</u>				
<u>AB</u>	+! LAVIPHARM	<u>0.3MG/24HR</u>	<u>N018891 003</u>	Oct 10, 1984

CLONIDINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>0.1MG/24HR</u>	<u>A090873 001</u>	May 06, 2014
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A090873 002</u>	May 06, 2014
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A090873 003</u>	May 06, 2014
<u>AB</u>	AVEVA	<u>0.1MG/24HR</u>	<u>A076157 001</u>	Aug 18, 2009
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076157 002</u>	Aug 18, 2009
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076157 003</u>	Aug 18, 2009
<u>AB</u>	DR REDDYS LABS SA	<u>0.1MG/24HR</u>	<u>A079090 001</u>	Aug 20, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A079090 002</u>	Aug 20, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A079090 003</u>	Aug 20, 2010
<u>AB</u>	MYLAN TECHNOLOGIES	<u>0.1MG/24HR</u>	<u>A076166 001</u>	Jul 16, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076166 002</u>	Jul 16, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076166 003</u>	Jul 16, 2010

PRESCRIPTION DRUG PRODUCT LIST

CLONIDINE

TABLET, EXTENDED RELEASE;ORAL

NEXICLON XR

ATHENA

EQ 0.17MG BASE

N022500 001 Dec 03, 2009

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

<u>AP</u>	FRESENIUS KABI USA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200673 001</u>	Jul 08, 2011
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A200673 002</u>	Jul 08, 2011
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200300 001</u>	Jan 26, 2011
<u>AP</u>	!	<u>5MG/10ML (0.5MG/ML)</u>	<u>A200300 002</u>	Jan 26, 2011
<u>AP</u>	XGEN PHARMS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A203167 001</u>	Oct 29, 2013
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A203167 002</u>	Oct 29, 2013
<u>AP</u>	ZYDUS PHARMS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A202601 001</u>	Feb 20, 2014
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A202601 002</u>	Feb 20, 2014

DURACLONAP + MYLAN INSTITUTIONAL1MG/10ML (0.1MG/ML)N020615 001 Oct 02, 1996

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A070974 001</u>	Dec 16, 1986
<u>AB</u>		<u>0.2MG</u>	<u>A070975 001</u>	Dec 16, 1986
<u>AB</u>		<u>0.3MG</u>	<u>A070976 001</u>	Dec 16, 1986
<u>AB</u>	ALEMBIC PHARMS LTD	<u>0.1MG</u>	<u>A091368 001</u>	Dec 06, 2011
<u>AB</u>		<u>0.2MG</u>	<u>A091368 002</u>	Dec 06, 2011
<u>AB</u>		<u>0.3MG</u>	<u>A091368 003</u>	Dec 06, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.1MG</u>	<u>A070886 002</u>	Aug 31, 1988
<u>AB</u>		<u>0.2MG</u>	<u>A070886 001</u>	Aug 31, 1988
<u>AB</u>		<u>0.3MG</u>	<u>A070886 003</u>	Aug 31, 1988
<u>AB</u>	CHARTWELL MOLECULES	<u>0.1MG</u>	<u>A071785 002</u>	Apr 05, 1988
<u>AB</u>		<u>0.2MG</u>	<u>A071785 003</u>	Apr 05, 1988
<u>AB</u>		<u>0.3MG</u>	<u>A071785 001</u>	Apr 05, 1988
<u>AB</u>	IMPAX LABS	<u>0.1MG</u>	<u>A078099 001</u>	Aug 27, 2009
<u>AB</u>		<u>0.2MG</u>	<u>A078099 002</u>	Aug 27, 2009
<u>AB</u>		<u>0.3MG</u>	<u>A078099 003</u>	Aug 27, 2009
<u>AB</u>	PRINSTON INC	<u>0.1MG</u>	<u>A077901 001</u>	Mar 09, 2007
<u>AB</u>		<u>0.2MG</u>	<u>A077901 002</u>	Mar 09, 2007
<u>AB</u>		<u>0.3MG</u>	<u>A077901 003</u>	Mar 09, 2007
<u>AB</u>	UNICHEM	<u>0.1MG</u>	<u>A078895 001</u>	Aug 26, 2009
<u>AB</u>		<u>0.2MG</u>	<u>A078895 002</u>	Aug 26, 2009
<u>AB</u>	!	<u>0.3MG</u>	<u>A078895 003</u>	Aug 26, 2009
<u>AB</u>	URL LABS	<u>0.1MG</u>	<u>A070923 003</u>	Sep 04, 1987
<u>AB</u>		<u>0.2MG</u>	<u>A070923 002</u>	Sep 04, 1987
<u>AB</u>		<u>0.3MG</u>	<u>A070923 001</u>	Sep 04, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>0.1MG</u>	<u>A202297 001</u>	Jun 13, 2013
<u>AB</u>		<u>0.2MG</u>	<u>A202297 002</u>	Jun 13, 2013
<u>AB</u>		<u>0.3MG</u>	<u>A202297 003</u>	Jun 13, 2013

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

<u>AB1</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A203320 001</u>	May 15, 2015
<u>AB1</u>	!	<u>0.1MG</u>	<u>A209686 001</u>	Nov 20, 2017
<u>AB1</u>	ANCHEN PHARMS	<u>0.1MG</u>	<u>A202984 001</u>	Sep 30, 2013
<u>AB1</u>	JUBILANT GENERICS	<u>0.1MG</u>	<u>A210338 001</u>	Jan 29, 2018
<u>AB1</u>	LUPIN LTD	<u>0.1MG</u>	<u>A209285 001</u>	Oct 23, 2017
<u>AB1</u>	NOVAST LABS	<u>0.1MG</u>	<u>A209675 001</u>	Mar 05, 2019
<u>AB1</u>	XIAMEN LP PHARM CO	<u>0.1MG</u>	<u>A209757 001</u>	Nov 20, 2017

CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 75MG BASE</u>	<u>A202925 001</u>	Mar 27, 2013
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202925 002</u>	Mar 27, 2013
<u>AB</u>	ACME LABS	<u>EQ 75MG BASE</u>	<u>A078004 001</u>	May 17, 2012
<u>AB</u>	ALKEM LABS LTD	<u>EQ 75MG BASE</u>	<u>A203632 001</u>	Dec 08, 2023
<u>AB</u>	AMNEAL PHARMS	<u>EQ 75MG BASE</u>	<u>A203751 001</u>	Apr 11, 2014
<u>AB</u>	APOTEX INC	<u>EQ 75MG BASE</u>	<u>A076274 001</u>	May 17, 2012
<u>AB</u>	!	<u>EQ 300MG BASE</u>	<u>A076274 002</u>	Mar 04, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 75MG BASE</u>	<u>A090540 001</u>	May 17, 2012
<u>AB</u>	DR REDDYS	<u>EQ 75MG BASE</u>	<u>A076273 001</u>	Jan 14, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 300MG BASE</u>	<u>A091023 001</u>	May 17, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 75MG BASE</u>	<u>A205345 001</u>	Aug 04, 2023
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A205345 002</u>	Aug 04, 2023

PRESCRIPTION DRUG PRODUCT LIST

CLOPIDOGREL BISULFATE

TABLET; ORAL

CLOPIDOGREL BISULFATE

AB	MACLEODS PHARMS LTD	EQ 75MG BASE	A202928 001	Feb 10, 2014
AB	POLYGEN PHARMS	EQ 75MG BASE	A213351 001	Jul 17, 2020
AB		EQ 300MG BASE	A213351 002	Jul 17, 2020
AB	PRINSTON INC	EQ 75MG BASE	A206376 001	May 07, 2018
AB		EQ 300MG BASE	A206376 002	May 07, 2018
AB	RISING	EQ 75MG BASE	A204359 001	Feb 02, 2017
AB	SCIEGEN PHARMS INC	EQ 75MG BASE	A204165 001	Sep 15, 2014
AB		EQ 300MG BASE	A204165 002	Sep 15, 2014
AB	TEVA	EQ 75MG BASE	A076999 001	May 17, 2012
AB	TORRENT PHARMS LTD	EQ 75MG BASE	A090844 001	May 17, 2012
PLAVIX				
AB	+ SANOFI AVENTIS US	EQ 75MG BASE	N020839 001	Nov 17, 1997
AB	+	EQ 300MG BASE	N020839 002	Sep 20, 2007

CLORAZEPATE DIPOTASSIUM

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

AB	AUROBINDO PHARMA	3.75MG	A071858 002	Jul 17, 1987
AB		7.5MG	A071858 003	Jul 17, 1987
AB	!	15MG	A071858 001	Jul 17, 1987
AB	COREPHARMA	3.75MG	A215566 001	Jun 14, 2022
AB		7.5MG	A215566 002	Jun 14, 2022
AB		15MG	A215566 003	Jun 14, 2022
AB	NOVITIUM PHARMA	3.75MG	A213730 001	Jun 16, 2022
AB		7.5MG	A213730 002	Jun 16, 2022
AB		15MG	A213730 003	Dec 16, 2022
AB	TARO	3.75MG	A075731 003	Apr 27, 2000
AB		7.5MG	A075731 002	Apr 27, 2000
AB		15MG	A075731 001	Apr 27, 2000
TRANXENE				
AB	+ AJENAT PHARMS	7.5MG	N017105 007	

CLOTRIMAZOLE

CREAM; TOPICAL

CLOTRIMAZOLE

AB	FOUGERA PHARMS	1%	A078338 001	Sep 02, 2008
AB	GLENMARK PHARMS	1%	A090219 001	Aug 03, 2010
AB	! TARO	1%	A072640 001	Aug 31, 1993

SOLUTION; TOPICAL

CLOTRIMAZOLE

AT	NOVITIUM PHARMA	1%	A209815 001	Feb 14, 2019
AT	SCIEGEN PHARMS INC	1%	A216569 001	Oct 16, 2023
AT	! TARO	1%	A074580 001	Jul 29, 1996
AT	TASMAN PHARMA	1%	A212281 001	Jul 25, 2019
AT	TEVA	1%	A073306 001	Feb 28, 1995

TROCHE/LOZENGE; ORAL

CLOTRIMAZOLE

AB	! HIKMA	10MG	A076387 001	Jul 29, 2004
AB	PADAGIS US	10MG	A076763 001	Oct 28, 2005

CLOZAPINE

SUSPENSION; ORAL

VERSACLOZ

+! TASMAN PHARMA

50MG/ML

N203479 001 Feb 06, 2013

TABLET; ORAL

CLOZAPINE

AB	ACCORD HLTHCARE	25MG	A202873 001	Nov 25, 2015
AB		100MG	A202873 002	Nov 25, 2015
AB	AUROBINDO PHARMA	25MG	A206433 001	Nov 29, 2016
AB		50MG	A206433 002	Nov 29, 2016
AB		100MG	A206433 003	Nov 29, 2016
AB		200MG	A206433 004	Nov 29, 2016
AB	IVAX SUB TEVA PHARMS	25MG	A074949 001	Nov 26, 1997
AB		50MG	A074949 004	Apr 25, 2005
AB		50MG	A076809 003	Dec 16, 2005
AB		100MG	A074949 002	Nov 26, 1997
AB		100MG	A076809 002	Dec 16, 2005
AB		200MG	A076809 001	Dec 16, 2005
AB	MYLAN	25MG	A075417 001	May 27, 1999
AB		50MG	A075417 004	Apr 15, 2010

PRESCRIPTION DRUG PRODUCT LIST

CLOZAPINE

TABLET; ORAL

CLOZAPINE

AB		<u>100MG</u>	<u>A075417 002</u>	May 27, 1999
AB		<u>200MG</u>	<u>A075417 005</u>	Apr 15, 2010
AB	SUN PHARM INDS INC	<u>25MG</u>	<u>A075713 001</u>	Nov 15, 2002
AB		<u>50MG</u>	<u>A075713 003</u>	Aug 19, 2005
AB		<u>100MG</u>	<u>A075713 002</u>	Nov 15, 2002
AB		<u>200MG</u>	<u>A075713 004</u>	Nov 07, 2017

CLOZARIL

AB	+	HERITAGE LIFE	<u>25MG</u>	<u>N019758 001</u>	Sep 26, 1989
AB	+		<u>50MG</u>	<u>N019758 003</u>	May 20, 2019
AB	+	!	<u>100MG</u>	<u>N019758 002</u>	Sep 26, 1989
AB	+		<u>200MG</u>	<u>N019758 004</u>	May 20, 2019

CLOZAPINE

IVAX SUB TEVA
PHARMS

A074949 003 Jul 31, 2003

TABLET, ORALLY DISINTEGRATING; ORAL

CLOZAPINE

AB		BARR LABS INC	<u>25MG</u>	<u>A090308 001</u>	Nov 25, 2015
AB	!		<u>100MG</u>	<u>A090308 002</u>	Nov 25, 2015
AB		MYLAN	<u>25MG</u>	<u>A201824 002</u>	Sep 15, 2015
AB			<u>100MG</u>	<u>A201824 003</u>	Sep 15, 2015
AB			<u>150MG</u>	<u>A201824 004</u>	Aug 07, 2023
AB			<u>200MG</u>	<u>A201824 005</u>	Aug 07, 2023
AB		TEVA PHARMS USA	<u>150MG</u>	<u>A203039 001</u>	Nov 25, 2015
AB			<u>200MG</u>	<u>A203039 002</u>	Nov 25, 2015
		BARR LABS INC	12.5MG	A090308 003	Apr 09, 2018

COBICISTAT

TABLET; ORAL

TYBOST

+	!	GILEAD SCIENCES INC	150MG	N203094 001	Sep 24, 2014
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COBICISTAT; DARUNAVIR

TABLET; ORAL

PREZCOBIX

+	!	JANSSEN PRODS	150MG; 800MG	N205395 001	Jan 29, 2015
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COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

SYM TUZA

+	!	JANSSEN PRODS	150MG; 800MG; 200MG; EQ 10MG BASE	N210455 001	Jul 17, 2018
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COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

GENVOYA

+	!	GILEAD SCIENCES INC	150MG; 150MG; 200MG; EQ 10MG BASE	N207561 001	Nov 05, 2015
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COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

STRIBILD

+	!	GILEAD SCIENCES INC	150MG; 150MG; 200MG; 300MG	N203100 001	Aug 27, 2012
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COBIMETINIB FUMARATE

TABLET; ORAL

COTELLIC

+	!	GENENTECH INC	EQ 20MG BASE	N206192 001	Nov 10, 2015
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COCAINE HYDROCHLORIDE

SOLUTION; NASAL

GOPRELTO

+	!	GENUS LIFESCIENCES	4%	N209963 001	Dec 14, 2017
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NUMBRINO

+	!	OMNIVIUM PHARMS	4%	N209575 001	Jan 10, 2020
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CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA	!	PHARM ASSOC	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>A040660 001</u>	Dec 07, 2006
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PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA		HIKMA	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>A040674 001</u>	Dec 23, 2014
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PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA		AMNEAL PHARMS	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>A200963 001</u>	Aug 26, 2015
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PRESCRIPTION DRUG PRODUCT LIST

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

AA	AMNEAL PHARMS	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A200894</u> <u>001</u>	Apr 24, 2013
AA	HIKMA	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A040151</u> <u>001</u>	Aug 26, 1997
AA	NOSTRUM LABS INC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A090180</u> <u>001</u>	Mar 17, 2010
AA	QUAGEN	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A214238</u> <u>001</u>	Oct 08, 2020
AA	! TRIS PHARMA INC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A200386</u> <u>001</u>	Jun 29, 2012
AA	WOCKHARDT BIO AG	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A088875</u> <u>001</u>	Dec 17, 1984
<u>PROMETHAZINE WITH CODEINE</u>				
AA	PHARM ASSOC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A040650</u> <u>001</u>	Jan 31, 2006

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

TRIACIN-C

! ANIMA

10MG/5ML; 30MG/5ML; 1.25MG/5ML

A088704 001 Mar 22, 1985

CODEINE SULFATE

TABLET; ORAL

CODEINE SULFATE

AB	+	HIKMA	<u>15MG</u>	<u>N022402</u> <u>001</u>	Jul 16, 2009
AB	+		<u>30MG</u>	<u>N022402</u> <u>002</u>	Jul 16, 2009
AB	+	!	<u>60MG</u>	<u>N022402</u> <u>003</u>	Jul 16, 2009
AB		LANNETT CO INC	<u>15MG</u>	<u>A203046</u> <u>001</u>	Jun 13, 2014
AB			<u>30MG</u>	<u>A203046</u> <u>002</u>	Jun 13, 2014
AB			<u>60MG</u>	<u>A203046</u> <u>003</u>	Jun 13, 2014

COLCHICINE

CAPSULE; ORAL

COLCHICINE

AB		PAR PHARM INC	<u>0.6MG</u>	<u>A208678</u> <u>001</u>	Nov 29, 2018
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MITIGARE

AB	+	!	HIKMA INTL PHARMS	<u>0.6MG</u>	<u>N204820</u> <u>001</u>	Sep 26, 2014
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SOLUTION; ORAL

GLOPERBA

+! SCILEX PHARMS

0.6MG/5ML

N210942 001 Jan 30, 2019

TABLET; ORAL

COLCHICINE

AB		ALKEM LABS LTD	<u>0.6MG</u>	<u>A211250</u> <u>001</u>	Feb 08, 2019
AB		AMNEAL PHARMS	<u>0.6MG</u>	<u>A204711</u> <u>001</u>	Sep 28, 2016
AB		AUROBINDO PHARMA LTD	<u>0.6MG</u>	<u>A215444</u> <u>001</u>	Jan 06, 2022
AB		DR REDDYS	<u>0.6MG</u>	<u>A209876</u> <u>001</u>	Sep 06, 2019
AB		GRANULES	<u>0.6MG</u>	<u>A210425</u> <u>001</u>	Feb 05, 2020
AB		HETERO LABS LTD V	<u>0.6MG</u>	<u>A208993</u> <u>001</u>	Aug 13, 2021
AB		MYLAN	<u>0.6MG</u>	<u>A209470</u> <u>001</u>	Sep 16, 2019
AB		PAR PHARM INC	<u>0.6MG</u>	<u>A203976</u> <u>001</u>	Aug 12, 2021
AB		STRIDES PHARMA	<u>0.6MG</u>	<u>A209173</u> <u>001</u>	Mar 10, 2022
AB		WATSON LABS INC	<u>0.6MG</u>	<u>A204461</u> <u>001</u>	Jul 31, 2019
AB		ZYDUS PHARMS	<u>0.6MG</u>	<u>A211519</u> <u>001</u>	Feb 19, 2019

COLCRYS

AB	+	!	TAKEDA PHARMS USA	<u>0.6MG</u>	<u>N022352</u> <u>001</u>	Jul 29, 2009
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COLCHICINE

ZYDUS PHARMS

0.3MG

A211519 002 Nov 14, 2019

LODOCO

+! AGEPHA PHARMA FZ

0.5MG

N215727 001 Jun 16, 2023

COLCHICINE; PROBENECID

TABLET; ORAL

COL-PROBENECID

AB	+	!	WATSON LABS	<u>0.5MG; 500MG</u>	<u>A084279</u> <u>001</u>	
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PROBENECID AND COLCHICINE

AB		NOVAST LABS	<u>0.5MG; 500MG</u>	<u>A040618</u> <u>001</u>	May 13, 2008
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AB		RISING	<u>0.5MG; 500MG</u>	<u>A217030</u> <u>001</u>	Oct 24, 2023
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COLESEVELAM HYDROCHLORIDE

FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

AB		ALKEM LABS LTD	<u>3.75GM/PACKET</u>	<u>A210316</u> <u>001</u>	May 06, 2019
AB		GLENMARK PHARMS LTD	<u>3.75GM/PACKET</u>	<u>A202190</u> <u>002</u>	Jul 16, 2018
AB		IMPAX	<u>3.75GM/PACKET</u>	<u>A212886</u> <u>001</u>	Jan 18, 2023

WELCHOL

AB	+	!	COSETTE	<u>3.75GM/PACKET</u>	<u>N022362</u> <u>002</u>	Oct 02, 2009
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COLESEVELAM HYDROCHLORIDE

GLENMARK PHARMS LTD

1.875GM/PACKET

A202190 001 Jul 16, 2018

PRESCRIPTION DRUG PRODUCT LIST

COLESEVELAM HYDROCHLORIDE

TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

AB	ALKEM LABS LTD	625MG	A209038 001	Oct 05, 2018
AB	BEIJING TIDE PHARM	625MG	A206036 001	Oct 14, 2021
AB	BIONPHARMA	625MG	A208670 001	Sep 13, 2019
AB	DR REDDYS	625MG	A210889 001	Oct 05, 2018
AB	GLENMARK PHARMS LTD	625MG	A203480 001	May 18, 2018
AB	IMPAX LABS INC	625MG	A091600 001	May 16, 2018
AB	INVENTIA	625MG	A212050 001	Dec 04, 2020
AB	ZHEJIANG JINGXIN	625MG	A209946 001	Jul 15, 2020
AB	ZYDUS PHARMS	625MG	A207765 001	Oct 07, 2019

WELCHOL

AB	+ COSETTE	625MG	N021176 001	May 26, 2000
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COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL

COLESTID

AB	+ PFIZER	5GM/SCOOPFUL	N017563 003	Sep 22, 1995
AB	+ !	5GM/PACKET	N017563 004	Sep 22, 1995

COLESTIPOL HYDROCHLORIDE

AB	IMPAX LABS	5GM/SCOOPFUL	A077277 001	May 02, 2006
AB		5GM/PACKET	A077277 002	May 02, 2006

FLAVORED COLESTID

+ PFIZER

5GM/PACKET

N017563 001

+

5GM/SCOOPFUL

N017563 002

TABLET; ORAL

COLESTID

AB	+ PFIZER	1GM	N020222 001	Jul 19, 1994
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COLESTIPOL HYDROCHLORIDE

AB	ANI PHARMS	1GM	A216517 001	Mar 10, 2023
AB	! IMPAX LABS	1GM	A077510 001	Oct 24, 2006
AB	ZYDUS PHARMS	1GM	A215223 001	Mar 10, 2022

COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

COLISTIMETHATE SODIUM

AP	AVET LIFESCIENCES	EQ 150MG BASE/VIAL	A202359 001	Sep 28, 2012
AP	FRESENIUS KABI USA	EQ 150MG BASE/VIAL	A065364 001	Apr 17, 2008
AP	NEXUS	EQ 150MG BASE/VIAL	A065177 001	Mar 19, 2004
AP	SAGENT PHARMS INC	EQ 150MG BASE/VIAL	A201365 001	Feb 19, 2014
AP	XELLIA PHARMS APS	EQ 150MG BASE/VIAL	A205356 001	May 29, 2015
AP	XGEN PHARMS	EQ 150MG BASE/VIAL	A064216 001	Feb 26, 1999

COLY-MYCIN M

AP	+ ! PAR STERILE PRODUCTS	EQ 150MG BASE/VIAL	N050108 002	
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COLISTIN SULFATE; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; THONZONIUM BROMIDE

SUSPENSION/DROPS; OTIC

COLY-MYCIN S

+! ENDO PHARMS INC

EQ 3MG BASE/ML; 10MG/ML; EQ 3.3MG
BASE/ML; 0.5MG/ML

N050356 001

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER

+! CUMBERLAND

20MG/100ML (0.2MG/ML)

N021697 002 Oct 08, 2008

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

PARAGARD T 380A

+! COOPERSURGICAL

309MG/COPPER

N018680 001 Nov 15, 1984

COPPER CU-64 DOTATATE

SOLUTION; INTRAVENOUS

DETECTNET

+! RADIOMEDIX

4mL (1mCi/ML)

N213227 001 Sep 03, 2020

CORTICOTROPIN

INJECTABLE; INJECTION

ACTHAR GEL

+! MALLINCKRODT ARD

80 UNITS/ML

N008372 008

PURIFIED CORTROPHIN GEL

+! ANI PHARMS

80 UNITS/ML

N008975 002

PRESCRIPTION DRUG PRODUCT LIST

COSYNTROPIN

INJECTABLE; INJECTION

CORTROSYN

AP +! AMPHASTAR PHARMS INC **0.25MG/VIAL** **N016750 001**

COSYNTROPIN

AP MYLAN INSTITUTIONAL **0.25MG/VIAL** **A090574 001** Dec 17, 2009
AP SANDOZ **0.25MG/VIAL** **A202147 001** Jun 29, 2012

CRISABOROLE

OINTMENT; TOPICAL

EUCRISA

+! ANACOR PHARMS INC 2% N207695 001 Dec 14, 2016

CRIZOTINIB

CAPSULE; ORAL

XALKORI

+ PF PRISM CV 200MG N202570 001 Aug 26, 2011

+! 250MG N202570 002 Aug 26, 2011

CAPSULE, PELLETS; ORAL

XALKORI

+ PF PRISM CV 20MG N217581 001 Sep 07, 2023

+ 50MG N217581 002 Sep 07, 2023

+! 150MG N217581 003 Sep 07, 2023

CROFELEMER

TABLET, DELAYED RELEASE; ORAL

MYTESI

+! NAPO PHARMS INC 125MG N202292 001 Dec 31, 2012

CROMOLYN SODIUM

CONCENTRATE; ORAL

CROMOLYN SODIUM

AA AILEX PHARMS LLC **100MG/5ML** **A209264 001** Oct 16, 2017

AA MICRO LABS LTD **100MG/5ML** **A202745 001** Apr 04, 2013

AA INDIA **100MG/5ML** **A202583 001** Oct 27, 2011

AA RISING **100MG/5ML** **A202583 001** Oct 27, 2011

GASTROCROM

AA +! MYLAN SPECIALITY LP **100MG/5ML** **N020479 001** Feb 29, 1996

SOLUTION; INHALATION

CROMOLYN SODIUM

AN AILEX PHARMS LLC **10MG/ML** **A209453 001** Oct 16, 2017

AN MICRO LABS **10MG/ML** **A213658 001** Apr 29, 2022

AN ! TEVA PHARMS **10MG/ML** **A075271 001** Jan 18, 2000

AN VIRTUS **10MG/ML** **A075437 001** Apr 21, 2000

AN WOCKHARDT BIO AG **10MG/ML** **A075346 001** Oct 25, 1999

SOLUTION/DROPS; OPHTHALMIC

CROMOLYN SODIUM

! SANDOZ 4% A075282 001 Jun 16, 1999

CROTAMITON

LOTION; TOPICAL

CROTAN

AT LEGACY PHARMA **10%** **A087204 001**

EURAX

AT +! JOURNEY **10%** **N009112 003**

CUPRIC CHLORIDE

INJECTABLE; INJECTION

CUPRIC CHLORIDE

AP EXELA PHARMA **EQ 0.4MG COPPER/ML** **A212071 001** Oct 31, 2022

CUPRIC CHLORIDE IN PLASTIC CONTAINER

AP +! HOSPIRA **EQ 0.4MG COPPER/ML** **N018960 001** Jun 26, 1986

CUPRIC SULFATE

INJECTABLE; INJECTION

CUPRIC SULFATE

! AM REGENT EQ 0.4MG COPPER/ML A216324 001 Dec 16, 2022

CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE

SOLUTION; INTRAVENOUS

MULTRYIS

+! AM REGENT EQ 60MCG COPPER/ML; EQ 3MCG BASE/ML; EQ 6MCG SELENIUM/ML; EQ 1000MCG BASE/ML (1ML) N209376 003 Jun 30, 2021

TRALEMENT

+! AM REGENT EQ 0.3MG COPPER/ML; EQ 55MCG BASE/ML; EQ N209376 001 Jul 02, 2020

PRESCRIPTION DRUG PRODUCT LIST

CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE

SOLUTION; INTRAVENOUS
TRALEMENT

60MCG SELENIUM/ML;EQ 3MG BASE/ML (1ML)			
+!	EQ 0.3MG COPPER/ML;EQ 55MCG BASE/ML;EQ	N209376 002	Dec 02, 2020
	60MCG SELENIUM/ML;EQ 3MG BASE/ML (5ML)		

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN

<u>AP</u>	+!	AM REGENT	<u>1MG/ML</u>	<u>A080737 001</u>	
<u>AP</u>		EUGIA PHARMA	<u>1MG/ML</u>	<u>A213874 001</u>	Dec 08, 2020
<u>AP</u>		GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A214390 001</u>	Sep 24, 2020
<u>AP</u>		MYLAN LABS LTD	<u>1MG/ML</u>	<u>A204829 001</u>	Jun 05, 2017
<u>AP</u>		SAGENT PHARMS INC	<u>1MG/ML</u>	<u>A215107 001</u>	Nov 15, 2022
<u>AP</u>		SANDOZ	<u>1MG/ML</u>	<u>A212915 001</u>	Jan 04, 2021
<u>AP</u>		SOLA PHARMS	<u>1MG/ML</u>	<u>A215417 001</u>	Apr 15, 2022
<u>AP</u>		SOMERSET THERAPS LLC	<u>1MG/ML</u>	<u>A206503 001</u>	Dec 11, 2015
<u>AP</u>			<u>1MG/ML</u>	<u>A209429 001</u>	Dec 18, 2018
<u>AP</u>		VITRUVIAS THERAP	<u>1MG/ML</u>	<u>A209255 001</u>	Dec 18, 2018
<u>AP</u>		WEST-WARD PHARMS INT	<u>1MG/ML</u>	<u>A080515 002</u>	
<u>AP</u>		ZYDUS PHARMS	<u>1MG/ML</u>	<u>A214655 001</u>	Apr 15, 2022

DODEX

<u>AP</u>		ACCORD HLTHCARE	<u>1MG/ML</u>	<u>A083022 001</u>	
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VIBISONE

<u>AP</u>	+!	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A080557 003</u>	
		SPRAY, METERED;NASAL			

CYANOCOBALAMIN

<u>AB</u>		LUPIN	<u>0.5MG/SPRAY</u>	<u>A210629 001</u>	Jun 30, 2023
<u>AB</u>		PADAGIS ISRAEL	<u>0.5MG/SPRAY</u>	<u>A212458 001</u>	Sep 09, 2020

NASCOBAL

<u>AB</u>	+!	ENDO PHARMS INC	<u>0.5MG/SPRAY</u>	<u>N021642 001</u>	Jan 31, 2005
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CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

AMRIX

<u>AB</u>	+	TEVA PHARMS INTL	<u>15MG</u>	<u>N021777 001</u>	Feb 01, 2007
<u>AB</u>	+!		<u>30MG</u>	<u>N021777 002</u>	Feb 01, 2007

CYCLOBENZAPRINE HYDROCHLORIDE

<u>AB</u>		TWI PHARMS INC	<u>15MG</u>	<u>A091281 001</u>	Jan 31, 2013
<u>AB</u>			<u>30MG</u>	<u>A091281 002</u>	Jan 31, 2013

TABLET;ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>5MG</u>	<u>A071611 002</u>	Feb 03, 2006
<u>AB</u>			<u>7.5MG</u>	<u>A071611 003</u>	Feb 03, 2006
<u>AB</u>			<u>10MG</u>	<u>A071611 001</u>	May 03, 1989
<u>AB</u>		ANDA REPOSITORY	<u>5MG</u>	<u>A073541 002</u>	Apr 06, 2006
<u>AB</u>			<u>10MG</u>	<u>A073541 001</u>	May 23, 1995
<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A078643 001</u>	Sep 26, 2008
<u>AB</u>			<u>10MG</u>	<u>A078643 002</u>	Sep 26, 2008
<u>AB</u>		CHARTWELL RX	<u>5MG</u>	<u>A078218 002</u>	Jun 19, 2015
<u>AB</u>			<u>7.5MG</u>	<u>A078218 003</u>	Nov 03, 2020
<u>AB</u>			<u>10MG</u>	<u>A078218 001</u>	Apr 18, 2008
<u>AB</u>		INVAGEN PHARMS	<u>5MG</u>	<u>A090478 001</u>	Jul 23, 2010
<u>AB</u>			<u>10MG</u>	<u>A090478 002</u>	Jul 23, 2010
<u>AB</u>		JUBILANT CADISTA	<u>5MG</u>	<u>A077563 001</u>	Apr 19, 2006
<u>AB</u>			<u>7.5MG</u>	<u>A077563 003</u>	Aug 25, 2017
<u>AB</u>			<u>10MG</u>	<u>A077563 002</u>	Apr 19, 2006
<u>AB</u>		KVK TECH	<u>5MG</u>	<u>A078048 001</u>	Feb 28, 2011
<u>AB</u>			<u>10MG</u>	<u>A078048 002</u>	Feb 28, 2011
<u>AB</u>		OXFORD PHARMS	<u>5MG</u>	<u>A077209 002</u>	Feb 03, 2006
<u>AB</u>			<u>10MG</u>	<u>A077209 001</u>	Oct 04, 2005
<u>AB</u>		PRINSTON INC	<u>5MG</u>	<u>A077797 001</u>	Feb 28, 2007
<u>AB</u>			<u>10MG</u>	<u>A077797 002</u>	Feb 28, 2007
<u>AB</u>		RUBICON	<u>5MG</u>	<u>A208170 001</u>	May 31, 2017
<u>AB</u>			<u>7.5MG</u>	<u>A208170 002</u>	May 31, 2017
<u>AB</u>	!		<u>10MG</u>	<u>A208170 003</u>	May 31, 2017
<u>AB</u>		SUN PHARM INDS LTD	<u>5MG</u>	<u>A078722 001</u>	May 12, 2008
<u>AB</u>			<u>7.5MG</u>	<u>A078722 002</u>	May 12, 2008
<u>AB</u>			<u>10MG</u>	<u>A078722 003</u>	May 12, 2008
<u>AB</u>		UNICHEM	<u>5MG</u>	<u>A213324 001</u>	Jul 06, 2020
<u>AB</u>			<u>7.5MG</u>	<u>A213324 002</u>	Jul 06, 2020

PRESCRIPTION DRUG PRODUCT LIST

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

AB 10MG **A213324 003** Jul 05, 2020

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPENTOLATE

AT RISING **1%** **A040164 001** Jan 13, 1997

AT **2%** **A040165 001** Jan 13, 1997

CYCLOGYL

AT +! ALCON LABS INC **1%** **A084110 001**

AT +! **2%** **A084108 001**

PENTOLAIR

AT BAUSCH AND LOMB **1%** **A040075 001** Apr 29, 1994

CYCLOGYL

+! ALCON LABS INC 0.5% A084109 001

CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYCLOMYDRIL

! ALCON LABS INC 0.2%; 1% A084300 001

CYCLOPHOSPHAMIDE

CAPSULE; ORAL

CYCLOPHOSPHAMIDE

AB ALEMBIC **25MG** **A215892 001** Nov 10, 2022

AB **50MG** **A215892 002** Nov 10, 2022

AB ANIMA **25MG** **A209872 001** May 07, 2018

AB **50MG** **A209872 002** May 07, 2018

AB CIPLA **25MG** **A211608 001** Jan 18, 2019

AB **50MG** **A211608 002** Jan 18, 2019

AB + HIKMA **25MG** **N203856 001** Sep 16, 2013

AB +! **50MG** **N203856 002** Sep 16, 2013

AB ZYDUS LIFESCIENCES **25MG** **A211552 001** Dec 13, 2023

AB **50MG** **A211552 002** Dec 13, 2023

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

AP AMNEAL **500MG/VIAL** **A210046 001** May 25, 2018

AP **1GM/VIAL** **A210046 002** May 25, 2018

AP **2GM/VIAL** **A210046 003** May 25, 2018

AP ! BAXTER HLTHCARE **500MG/VIAL** **A040745 001** May 21, 2008

AP ! **1GM/VIAL** **A040745 002** May 21, 2008

AP ! **2GM/VIAL** **A040745 003** May 21, 2008

AP HENGRUI PHARMA **500MG/VIAL** **A204555 001** Oct 31, 2014

AP **1GM/VIAL** **A204555 002** Oct 31, 2014

AP **2GM/VIAL** **A204555 003** Oct 31, 2014

AP RK PHARMA **500MG/VIAL** **A216958 001** Dec 18, 2023

AP **1GM/VIAL** **A216958 002** Dec 18, 2023

AP **2GM/VIAL** **A216958 003** Dec 18, 2023

AP SAGENT PHARMS INC **500MG/VIAL** **A214529 001** Jul 17, 2023

AP **1GM/VIAL** **A214529 002** Jul 17, 2023

AP **2GM/VIAL** **A214529 003** Jul 17, 2023

AP SUNNY **500MG/VIAL** **A215089 001** Oct 26, 2023

AP **1GM/VIAL** **A215089 002** Oct 26, 2023

AP **2GM/VIAL** **A215089 003** Oct 26, 2023

AP XGEN PHARMS **500MG/VIAL** **A211757 001** Oct 18, 2022

AP **1GM/VIAL** **A211757 002** Oct 18, 2022

AP **2GM/VIAL** **A211757 003** Oct 18, 2022

SOLUTION; INTRAVENOUS

CYCLOPHOSPHAMIDE

+! DR REDDYS 500MG/ML (500MG/ML) N210852 001 Jun 07, 2023

+! 1GM/2ML (500MG/ML) N210852 002 Jun 07, 2023

+! 2GM/4ML (500MG/ML) N210852 003 Jun 07, 2023

+! EUGIA PHARMA 500MG/2.5ML (200MG/ML) N210735 001 Aug 25, 2021

+! SPECLTS 1GM/5ML (200MG/ML) N210735 002 Aug 25, 2021

+! 2GM/10ML (200MG/ML) N210735 003 Nov 20, 2023

+! INGENUS PHARMS LLC 500MG/2.5ML (200MG/ML) N212501 001 Jul 30, 2020

+! 1GM/5ML (200MG/ML) N212501 002 Jul 30, 2020

+! 2GM/10ML (200MG/ML) N212501 003 Nov 19, 2021

+! SANDOZ 500MG/5ML (100MG/ML) N217150 001 Sep 12, 2023

+! 1GM/10ML (100MG/ML) N217150 002 Sep 12, 2023

+! 2GM/20ML (100MG/ML) N217150 003 Sep 12, 2023

PRESCRIPTION DRUG PRODUCT LIST

CYCLOPHOSPHAMIDE

TABLET; ORAL

CYTOXAN

+ BAXTER HLTHCARE
+!

25MG
50MG

N012141 002
N012141 001

CYCLOSERINE

CAPSULE; ORAL

SEROMYCIN

! CEROVENE INC

250MG

A060593 001

CYCLOSPORINE

CAPSULE; ORAL

CYCLOSPORINE

AB1	APOTEX	25MG	A210721 001	Jul 10, 2019
AB1		50MG	A210721 002	Jul 10, 2019
AB1		100MG	A210721 003	Jul 10, 2019
AB1	DR REDDYS LABS SA	25MG	A065044 002	Dec 20, 2000
AB1		100MG	A065044 001	Dec 20, 2000
AB1	IVAX SUB TEVA PHARMS	25MG	A065110 003	Mar 29, 2005
AB1		50MG	A065110 001	Mar 29, 2005
AB1		100MG	A065110 002	Mar 29, 2005
AB1	SANDOZ	25MG	A065017 002	Jan 13, 2000
AB1		100MG	A065017 001	Jan 13, 2000
AB1	STRIDES PHARMA	25MG	A216046 001	Aug 02, 2022
AB1		50MG	A216046 002	Aug 02, 2022
AB1		100MG	A216046 003	Aug 02, 2022
	GENGRAF			
AB1	ABEVIE	25MG	A065003 001	May 12, 2000
AB1		100MG	A065003 003	May 12, 2000
	NEORAL			
AB1	+ NOVARTIS	25MG	N050715 001	Jul 14, 1995
AB1	+!	100MG	N050715 002	Jul 14, 1995
	CYCLOSPORINE			
AB2	APOTEX	25MG	A065040 001	May 09, 2002
AB2		100MG	A065040 002	May 09, 2002
	SANDIMMUNE			
AB2	+ NOVARTIS	25MG	N050625 001	Mar 02, 1990
AB2	+!	100MG	N050625 002	Mar 02, 1990
BX	+	50MG	N050625 003	Nov 23, 1992
	EMULSION; OPHTHALMIC			
	CYCLOSPORINE			
AB	APOTEX	0.05%	A207606 001	Jan 12, 2023
AB	MYLAN	0.05%	A205894 001	Feb 02, 2022
AB	TEVA PHARMS USA INC	0.05%	A203880 001	Dec 14, 2023
	RESTASIS			
AB	+! ABBEVIE	0.05%	N050790 001	Dec 23, 2002
	RESTASIS MULTIDOSE			
	+! ABBEVIE	0.05%	N050790 002	Oct 27, 2016
	VERKAZIA			
	+! HARROW EYE	0.1%	N214965 001	Jun 23, 2021
	INJECTABLE; INJECTION			
	CYCLOSPORINE			
AP	HIKMA	50MG/ML	A065004 001	Oct 29, 1999
AP	PADAGIS US	50MG/ML	A065151 001	Oct 07, 2003
	SANDIMMUNE			
AP	+! NOVARTIS	50MG/ML	N050573 001	Nov 14, 1983
	SOLUTION; OPHTHALMIC			
	CEQUA			
	+! SUN PHARM	0.09%	N210913 001	Aug 14, 2018
	VEVYE			
	+! HARROW EYE	0.1%	N217469 001	May 30, 2023
	SOLUTION; ORAL			
	CYCLOSPORINE			
AB1	ABEVIE	100MG/ML	A065025 001	Mar 03, 2000
AB1	IVAX SUB TEVA PHARMS	100MG/ML	A065078 001	Mar 25, 2005
	NEORAL			
AB1	+! NOVARTIS	100MG/ML	N050716 001	Jul 14, 1995
	SANDIMMUNE			
	+! NOVARTIS	100MG/ML	N050574 001	Nov 14, 1983

PRESCRIPTION DRUG PRODUCT LIST

CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>	CHARTWELL MOLECULAR	<u>2MG/5ML</u>	<u>A203191</u>	<u>001</u>	Jul 13, 2017
<u>AA</u>	ELYSIUM	<u>2MG/5ML</u>	<u>A209108</u>	<u>001</u>	Oct 16, 2018
<u>AA</u>	! LYNE	<u>2MG/5ML</u>	<u>A040668</u>	<u>001</u>	Jun 28, 2006
<u>AA</u>	PATRIN	<u>2MG/5ML</u>	<u>A204823</u>	<u>001</u>	Dec 27, 2016
<u>AA</u>	PHARM ASSOC	<u>2MG/5ML</u>	<u>A091295</u>	<u>001</u>	Mar 28, 2013
<u>AA</u>	QUAGEN	<u>2MG/5ML</u>	<u>A212423</u>	<u>001</u>	May 22, 2019

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>	APPCO	<u>4MG</u>	<u>A206553</u>	<u>001</u>	Nov 29, 2016
<u>AA</u>	BEXIMCO PHARMS USA	<u>4MG</u>	<u>A206676</u>	<u>001</u>	Apr 12, 2019
<u>AA</u>	CHARTWELL RX	<u>4MG</u>	<u>A088212</u>	<u>001</u>	May 26, 1983
<u>AA</u>	KENTON	<u>4MG</u>	<u>A040644</u>	<u>001</u>	May 30, 2006
<u>AA</u>	MOUNTAIN	<u>4MG</u>	<u>A040537</u>	<u>001</u>	Sep 30, 2003
<u>AA</u>	NOVAST LABS	<u>4MG</u>	<u>A205087</u>	<u>001</u>	Sep 23, 2015
<u>AA</u>	QUAGEN	<u>4MG</u>	<u>A212491</u>	<u>001</u>	Feb 24, 2021
<u>AA</u>	RISING	<u>4MG</u>	<u>A207555</u>	<u>001</u>	Jan 31, 2017
<u>AA</u>	STRIDES PHARMA	<u>4MG</u>	<u>A209172</u>	<u>001</u>	Apr 11, 2018
<u>AA</u>	! ZYDUS PHARMS	<u>4MG</u>	<u>A208938</u>	<u>001</u>	May 19, 2017

CYSTEAMINE BITARTRATE

CAPSULE; ORAL

CYSTAGON

+	MYLAN	EQ 50MG BASE	N020392	001	Aug 15, 1994
+	!	EQ 150MG BASE	N020392	002	Aug 15, 1994

CAPSULE, DELAYED RELEASE; ORAL

PROCYSBI

+	HORIZON	EQ 25MG BASE	N203389	001	Apr 30, 2013
+	!	EQ 75MG BASE	N203389	002	Apr 30, 2013

GRANULE, DELAYED RELEASE; ORAL

PROCYSBI

+	HORIZON	EQ 75MG BASE/PACKET	N213491	001	Feb 14, 2020
+	!	EQ 300MG BASE/PACKET	N213491	002	Feb 14, 2020

CYSTEAMINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYSTADROPS

+	! RECORDATI RARE	EQ 0.37% BASE	N211302	001	Aug 19, 2020
+	! LEADIANT BIOSCI INC	EQ 0.44% BASE	N200740	001	Oct 02, 2012

CYSTEINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

ELCYS

+	! EXELA PHARMA	500MG/10ML (50MG/ML)	N210660	001	Apr 16, 2019
+	!	2500MG/50ML (50MG/ML)	N210660	002	Dec 04, 2023

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

<u>AP</u>	! FRESENIUS KABI USA	<u>100MG/ML</u>	<u>A076512</u>	<u>001</u>	Jan 15, 2004
<u>AP</u>	GLAND PHARMA LTD	<u>100MG/VIAL</u>	<u>A211937</u>	<u>001</u>	Dec 23, 2019
<u>AP</u>		<u>2GM/VIAL</u>	<u>A211938</u>	<u>001</u>	Dec 23, 2019
<u>AP</u>	HIKMA	<u>100MG/VIAL</u>	<u>A071471</u>	<u>001</u>	Aug 02, 1989
<u>AP</u>	! HOSPIRA	<u>20MG/ML</u>	<u>A071868</u>	<u>001</u>	Jun 04, 1990
<u>AP</u>	!	<u>20MG/ML</u>	<u>A072168</u>	<u>001</u>	Aug 31, 1990
<u>AP</u>	!	<u>20MG/ML</u>	<u>A072945</u>	<u>001</u>	Feb 28, 1994
<u>AP</u>		<u>100MG/ML</u>	<u>A075383</u>	<u>001</u>	Nov 22, 1999
<u>AP</u>	MEITHEAL	<u>20MG/ML</u>	<u>A208485</u>	<u>001</u>	Feb 28, 2022
<u>AP</u>		<u>100MG/ML</u>	<u>A205696</u>	<u>001</u>	Jul 17, 2018
<u>AP</u>	RISING	<u>20MG/ML</u>	<u>A200915</u>	<u>001</u>	Dec 13, 2011
<u>AP</u>		<u>100MG/ML</u>	<u>A201784</u>	<u>001</u>	Jan 30, 2012
<u>AP</u>	! WEST-WARD PHARMS	<u>2GM/VIAL</u>	<u>A074245</u>	<u>002</u>	Aug 31, 1994
	INT				
!	HIKMA	500MG/VIAL	A071472	001	Aug 02, 1989
!	WEST-WARD PHARMS	1GM/VIAL	A074245	001	Aug 31, 1994
	INT				

PRESCRIPTION DRUG PRODUCT LIST

CYTARABINE; DAUNORUBICIN

POWDER; INTRAVENOUS

VYXEOS

+! CELATOR PHARMS 100MG;44MG

N209401 001 Aug 03, 2017

DABIGATRAN ETEXILATE MESYLATE

CAPSULE; ORAL

DABIGATRAN ETEXILATE MESYLATEAB ALKEM LABS LTD EQ 75MG BASE A208040 001 Mar 11, 2020AB EQ 150MG BASE A208040 002 Mar 11, 2020AB APOTEX EQ 75MG BASE A208070 001 Dec 15, 2023AB EQ 110MG BASE A208070 002 Dec 15, 2023AB EQ 150MG BASE A208070 003 Dec 15, 2023AB HETERO LABS LTD III EQ 75MG BASE A207961 001 May 06, 2020AB EQ 150MG BASE A207961 002 May 06, 2020PRADAXAAB + BOEHRINGER EQ 75MG BASE N022512 001 Oct 19, 2010
INGELHEIMAB + EQ 110MG BASE N022512 003 Nov 20, 2015AB +! EQ 150MG BASE N022512 002 Oct 19, 2010

PELLETS; ORAL

PRADAXA

+ BOEHRINGER EQ 20MG BASE/PACKET N214358 001 Jun 21, 2021
INGELHEIM

+ EQ 30MG BASE/PACKET N214358 002 Jun 21, 2021

+ EQ 40MG BASE/PACKET N214358 003 Jun 21, 2021

+ EQ 50MG BASE/PACKET N214358 004 Jun 21, 2021

+ EQ 110MG BASE/PACKET N214358 005 Jun 21, 2021

+! EQ 150MG BASE/PACKET N214358 006 Jun 21, 2021

DABRAFENIB MESYLATE

CAPSULE; ORAL

TAFINLAR

+ NOVARTIS EQ 50MG BASE N202806 001 May 29, 2013

+! EQ 75MG BASE N202806 002 May 29, 2013

TABLET, FOR SUSPENSION; ORAL

TAFINLAR

+! NOVARTIS EQ 10MG BASE N217514 001 Mar 16, 2023

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINEAP ! FRESENIUS KABI USA 200MG/VIAL A075371 002 Aug 27, 1999AP HIKMA 200MG/VIAL A075812 001 Jun 15, 2001AP 500MG/VIAL A075812 002 Oct 31, 2002AP HOSPIRA 200MG/VIAL A075940 001 Oct 18, 2001AP MEITHEAL 200MG/VIAL A075259 002 Aug 27, 1998AP ! 500MG/VIAL A075259 001 Sep 22, 2000

! FRESENIUS KABI USA 100MG/VIAL A075371 001 Aug 27, 1999

DACOMITINIB

TABLET; ORAL

VIZIMPRO

+ PFIZER 15MG N211288 001 Sep 27, 2018

+ 30MG N211288 002 Sep 27, 2018

+! 45MG N211288 003 Sep 27, 2018

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGENAP +! RECORDATI RARE 0.5MG/VIAL N050682 001DACTINOMYCINAP EUGIA PHARMA 0.5MG/VIAL A203385 001 Nov 09, 2017AP HISUN PHARM 0.5MG/VIAL A207232 001 Jul 16, 2019
HANGZHOUAP MEITHEAL 0.5MG/VIAL A213463 001 Nov 13, 2020AP XGEN PHARMS 0.5MG/VIAL A203999 001 May 20, 2019DALBAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

DALVANCE

+! ABBVIE EQ 500MG BASE/VIAL N021883 001 May 23, 2014

PRESCRIPTION DRUG PRODUCT LIST

DALFAMPRIDINE

TABLET, EXTENDED RELEASE;ORAL

AMPYRA

AB	+ !	ACORDA	10MG	N022250	001	Jan 22, 2010
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DALFAMPRIDINE

AB		ACCORD HLTHCARE	10MG	A206863	001	Jul 11, 2018
AB		ACTAVIS LABS FL INC	10MG	A206836	001	Jan 23, 2017
AB		ALKEM LABS LTD	10MG	A206765	001	Jul 30, 2018
AB		AUROBINDO PHARMA	10MG	A206811	001	Jan 23, 2017
AB		MICRO LABS	10MG	A210158	001	Mar 11, 2019
AB		SUN PHARM	10MG	A208292	001	May 21, 2019

DALFOPRISTIN; QUINUPRISTININJECTABLE; INTRAVENOUS
SYNERCID

+ !	KING PHARMS	350MG/VIAL;150MG/VIAL	N050748	001	Sep 21, 1999
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DALTEPARIN SODIUMINJECTABLE; SUBCUTANEOUS
FRAGMIN

+	PFIZER	2,500IU/0.2ML (12,500IU/ML)	N020287	001	Dec 22, 1994
+		5,000IU/0.2ML (25,000IU/ML)	N020287	003	Mar 18, 1996
+		7,500IU/0.3ML (25,000IU/ML)	N020287	005	Apr 04, 2002
+		10,000IU/ML (10,000IU/ML)	N020287	004	Jan 30, 1998
+		10,000IU/4ML (2,500IU/ML)	N020287	012	Mar 15, 2022
+		12,500IU/0.5ML (25,000IU/ML)	N020287	009	May 01, 2007
+		15,000IU/0.6ML (25,000IU/ML)	N020287	010	May 01, 2007
+		18,000IU/0.72ML (25,000IU/ML)	N020287	011	May 01, 2007
+ !		95,000IU/3.8ML (25,000IU/ML)	N020287	006	Apr 04, 2002

DANAZOL

CAPSULE; ORAL

DANAZOL

AB		BARR	50MG	A074582	003	May 29, 1998
AB			100MG	A074582	002	May 29, 1998
AB	!		200MG	A074582	001	Aug 09, 1996
AB		LANNETT CO INC	50MG	A077246	002	Apr 19, 2007
AB			100MG	A077246	003	Apr 19, 2007
AB			200MG	A077246	001	Sep 28, 2005

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIUM

AB	+	PAR STERILE PRODUCTS	25MG	N017443	001	
AB	+		50MG	N017443	003	
AB	+ !		100MG	N017443	002	

DANTROLENE SODIUM

AB		ELITE LABS INC	25MG	A076686	001	Oct 24, 2005
AB			50MG	A076686	002	Oct 24, 2005
AB			100MG	A076686	003	Oct 24, 2005
AB		IMPAX LABS	25MG	A076856	001	Mar 01, 2005
AB			50MG	A076856	002	Mar 01, 2005
AB			100MG	A076856	003	Mar 01, 2005

FOR SUSPENSION; INTRAVENOUS

RYANODEX

+ !	EAGLE PHARMS	250MG/VIAL	N205579	001	Jul 22, 2014
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INJECTABLE; INJECTION

DANTRIUM

AP	+ !	PAR STERILE PRODUCTS	20MG/VIAL	N018264	001	
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DANTROLENE SODIUM

AP		HIKMA	20MG/VIAL	A204762	001	Jun 19, 2017
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REVONTO

AP		USWM	20MG/VIAL	A078378	001	Jul 24, 2007
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DAPAGLIFLOZIN

TABLET; ORAL

FARXIGA

+	ASTRAZENECA AB	5MG	N202293	001	Jan 08, 2014
+ !		10MG	N202293	002	Jan 08, 2014

PRESCRIPTION DRUG PRODUCT LIST

DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

XIGDUO XR

+	ASTRAZENECA AB	2.5MG;1GM	N205649	005	Jul 28, 2017
+		5MG;500MG	N205649	001	Oct 29, 2014
+		5MG;1GM	N205649	002	Oct 29, 2014
+		10MG;500MG	N205649	003	Oct 29, 2014
+	!	10MG;1GM	N205649	004	Oct 29, 2014

DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE

TABLET;ORAL

QTERN

+	ASTRAZENECA AB	5MG;EQ 5MG BASE	N209091	002	May 02, 2019
+	!	10MG;EQ 5MG BASE	N209091	001	Feb 27, 2017

DAPRODUSTAT

TABLET;ORAL

JESDUVROQ

+	GLAXOSMITHKLINE	1MG	N216951	001	Feb 01, 2023
+		2MG	N216951	002	Feb 01, 2023
+		4MG	N216951	003	Feb 01, 2023
+		6MG	N216951	004	Feb 01, 2023
+	!	8MG	N216951	005	Feb 01, 2023

DAPSONE

GEL;TOPICAL

ACZONE

<u>AB</u>	+	!	ABEVIE	<u>5%</u>	<u>N021794</u>	<u>001</u>	Jul 07, 2005
<u>AB</u>	+	!	ALMIRALL	<u>7.5%</u>	<u>N207154</u>	<u>001</u>	Feb 24, 2016

DAPSONE

<u>AB</u>			ALEMBIC	<u>7.5%</u>	<u>A215718</u>	<u>001</u>	Oct 27, 2023
<u>AB</u>			AMNEAL	<u>5%</u>	<u>A209890</u>	<u>001</u>	Oct 19, 2023
<u>AB</u>				<u>7.5%</u>	<u>A212701</u>	<u>001</u>	May 31, 2023
<u>AB</u>			BELOTECA	<u>5%</u>	<u>A213907</u>	<u>001</u>	Jun 06, 2023
<u>AB</u>			COSETTE	<u>5%</u>	<u>A210178</u>	<u>001</u>	Mar 31, 2022
<u>AB</u>			ENCUBE	<u>5%</u>	<u>A212383</u>	<u>001</u>	Oct 13, 2023
<u>AB</u>			MYLAN	<u>7.5%</u>	<u>A213847</u>	<u>001</u>	Feb 04, 2022
<u>AB</u>			PADAGIS ISRAEL	<u>5%</u>	<u>A215087</u>	<u>001</u>	Oct 31, 2023
<u>AB</u>				<u>7.5%</u>	<u>A212657</u>	<u>001</u>	Nov 01, 2023
<u>AB</u>			TARO	<u>5%</u>	<u>A209506</u>	<u>001</u>	Oct 16, 2017
<u>AB</u>				<u>7.5%</u>	<u>A210191</u>	<u>001</u>	Jun 26, 2019
<u>AB</u>			TORRENT	<u>7.5%</u>	<u>A214722</u>	<u>001</u>	Feb 10, 2022

TABLET;ORAL

DAPSONE

<u>AB</u>			ACTAVIS LLC	<u>25MG</u>	<u>A204380</u>	<u>001</u>	Mar 23, 2017
<u>AB</u>				<u>100MG</u>	<u>A204380</u>	<u>002</u>	Mar 23, 2017
<u>AB</u>	+		EVEREST LIFE SCI	<u>25MG</u>	<u>A086841</u>	<u>001</u>	
<u>AB</u>	+			<u>100MG</u>	<u>A086842</u>	<u>001</u>	
<u>AB</u>			NOSTRUM LABS INC	<u>25MG</u>	<u>A203887</u>	<u>001</u>	May 06, 2016
<u>AB</u>				<u>100MG</u>	<u>A203887</u>	<u>002</u>	May 06, 2016
<u>AB</u>			NOVITIUM PHARMA	<u>25MG</u>	<u>A206505</u>	<u>001</u>	Dec 01, 2016
<u>AB</u>	!			<u>100MG</u>	<u>A206505</u>	<u>002</u>	Dec 01, 2016
<u>AB</u>			RISING	<u>25MG</u>	<u>A207165</u>	<u>002</u>	Jul 20, 2023
<u>AB</u>				<u>100MG</u>	<u>A207165</u>	<u>001</u>	May 08, 2019

DAPTOMYCIN

POWDER; INTRAVENOUS

DAPTOMYCIN

<u>AP</u>			ACCORD HLTHCARE	<u>350MG/VIAL</u>	<u>A212667</u>	<u>001</u>	Jul 12, 2019
<u>AP</u>				<u>500MG/VIAL</u>	<u>A211961</u>	<u>001</u>	Jun 24, 2019
<u>AP</u>			ASPIRO	<u>350MG/VIAL</u>	<u>A216445</u>	<u>001</u>	Dec 23, 2022
<u>AP</u>				<u>500MG/VIAL</u>	<u>A216413</u>	<u>001</u>	Sep 23, 2022
<u>AP</u>			BE PHARMS	<u>350MG/VIAL</u>	<u>A213425</u>	<u>001</u>	Aug 20, 2020
<u>AP</u>				<u>500MG/VIAL</u>	<u>A212513</u>	<u>001</u>	Jun 26, 2019
<u>AP</u>			DR REDDYS	<u>350MG/VIAL</u>	<u>A211403</u>	<u>001</u>	Aug 31, 2020
<u>AP</u>				<u>500MG/VIAL</u>	<u>A208375</u>	<u>001</u>	May 01, 2019
<u>AP</u>			EUGIA PHARMA	<u>500MG/VIAL</u>	<u>A213171</u>	<u>001</u>	Sep 02, 2021
<u>AP</u>			FRESENIUS KABI USA	<u>500MG/VIAL</u>	<u>A206077</u>	<u>001</u>	Apr 11, 2018
<u>AP</u>			HAINAN POLY PHARM	<u>500MG/VIAL</u>	<u>A215890</u>	<u>001</u>	Aug 18, 2022
<u>AP</u>			HANGZHOU ZHONGMEI	<u>500MG/VIAL</u>	<u>A215215</u>	<u>001</u>	Nov 26, 2021
<u>AP</u>			HENGRUI PHARMA	<u>500MG/VIAL</u>	<u>A212022</u>	<u>001</u>	Aug 22, 2019
<u>AP</u>			HISUN PHARM	<u>500MG/VIAL</u>	<u>A212250</u>	<u>001</u>	Apr 21, 2021
<u>AP</u>			HANGZHOU				
<u>AP</u>			MEITHEAL	<u>350MG/VIAL</u>	<u>A213786</u>	<u>001</u>	Jun 29, 2021
<u>AP</u>				<u>500MG/VIAL</u>	<u>A213623</u>	<u>001</u>	Jun 29, 2021

PRESCRIPTION DRUG PRODUCT LIST

DAPTOMYCIN

POWDER; INTRAVENOUS

DAPTOMYCIN

<u>AP</u>	MYLAN	<u>500MG/VIAL</u>	<u>A213966</u>	<u>001</u>	Aug 07, 2023
<u>AP</u>	MYLAN LABS LTD	<u>500MG/VIAL</u>	<u>A205037</u>	<u>001</u>	Jun 05, 2018
<u>AP</u>	! QILU PHARM HAINAN	<u>500MG/VIAL</u>	<u>A215316</u>	<u>001</u>	Aug 24, 2021
<u>AP</u>	+! SAGENT PHARMS INC	<u>350MG/VIAL</u>	<u>N208385</u>	<u>001</u>	Sep 12, 2017
<u>AP</u>		<u>500MG/VIAL</u>	<u>A207104</u>	<u>001</u>	Nov 15, 2019
<u>AP</u>	TEVA PHARMS USA	<u>500MG/VIAL</u>	<u>A091039</u>	<u>001</u>	Mar 25, 2016
<u>AP</u>	XELLIA PHARMS APS	<u>500MG/VIAL</u>	<u>A206005</u>	<u>001</u>	Jun 15, 2016
	+! HOSPIRA	350MG/VIAL	N210282	001	Jun 21, 2021
	+!	500MG/VIAL	N210282	002	Jun 21, 2021
	+! XELLIA PHARMS APS	350MG/VIAL	N209949	001	Oct 20, 2017
	+!	350MG/VIAL	N217415	001	Jan 30, 2023
	+!	500MG/VIAL	N217415	002	Jan 30, 2023

SOLUTION; INTRAVENOUS

DAPTOMYCIN IN 0.9% SODIUM CHLORIDE

	+! BAXTER HLTHCARE CORP	350MG/50ML (7MG/ML)	N213645	002	Feb 27, 2023
	+!	500MG/50ML (10MG/ML)	N213645	003	Feb 27, 2023
	+!	700MG/100ML (7MG/ML)	N213645	004	Feb 27, 2023
	+!	1GM/100ML (10MG/ML)	N213645	005	Feb 27, 2023

DARIDOREXANT HYDROCHLORIDE

TABLET; ORAL

QUVIVIQ

	+ IDORSIA	EQ 25MG BASE	N214985	001	Apr 07, 2022
	+!	EQ 50MG BASE	N214985	002	Apr 07, 2022

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

DARIFENACIN

<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 7.5MG BASE</u>	<u>A207302</u>	<u>001</u>	Jul 28, 2017
<u>AB</u>	!	<u>EQ 15MG BASE</u>	<u>A207302</u>	<u>002</u>	Jul 28, 2017

DARIFENACIN HYDROBROMIDE

<u>AB</u>	ALEMBIC	<u>EQ 7.5MG BASE</u>	<u>A207681</u>	<u>001</u>	Dec 08, 2017
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A207681</u>	<u>002</u>	Dec 08, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 7.5MG BASE</u>	<u>A206743</u>	<u>001</u>	Sep 19, 2016
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A206743</u>	<u>002</u>	Sep 19, 2016
<u>AB</u>	CIPLA	<u>EQ 7.5MG BASE</u>	<u>A207664</u>	<u>001</u>	Sep 01, 2016
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A207664</u>	<u>002</u>	Sep 01, 2016
<u>AB</u>	POLYGEN PHARMS	<u>EQ 7.5MG BASE</u>	<u>A211045</u>	<u>001</u>	Jan 06, 2020
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A211045</u>	<u>002</u>	Jan 06, 2020
<u>AB</u>	TORRENT	<u>EQ 7.5MG BASE</u>	<u>A205209</u>	<u>001</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A205209</u>	<u>002</u>	Nov 17, 2016

DAROLUTAMIDE

TABLET; ORAL

NUBEQA

	+! BAYER HEALTHCARE	300MG	N212099	001	Jul 30, 2019
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DARUNAVIR

SUSPENSION; ORAL

PREZISTA

	+! JANSSEN PRODS	100MG/ML	N202895	001	Dec 16, 2011
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TABLET; ORAL

DARUNAVIR

<u>AB</u>	AMNEAL	<u>600MG</u>	<u>A212493</u>	<u>001</u>	Dec 08, 2023
<u>AB</u>		<u>800MG</u>	<u>A212493</u>	<u>002</u>	Dec 08, 2023
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A210677</u>	<u>001</u>	Nov 28, 2023
<u>AB</u>		<u>800MG</u>	<u>A210677</u>	<u>002</u>	Nov 28, 2023
<u>AB</u>	CIPLA	<u>600MG</u>	<u>A206288</u>	<u>001</u>	Nov 28, 2023
<u>AB</u>		<u>800MG</u>	<u>A206288</u>	<u>002</u>	Nov 28, 2023
<u>AB</u>	DR REDDYS	<u>600MG</u>	<u>A211578</u>	<u>001</u>	Nov 28, 2023
<u>AB</u>		<u>800MG</u>	<u>A211578</u>	<u>002</u>	Nov 28, 2023
<u>AB</u>	HETERO LABS LTD III	<u>600MG</u>	<u>A202083</u>	<u>002</u>	Sep 14, 2023
<u>AB</u>	LUPIN LTD	<u>600MG</u>	<u>A202073</u>	<u>001</u>	Sep 29, 2022
<u>AB</u>		<u>800MG</u>	<u>A202073</u>	<u>002</u>	Sep 29, 2022
<u>AB</u>	MSN	<u>600MG</u>	<u>A215389</u>	<u>001</u>	Nov 28, 2023
<u>AB</u>		<u>800MG</u>	<u>A215389</u>	<u>002</u>	Nov 28, 2023
<u>AB</u>	TEVA PHARMS USA	<u>600MG</u>	<u>A202118</u>	<u>001</u>	Nov 21, 2017
<u>AB</u>	ZYDUS PHARMS	<u>600MG</u>	<u>A214085</u>	<u>001</u>	Dec 13, 2023
<u>AB</u>		<u>800MG</u>	<u>A214085</u>	<u>002</u>	Dec 13, 2023

PRESCRIPTION DRUG PRODUCT LIST

DARUNAVIR

TABLET; ORAL

PREZISTA

<u>AB</u>	+	JANSSEN PRODS	<u>600MG</u>	<u>N021976</u>	<u>002</u>	Feb 25, 2008
<u>AB</u>	+	!	<u>800MG</u>	<u>N021976</u>	<u>006</u>	Nov 09, 2012
		DARUNAVIR				
		HETERO LABS LTD III	400MG	A202083	001	Sep 14, 2023
		PREZISTA				
		+	75MG	N021976	004	Dec 18, 2008
		+	150MG	N021976	005	Dec 18, 2008

DASATINIB

TABLET; ORAL

PHYRAGO

		+	NANOCOPOEIA	20MG	N216099	001	Dec 05, 2023
		+		50MG	N216099	002	Dec 05, 2023
		+		70MG	N216099	003	Dec 05, 2023
		+		80MG	N216099	004	Dec 05, 2023
		+	!	100MG	N216099	005	Dec 05, 2023
		+		140MG	N216099	006	Dec 05, 2023

SPRYCEL

		+	BRISTOL MYERS SQUIBB	20MG	N021986	001	Jun 28, 2006
		+		50MG	N021986	002	Jun 28, 2006
		+		70MG	N021986	003	Jun 28, 2006
		+		80MG	N021986	005	Oct 28, 2010
		+	!	100MG	N021986	004	May 30, 2008
		+		140MG	N021986	006	Oct 28, 2010

DASIGLUCAGON HYDROCHLORIDE

SOLUTION; SUBCUTANEOUS

ZEGALOGUE

		+	!	ZEALAND PHARMA	EQ 0.6MG BASE/0.6ML (EQ 0.6MG BASE/0.6ML)	N214231	001	Mar 22, 2021
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ZEGALOGUE (AUTOINJECTOR)

		+	!	ZEALAND PHARMA	EQ 0.6MG BASE/0.6ML (EQ 0.6MG BASE/0.6ML)	N214231	002	Mar 22, 2021
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DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

CERUBIDINE

<u>AP</u>	!	HIKMA	<u>EQ 20MG BASE/VIAL</u>	<u>A064103</u>	<u>001</u>	Feb 03, 1995
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DAUNORUBICIN HYDROCHLORIDE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 20MG BASE/VIAL</u>	<u>A065000</u>	<u>001</u>	May 25, 1999	
<u>AP</u>	+	!	HIKMA	<u>EQ 5MG BASE/ML</u>	<u>N050731</u>	001	Jan 30, 1998
<u>AP</u>		HISUN PHARM HANGZHOU	<u>EQ 5MG BASE/ML</u>	<u>A208759</u>	<u>001</u>	Apr 12, 2019	
<u>AP</u>		MEITHEAL FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A065035</u>	<u>001</u>	Jan 24, 2000	
			EQ 5MG BASE/VIAL	A065034	001	Nov 20, 2001	

DECITABINE

INJECTABLE; INTRAVENOUS

DECITABINE

<u>AP</u>	!	ACCORD HLTHCARE	<u>50MG/VIAL</u>	<u>A203475</u>	<u>001</u>	Feb 27, 2017
<u>AP</u>		CHEMI SPA	<u>50MG/VIAL</u>	<u>A206033</u>	<u>001</u>	Sep 22, 2017
<u>AP</u>		DR REDDYS	<u>50MG/VIAL</u>	<u>A203131</u>	<u>001</u>	Jul 11, 2013
<u>AP</u>		EUGIA PHARMA	<u>50MG/VIAL</u>	<u>A214569</u>	<u>001</u>	Sep 20, 2021
<u>AP</u>		GLAND	<u>50MG/VIAL</u>	<u>A205539</u>	<u>001</u>	Nov 23, 2020
<u>AP</u>		JIANGSU HANSON PHARM	<u>50MG/VIAL</u>	<u>A213472</u>	<u>001</u>	Apr 15, 2022
<u>AP</u>		LUPIN LTD	<u>50MG/VIAL</u>	<u>A210756</u>	<u>001</u>	Nov 09, 2018
<u>AP</u>		MEITHEAL	<u>50MG/VIAL</u>	<u>A212959</u>	<u>001</u>	Jul 02, 2021
<u>AP</u>		MSN	<u>50MG/VIAL</u>	<u>A212265</u>	<u>001</u>	Aug 28, 2019
<u>AP</u>		NIVAGEN PHARMS INC	<u>50MG/VIAL</u>	<u>A212117</u>	<u>001</u>	Dec 07, 2020
<u>AP</u>		NOVAST LABS	<u>50MG/VIAL</u>	<u>A210984</u>	<u>001</u>	Sep 16, 2019
<u>AP</u>		PHARMASCIENCE INC	<u>50MG/VIAL</u>	<u>A204607</u>	<u>001</u>	May 31, 2017
<u>AP</u>		QILU PHARM HAINAN	<u>50MG/VIAL</u>	<u>A212826</u>	<u>001</u>	Apr 12, 2021
<u>AP</u>		SAGENT PHARMS INC	<u>50MG/VIAL</u>	<u>A207100</u>	<u>001</u>	Mar 16, 2018
<u>AP</u>		SANDOZ	<u>50MG/VIAL</u>	<u>A202969</u>	<u>001</u>	Aug 28, 2014
<u>AP</u>		WOCKHARDT BIO AG	<u>50MG/VIAL</u>	<u>A209056</u>	<u>001</u>	Apr 09, 2019
<u>AP</u>		ZYDUS PHARMS	<u>50MG/VIAL</u>	<u>A214486</u>	<u>001</u>	Nov 19, 2021

POWDER; INTRAVENOUS

DECITABINE

		+	!	SUN PHARM	50MG/VIAL	N205582	001	Jan 28, 2014
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PRESCRIPTION DRUG PRODUCT LIST

DEFERASIROX

GRANULE; ORAL

DEFERASIROX

<u>AB</u>	ALKEM LABS LTD	<u>90MG</u>	<u>A213374 001</u>	Jul 14, 2020
<u>AB</u>		<u>180MG</u>	<u>A213374 002</u>	Jul 14, 2020
<u>AB</u>		<u>360MG</u>	<u>A213374 003</u>	Jul 14, 2020
<u>AB</u>	ANNORA PHARMA	<u>90MG</u>	<u>A216229 003</u>	Mar 16, 2023
<u>AB</u>		<u>180MG</u>	<u>A216229 001</u>	Sep 22, 2022
<u>AB</u>		<u>360MG</u>	<u>A216229 002</u>	Sep 22, 2022
<u>AB</u>	AUCTA	<u>90MG</u>	<u>A214559 001</u>	Mar 09, 2021
<u>AB</u>		<u>180MG</u>	<u>A214559 002</u>	Mar 09, 2021
<u>AB</u>		<u>360MG</u>	<u>A214559 003</u>	Mar 09, 2021
<u>AB</u>	CIPLA	<u>90MG</u>	<u>A215026 001</u>	Feb 23, 2022
<u>AB</u>		<u>180MG</u>	<u>A215026 002</u>	Feb 23, 2022
<u>AB</u>		<u>360MG</u>	<u>A215026 003</u>	Feb 23, 2022
<u>AB</u>	MSN	<u>90MG</u>	<u>A214650 003</u>	Apr 20, 2022
<u>AB</u>		<u>180MG</u>	<u>A214650 001</u>	Mar 17, 2021
<u>AB</u>		<u>360MG</u>	<u>A214650 002</u>	Mar 17, 2021
<u>AB</u>	TEVA PHARMS USA	<u>90MG</u>	<u>A214180 001</u>	Nov 19, 2021
<u>AB</u>		<u>180MG</u>	<u>A214180 002</u>	Nov 19, 2021
<u>AB</u>		<u>360MG</u>	<u>A214180 003</u>	Nov 19, 2021
<u>JADENU SPRINKLE</u>				
<u>AB</u>	+ NOVARTIS	<u>90MG</u>	<u>N207968 001</u>	May 18, 2017
<u>AB</u>	+	<u>180MG</u>	<u>N207968 002</u>	May 18, 2017
<u>AB</u>	+!	<u>360MG</u>	<u>N207968 003</u>	May 18, 2017

TABLET; ORAL

DEFERASIROX

<u>AB</u>	ACTAVIS ELIZABETH	<u>90MG</u>	<u>A208697 001</u>	Dec 13, 2019
<u>AB</u>		<u>180MG</u>	<u>A208697 002</u>	Dec 13, 2019
<u>AB</u>		<u>360MG</u>	<u>A208697 003</u>	Dec 13, 2019
<u>AB</u>	ALEMBIC	<u>90MG</u>	<u>A211824 001</u>	Nov 20, 2019
<u>AB</u>		<u>180MG</u>	<u>A211824 003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A211824 002</u>	Nov 20, 2019
<u>AB</u>	ALKEM LABS LTD	<u>90MG</u>	<u>A210555 001</u>	Mar 30, 2020
<u>AB</u>		<u>180MG</u>	<u>A210555 003</u>	Jul 02, 2020
<u>AB</u>		<u>360MG</u>	<u>A210555 002</u>	Mar 30, 2020
<u>AB</u>	ANNORA PHARMA	<u>90MG</u>	<u>A214341 001</u>	May 14, 2021
<u>AB</u>		<u>180MG</u>	<u>A214341 002</u>	May 14, 2021
<u>AB</u>		<u>360MG</u>	<u>A214341 003</u>	May 14, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>90MG</u>	<u>A214474 001</u>	Oct 16, 2023
<u>AB</u>		<u>180MG</u>	<u>A214474 002</u>	Oct 16, 2023
<u>AB</u>		<u>360MG</u>	<u>A214474 003</u>	Oct 16, 2023
<u>AB</u>	CHARTWELL RX	<u>90MG</u>	<u>A212669 001</u>	May 27, 2021
<u>AB</u>		<u>180MG</u>	<u>A212669 002</u>	May 27, 2021
<u>AB</u>		<u>360MG</u>	<u>A212669 003</u>	May 27, 2021
<u>AB</u>	MSN	<u>90MG</u>	<u>A210945 001</u>	Nov 20, 2019
<u>AB</u>		<u>180MG</u>	<u>A210945 003</u>	Jun 16, 2020
<u>AB</u>		<u>360MG</u>	<u>A210945 002</u>	Nov 20, 2019
<u>AB</u>	PIRAMAL HLTHCARE UK	<u>90MG</u>	<u>A212995 001</u>	Dec 30, 2019
<u>AB</u>		<u>180MG</u>	<u>A212995 003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A212995 002</u>	Dec 30, 2019
<u>AB</u>	SUN PHARM	<u>90MG</u>	<u>A211641 001</u>	Jan 02, 2020
<u>AB</u>		<u>180MG</u>	<u>A211641 003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A211641 002</u>	Jan 02, 2020
<u>AB</u>	ZYDUS PHARMS	<u>90MG</u>	<u>A211383 001</u>	Nov 20, 2019
<u>AB</u>		<u>180MG</u>	<u>A211383 003</u>	Jun 15, 2020
<u>AB</u>		<u>360MG</u>	<u>A211383 002</u>	Nov 20, 2019
<u>JADENU</u>				
<u>AB</u>	+ NOVARTIS PHARMS CORP	<u>90MG</u>	<u>N206910 001</u>	Mar 30, 2015
<u>AB</u>	+	<u>180MG</u>	<u>N206910 002</u>	Mar 30, 2015
<u>AB</u>	+!	<u>360MG</u>	<u>N206910 003</u>	Mar 30, 2015

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

<u>AB</u>	ACTAVIS ELIZABETH	<u>125MG</u>	<u>A203560 001</u>	Jan 26, 2016
<u>AB</u>		<u>250MG</u>	<u>A203560 002</u>	Jan 26, 2016
<u>AB</u>		<u>500MG</u>	<u>A203560 003</u>	Jan 26, 2016
<u>AB</u>	ALEMBIC	<u>125MG</u>	<u>A210060 001</u>	Nov 20, 2019
<u>AB</u>		<u>250MG</u>	<u>A210060 002</u>	Nov 20, 2019
<u>AB</u>		<u>500MG</u>	<u>A210060 003</u>	Nov 20, 2019
<u>AB</u>	ALKEM LABS LTD	<u>125MG</u>	<u>A210519 001</u>	Nov 20, 2019
<u>AB</u>		<u>250MG</u>	<u>A210519 002</u>	Nov 20, 2019

PRESCRIPTION DRUG PRODUCT LIST

DEFERASIROX

TABLET, FOR SUSPENSION;ORAL

DEFERASIROX

<u>AB</u>		<u>500MG</u>	<u>A210519 003</u>	Nov 20, 2019
<u>AB</u>	BIONPHARMA	<u>125MG</u>	<u>A210920 001</u>	Nov 20, 2019
<u>AB</u>		<u>250MG</u>	<u>A210920 002</u>	Nov 20, 2019
<u>AB</u>		<u>500MG</u>	<u>A210920 003</u>	Nov 20, 2019
<u>AB</u>	ICHNOS	<u>125MG</u>	<u>A209433 001</u>	Jan 06, 2020
<u>AB</u>		<u>250MG</u>	<u>A209433 002</u>	Jan 06, 2020
<u>AB</u>		<u>500MG</u>	<u>A209433 003</u>	Jan 06, 2020
<u>AB</u>	MSN	<u>125MG</u>	<u>A209878 001</u>	Nov 20, 2019
<u>AB</u>		<u>250MG</u>	<u>A209878 002</u>	Nov 20, 2019
<u>AB</u>		<u>500MG</u>	<u>A209878 003</u>	Nov 20, 2019
<u>AB</u>	SUN PHARM	<u>125MG</u>	<u>A209782 001</u>	Nov 20, 2019
<u>AB</u>		<u>250MG</u>	<u>A209782 002</u>	Nov 20, 2019
<u>AB</u>		<u>500MG</u>	<u>A209782 003</u>	Nov 20, 2019

EXJADE

<u>AB</u>	+	NOVARTIS	<u>125MG</u>	<u>N021882 001</u>	Nov 02, 2005
<u>AB</u>	+		<u>250MG</u>	<u>N021882 002</u>	Nov 02, 2005
<u>AB</u>	+	!	<u>500MG</u>	<u>N021882 003</u>	Nov 02, 2005

DEFERIPRONE

SOLUTION;ORAL

FERRIPROX

+! CHIESI 100MG/ML N208030 001 Sep 09, 2015

TABLET;ORAL

DEFERIPRONE

<u>AB</u>		HIKMA	<u>500MG</u>	<u>A213239 001</u>	Mar 29, 2021
<u>AB</u>	!		<u>1GM</u>	<u>A213239 002</u>	Feb 08, 2022
<u>AB</u>		TARO	<u>500MG</u>	<u>A208800 001</u>	Feb 08, 2019
<u>AB</u>			<u>1GM</u>	<u>A208800 002</u>	Nov 22, 2023

FERRIPROX

<u>AB</u>	+	CHIESI	<u>500MG</u>	<u>N021825 001</u>	Oct 14, 2011
<u>AB</u>	+		<u>1GM</u>	<u>N021825 002</u>	Jul 25, 2019
	+	!	1GM	N212269 001	May 19, 2020

DEFEROXAMINE MESYLATE

INJECTABLE;INJECTION

DEFEROXAMINE MESYLATE

<u>AP</u>		FRESENIUS KABI USA	<u>500MG/VIAL</u>	<u>A078718 001</u>	Sep 15, 2009
<u>AP</u>			<u>2GM/VIAL</u>	<u>A078718 002</u>	Sep 15, 2009
<u>AP</u>		GLAND PHARMA LTD	<u>500MG/VIAL</u>	<u>A207384 001</u>	Sep 29, 2017
<u>AP</u>			<u>2GM/VIAL</u>	<u>A207384 002</u>	Sep 29, 2017
<u>AP</u>		HOSPIRA	<u>500MG/VIAL</u>	<u>A076019 001</u>	Mar 17, 2004
<u>AP</u>	!		<u>2GM/VIAL</u>	<u>A076019 002</u>	Mar 17, 2004
<u>AP</u>		WEST-WARD PHARMS INT	<u>500MG/VIAL</u>	<u>A078086 001</u>	May 30, 2007
<u>AP</u>			<u>2GM/VIAL</u>	<u>A078086 002</u>	May 30, 2007
<u>AP</u>	+	!	NOVARTIS	<u>500MG/VIAL</u>	<u>N016267 001</u>

DEFIBROTIDE SODIUM

SOLUTION;INTRAVENOUS

DEFITELIO

+! JAZZ PHARMS INC 200MG/2.5ML (80MG/ML) N208114 001 Mar 30, 2016

DEFLAZACORT

SUSPENSION;ORAL

EMFLAZA

+! PTC THERAP 22.75MG/ML N208685 001 Feb 09, 2017

TABLET;ORAL

EMFLAZA

+	PTC THERAP	6MG	N208684 001	Feb 09, 2017
+		18MG	N208684 002	Feb 09, 2017
+		30MG	N208684 003	Feb 09, 2017
+	!	36MG	N208684 004	Feb 09, 2017

DEGARELIX ACETATE

POWDER;SUBCUTANEOUS

FIRMAGON

+	FERRING	EQ 80MG BASE/VIAL	N022201 001	Dec 24, 2008
+	!	EQ 120MG BASE/VIAL	N022201 002	Dec 24, 2008

PRESCRIPTION DRUG PRODUCT LIST

DELAFLOXACIN MEGLUMINE

POWDER; INTRAVENOUS

BAXDELA

+! MELINTA

EQ 300MG BASE/VIAL

N208611 001 Jun 19, 2017

TABLET; ORAL

BAXDELA

+! MELINTA

EQ 450MG BASE

N208610 001 Jun 19, 2017

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DEMECLOCYCLINE HYDROCHLORIDE

AB	AMNEAL PHARM	150MG	<u>A065425 001</u>	Feb 27, 2008
AB	!	300MG	<u>A065425 002</u>	Feb 27, 2008
AB	EPIC PHARMA LLC	150MG	<u>A065389 001</u>	Dec 01, 2008
AB		150MG	<u>A065447 001</u>	Aug 18, 2015
AB		300MG	<u>A065389 002</u>	Dec 01, 2008
AB		300MG	<u>A065447 002</u>	Aug 18, 2015

DEOXYCHOLIC ACID

SOLUTION; SUBCUTANEOUS

DEOXYCHOLIC ACID

AP	SLAYBACK PHARMA LLC	20MG/2ML (10MG/ML)	<u>A212296 001</u>	Apr 02, 2021
AP	+! KYTHERA BIOPHARMS	20MG/2ML (10MG/ML)	<u>N206333 001</u>	Apr 29, 2015

DESFLURANE

LIQUID; INHALATION

DESFLURANE

AN	SHANGHAI HENGRUI	100%	<u>A208234 001</u>	Feb 26, 2018
AN	+! BAXTER HLTHCARE	100%	<u>N020118 001</u>	Sep 18, 1992

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

AB	ACTAVIS TOTOWA	10MG	<u>A074430 001</u>	Feb 09, 1996
AB		25MG	<u>A071601 001</u>	Jun 05, 1987
AB		50MG	<u>A071588 001</u>	Jun 05, 1987
AB		75MG	<u>A071602 001</u>	Oct 05, 1987
AB		100MG	<u>A071766 001</u>	Oct 05, 1987
AB		150MG	<u>A074430 002</u>	Feb 09, 1996
AB	ALEMBIC	10MG	<u>A209785 001</u>	Jul 07, 2021
AB		25MG	<u>A209785 002</u>	Jul 07, 2021
AB		50MG	<u>A209785 003</u>	Jul 07, 2021
AB		75MG	<u>A209785 004</u>	Jul 07, 2021
AB		100MG	<u>A209785 005</u>	Jul 07, 2021
AB		150MG	<u>A209785 006</u>	Jul 07, 2021
AB	AMNEAL PHARMS CO	10MG	<u>A208105 001</u>	Mar 17, 2016
AB		25MG	<u>A208105 002</u>	Mar 17, 2016
AB		50MG	<u>A208105 003</u>	Mar 17, 2016
AB		75MG	<u>A208105 004</u>	Mar 17, 2016
AB		100MG	<u>A208105 005</u>	Mar 17, 2016
AB		150MG	<u>A208105 006</u>	Mar 17, 2016
AB	HERITAGE PHARMS	10MG	<u>A207433 001</u>	May 05, 2016
AB		25MG	<u>A207433 002</u>	May 05, 2016
AB		50MG	<u>A207433 003</u>	May 05, 2016
AB		75MG	<u>A207433 004</u>	May 05, 2016
AB		100MG	<u>A207433 005</u>	May 05, 2016
AB		150MG	<u>A207433 006</u>	May 05, 2016
AB	NOVAST LABS	10MG	<u>A204963 001</u>	Dec 26, 2017
AB		25MG	<u>A204963 002</u>	Dec 26, 2017
AB		50MG	<u>A204963 003</u>	Dec 26, 2017
AB		75MG	<u>A204963 004</u>	Dec 26, 2017
AB		100MG	<u>A204963 005</u>	Dec 26, 2017
AB		150MG	<u>A204963 006</u>	Dec 26, 2017
AB	SANDOZ	10MG	<u>A072103 002</u>	May 24, 1988
AB		25MG	<u>A072103 003</u>	May 24, 1988
AB		50MG	<u>A072103 004</u>	May 24, 1988
AB		75MG	<u>A072103 005</u>	Jun 20, 1988
AB		100MG	<u>A072103 001</u>	Jun 20, 1988
AB		150MG	<u>A072103 006</u>	Jun 20, 1988
<u>NORPRAMIN</u>				
AB	+ VALIDUS PHARMS	10MG	<u>N014399 007</u>	Feb 11, 1982
AB	+	25MG	<u>N014399 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

DESIPRAMINE HYDROCHLORIDE

TABLET;ORAL

NORPRAMIN

AB	+		50MG	<u>N014399</u>	<u>003</u>	
AB	+		75MG	<u>N014399</u>	<u>004</u>	
AB	+	!	100MG	<u>N014399</u>	<u>005</u>	
AB	+		150MG	<u>N014399</u>	<u>006</u>	

DESLORATADINE

TABLET;ORAL

CLARINEX

AB	+	!	ORGANON	5MG	<u>N021165</u>	<u>001</u>	Dec 21, 2001
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DESLORATADINE

AB			BELCHER PHARMS	5MG	<u>A078355</u>	<u>001</u>	Apr 19, 2012
AB			DR REDDYS LABS LTD	5MG	<u>A078365</u>	<u>001</u>	Mar 08, 2011
AB			LUPIN PHARMS	5MG	<u>A078352</u>	<u>001</u>	Oct 25, 2010
AB			ORBION PHARMS	5MG	<u>A078357</u>	<u>001</u>	Feb 19, 2010

TABLET, ORALLY DISINTEGRATING;ORAL

DESLORATADINE

			REDDYS	2.5MG	A078367	001	Jul 12, 2010
		!		5MG	A078367	002	Jul 12, 2010

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CLARINEX-D 12 HOUR

		+	!	ORGANON LLC	2.5MG;120MG	N021313	001	Feb 01, 2006
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DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR

		!		DR REDDYS LABS LTD	5MG;240MG	A078366	001	Apr 26, 2011
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DESMOPRESSIN ACETATE

INJECTABLE;INJECTION

DDAVP

AP	+	!	FERRING PHARMS INC	0.004MG/ML	<u>N018938</u>	<u>001</u>	Mar 30, 1984
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DESMOPRESSIN ACETATE

AP			DR REDDYS	0.004MG/ML	<u>A215961</u>	<u>001</u>	Aug 19, 2022
AP			GLAND	0.004MG/ML	<u>A216904</u>	<u>001</u>	Mar 20, 2023
AP			GLAND PHARMA LTD	0.004MG/ML	<u>A216922</u>	<u>001</u>	Nov 16, 2022
AP			MEITHEAL	0.004MG/ML	<u>A074888</u>	<u>001</u>	Oct 15, 1997
AP			SAGENT PHARMS INC	0.004MG/ML	<u>A204695</u>	<u>001</u>	Aug 22, 2017
AP				0.004MG/ML	<u>A204751</u>	<u>001</u>	Aug 22, 2017
AP			SUN PHARM INDS LTD	0.004MG/ML	<u>A091280</u>	<u>001</u>	Jan 25, 2013
AP			UBI	0.004MG/ML	<u>A210218</u>	<u>001</u>	Feb 14, 2020
AP				0.004MG/ML	<u>A210223</u>	<u>001</u>	Sep 17, 2020

SPRAY, METERED;NASAL

DESMOPRESSIN ACETATE

AB	!		BAUSCH	0.01MG/SPRAY	<u>A074830</u>	<u>001</u>	Jan 25, 1999
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DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)

AB	!		APOTEX	0.01MG/SPRAY	<u>A076703</u>	<u>001</u>	Jan 27, 2005
AB			ZYDUS PHARMS	0.01MG/SPRAY	<u>A091345</u>	<u>001</u>	Oct 03, 2017

TABLET;ORAL

DDAVP

AB	+		FERRING PHARMS INC	0.1MG	<u>N019955</u>	<u>001</u>	Sep 06, 1995
AB	+	!		0.2MG	<u>N019955</u>	<u>002</u>	Sep 06, 1995

DESMOPRESSIN ACETATE

AB			ABHAI LLC	0.1MG	<u>A210371</u>	<u>001</u>	Jan 28, 2019
AB				0.2MG	<u>A210371</u>	<u>002</u>	Jan 28, 2019
AB			ACTAVIS LABS FL INC	0.1MG	<u>A076470</u>	<u>001</u>	Jul 01, 2005
AB				0.2MG	<u>A076470</u>	<u>002</u>	Jul 01, 2005
AB			APOTEX INC	0.1MG	<u>A077414</u>	<u>001</u>	Mar 07, 2006
AB				0.2MG	<u>A077414</u>	<u>002</u>	Mar 07, 2006
AB			GLENMARK PHARMS LTD	0.1MG	<u>A201831</u>	<u>001</u>	May 28, 2015
AB				0.2MG	<u>A201831</u>	<u>002</u>	May 28, 2015
AB			HERITAGE PHARMA	0.1MG	<u>A207880</u>	<u>001</u>	May 26, 2017
AB				0.2MG	<u>A207880</u>	<u>002</u>	May 26, 2017
AB			NOVAST LABS	0.1MG	<u>A208357</u>	<u>001</u>	Jun 06, 2019
AB				0.2MG	<u>A208357</u>	<u>002</u>	Jun 06, 2019

DESOGESTREL; ETHINYL ESTRADIOL

TABLET;ORAL-28

BEKYREE

AB			LUPIN LTD	0.15MG,N/A;0.02MG,0.01MG	<u>A202226</u>	<u>001</u>	Aug 12, 2015
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CYCLESSA

AB	+	!	ASPEN GLOBAL INC	0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG	<u>N021090</u>	<u>001</u>	Dec 20, 2000
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PRESCRIPTION DRUG PRODUCT LIST

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-28

DESOGESTREL AND ETHINYL ESTRADIOL

<u>AB</u>	<u>!</u>	DURAMED PHARMS BARR	<u>0.15MG;0.03MG</u>	<u>A075256</u>	<u>002</u>	Aug 12, 1999
<u>AB</u>		MYLAN LABS LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202296</u>	<u>001</u>	Aug 30, 2013
<u>AB</u>		NAARI PTE LTD	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A209170</u>	<u>001</u>	Jun 05, 2017
<u>AB</u>		NOVAST LABS	<u>0.15MG;0.03MG</u>	<u>A091234</u>	<u>001</u>	Jul 12, 2013
<u>AB</u>		WATSON LABS	<u>0.15MG;0.03MG</u>	<u>A076915</u>	<u>001</u>	Jul 29, 2005
<u>ENSKYCE</u>						
<u>AB</u>		LUPIN LTD	<u>0.15MG;0.03MG</u>	<u>A201887</u>	<u>001</u>	Mar 07, 2013
<u>ISIBLOOM</u>						
<u>AB</u>		XIROMED	<u>0.15MG;0.03MG</u>	<u>A202789</u>	<u>001</u>	Aug 12, 2015
<u>KALLIGA</u>						
<u>AB</u>		AUROBINDO PHARMA	<u>0.15MG;0.03MG</u>	<u>A207081</u>	<u>001</u>	May 17, 2017
<u>KARIYA</u>						
<u>AB</u>	<u>!</u>	BARR	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A075863</u>	<u>001</u>	Apr 05, 2002
<u>KIMIDESS</u>						
<u>AB</u>		VINTAGE PHARMS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A076681</u>	<u>001</u>	Apr 30, 2015
<u>PIMTREA</u>						
<u>AB</u>		NOVAST LABS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A091247</u>	<u>001</u>	Aug 01, 2013
<u>SIMLIYA</u>						
<u>AB</u>		AUROBINDO PHARMA	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A206853</u>	<u>001</u>	Mar 22, 2017
<u>VELIVET</u>						
<u>AB</u>		DURAMED PHARMS BARR	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>A076455</u>	<u>001</u>	Feb 24, 2004
<u>VIORELE</u>						
<u>AB</u>		GLENMARK GENERICS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A091346</u>	<u>001</u>	Apr 02, 2012
<u>VOLNEA</u>						
<u>AB</u>		XIROMED	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A202689</u>	<u>001</u>	Sep 09, 2016

DESONIDE

CREAM; TOPICAL

DESONIDE

<u>AB</u>		ALEMBIC	<u>0.05%</u>	<u>A214396</u>	<u>001</u>	Dec 08, 2022
<u>AB</u>		CADILA	<u>0.05%</u>	<u>A210198</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>		COSETTE	<u>0.05%</u>	<u>A074027</u>	<u>001</u>	Sep 28, 1992
<u>AB</u>		GLENMARK PHARMS	<u>0.05%</u>	<u>A209729</u>	<u>001</u>	Jul 24, 2017
<u>AB</u>	<u>+</u>	PADAGIS US	<u>0.05%</u>	<u>N017010</u>	<u>001</u>	
<u>AB</u>		TARO	<u>0.05%</u>	<u>A073548</u>	<u>001</u>	Jun 30, 1992

DESOWEN

<u>AB</u>		GALDERMA LABS LP	<u>0.05%</u>	<u>N019048</u>	<u>001</u>	Dec 14, 1984
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GEL; TOPICAL

DESONIDE

<u>!</u>		CINTEX SVCS	0.05%	A202470	001	May 11, 2020
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LOTION; TOPICAL

DESONIDE

<u>AB</u>		ALEMBIC	<u>0.05%</u>	<u>A213632</u>	<u>001</u>	Aug 24, 2020
<u>AB</u>		FOUGERA PHARMS	<u>0.05%</u>	<u>A075860</u>	<u>001</u>	Mar 19, 2002
<u>AB</u>		GLENMARK PHARMS	<u>0.05%</u>	<u>A209494</u>	<u>001</u>	Sep 26, 2017
<u>AB</u>		TARO	<u>0.05%</u>	<u>A202161</u>	<u>001</u>	Oct 31, 2014

DESOWEN

<u>AB</u>	<u>!</u>	GALDERMA LABS LP	<u>0.05%</u>	<u>A072354</u>	<u>001</u>	Jan 24, 1992
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OINTMENT; TOPICAL

DESONIDE

<u>AB</u>		ALEMBIC	<u>0.05%</u>	<u>A212473</u>	<u>001</u>	Oct 23, 2019
<u>AB</u>		ENCUBE ETHICALS	<u>0.05%</u>	<u>A210998</u>	<u>001</u>	Jan 30, 2019
<u>AB</u>		FOUGERA PHARMS	<u>0.05%</u>	<u>A075751</u>	<u>001</u>	Mar 12, 2001
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.05%</u>	<u>A209996</u>	<u>001</u>	Sep 15, 2017
<u>AB</u>		HIKMA	<u>0.05%</u>	<u>A208836</u>	<u>001</u>	Mar 27, 2017
<u>AB</u>	<u>+</u>	PADAGIS US	<u>0.05%</u>	<u>N017426</u>	<u>001</u>	
<u>AB</u>		TARO	<u>0.05%</u>	<u>A074254</u>	<u>001</u>	Aug 03, 1994

DESOWEN

<u>AB</u>		GALDERMA LABS LP	<u>0.05%</u>	<u>A071425</u>	<u>001</u>	Jun 15, 1988
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DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

<u>AB</u>		ACTAVIS MID	<u>0.25%</u>	<u>A205082</u>	<u>001</u>	Sep 04, 2015
		ATLANTIC				
<u>AB</u>		CADILA	<u>0.25%</u>	<u>A205620</u>	<u>001</u>	Sep 28, 2018
<u>AB</u>		COSETTE	<u>0.25%</u>	<u>A209595</u>	<u>001</u>	Mar 04, 2020
<u>AB</u>		FOUGERA PHARMS	<u>0.25%</u>	<u>A078369</u>	<u>001</u>	Jun 29, 2010
<u>AB</u>		LUPIN	<u>0.05%</u>	<u>A208163</u>	<u>001</u>	Jan 10, 2017
<u>AB</u>			<u>0.25%</u>	<u>A208164</u>	<u>001</u>	Jan 09, 2017

PRESCRIPTION DRUG PRODUCT LIST

DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

<u>AB</u>	PADAGIS ISRAEL	<u>0.25%</u>	<u>A076510 001</u>	Jul 01, 2003
<u>AB</u>	RISING	<u>0.05%</u>	<u>A210980 001</u>	Dec 21, 2018
<u>AB</u>		<u>0.25%</u>	<u>A205594 001</u>	Jul 02, 2018

TOPICORT

<u>AB</u>	! TARO	<u>0.05%</u>	<u>A073210 001</u>	Nov 30, 1990
<u>AB</u>	!	<u>0.25%</u>	<u>A073193 001</u>	Nov 30, 1990

GEL; TOPICAL

DESOXIMETASONE

<u>AB</u>	EPIC PHARMA LLC	<u>0.05%</u>	<u>A090727 001</u>	Mar 10, 2011
<u>AB</u>	PADAGIS US	<u>0.05%</u>	<u>A077552 001</u>	Jan 09, 2006
<u>AB</u>	RISING	<u>0.05%</u>	<u>A204675 001</u>	Aug 12, 2016

TOPICORT

<u>AB</u>	! TARO	<u>0.05%</u>	<u>A074904 001</u>	Jul 14, 1998
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OINTMENT; TOPICAL

DESOXIMETASONE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>0.25%</u>	<u>A204965 001</u>	Nov 07, 2016
<u>AB</u>	CADILA	<u>0.25%</u>	<u>A205206 001</u>	Sep 19, 2017
<u>AB</u>	COSETTE	<u>0.25%</u>	<u>A206740 001</u>	Dec 23, 2016
<u>AB</u>	EPIC PHARMA LLC	<u>0.25%</u>	<u>A201005 001</u>	Apr 24, 2014
<u>AB</u>	FOUGERA PHARMS	<u>0.25%</u>	<u>A078657 001</u>	Sep 28, 2012
<u>AB</u>	! GLENMARK GENERICS	<u>0.25%</u>	<u>A202838 001</u>	Sep 20, 2013
<u>AB</u>	LUPIN	<u>0.05%</u>	<u>A208044 001</u>	Dec 12, 2016
<u>AB</u>		<u>0.25%</u>	<u>A208104 001</u>	Dec 01, 2016
<u>AB</u>	NOVEL LABS INC	<u>0.25%</u>	<u>A206792 001</u>	May 10, 2016
<u>AB</u>	PADAGIS ISRAEL	<u>0.25%</u>	<u>A077770 001</u>	Apr 20, 2015
<u>AB</u>	RISING	<u>0.25%</u>	<u>A204272 001</u>	Nov 30, 2016
<u>AB</u>	THE J MOLNER	<u>0.05%</u>	<u>A209973 001</u>	Oct 23, 2018

TOPICORT

<u>AB</u>	+! TARO	<u>0.05%</u>	<u>N018594 001</u>	Jan 17, 1985
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SPRAY; TOPICAL

DESOXIMETASONE

<u>AT</u>	LUPIN	<u>0.25%</u>	<u>A208124 001</u>	Mar 16, 2018
<u>AT</u>	PADAGIS ISRAEL	<u>0.25%</u>	<u>A206441 001</u>	Jan 20, 2017

TOPICORT

<u>AT</u>	+! TARO	<u>0.25%</u>	<u>N204141 001</u>	Apr 11, 2013
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DESVENLAFAXINE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE

+	ALEMBIC PHARMS LTD	50MG	N204150 001	Mar 04, 2013
+	!	100MG	N204150 002	Mar 04, 2013

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE SUCCINATE

<u>AB</u>	ACTAVIS LABS FL	<u>EQ 25MG BASE</u>	<u>A204065 001</u>	Jul 29, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204065 002</u>	Jul 29, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204065 003</u>	Jul 29, 2016
<u>AB</u>	ALEMBIC	<u>EQ 25MG BASE</u>	<u>A204003 003</u>	Sep 14, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204003 001</u>	Jun 29, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204003 002</u>	Jun 29, 2015
<u>AB</u>	HIKMA	<u>EQ 25MG BASE</u>	<u>A204082 002</u>	Aug 28, 2017
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204082 001</u>	Feb 16, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204083 001</u>	Feb 16, 2016
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>EQ 50MG BASE</u>	<u>A204805 001</u>	May 07, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204805 002</u>	May 07, 2019
<u>AB</u>	LUPIN LTD	<u>EQ 25MG BASE</u>	<u>A204172 003</u>	Apr 13, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204172 001</u>	Jun 29, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204172 002</u>	Jun 29, 2015
<u>AB</u>	RUBICON	<u>EQ 25MG BASE</u>	<u>A204028 003</u>	Dec 07, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204028 001</u>	Jun 29, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204028 002</u>	Jun 29, 2015
<u>AB</u>	YICHANG HUMANWELL	<u>EQ 25MG BASE</u>	<u>A210014 003</u>	Oct 13, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210014 001</u>	Oct 01, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210014 002</u>	Oct 01, 2018
<u>AB</u>	ZYDUS PHARMS	<u>EQ 25MG BASE</u>	<u>A204020 003</u>	Nov 30, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204020 001</u>	Oct 11, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204020 002</u>	Oct 11, 2017

PRESCRIPTION DRUG PRODUCT LIST

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

PRISTIQ

<u>AB</u>	+	PF PRISM CV	<u>EQ 25MG BASE</u>	<u>N021992 003</u>	Aug 20, 2014
<u>AB</u>	+		<u>EQ 50MG BASE</u>	<u>N021992 001</u>	Feb 29, 2008
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N021992 002</u>	Feb 29, 2008

DEUCRAVACITINIB

TABLET;ORAL

SOTYKTU

+	!	BRISTOL	6MG	N214958 001	Sep 09, 2022
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DEUTETRABENAZINE

TABLET;ORAL

AUSTEDO

+		TEVA BRANDED PHARM	6MG	N208082 001	Apr 03, 2017
+			9MG	N208082 002	Apr 03, 2017
+	!		12MG	N208082 003	Apr 03, 2017

TABLET, EXTENDED RELEASE;ORAL

AUSTEDO XR

+		TEVA	6MG	N216354 001	Feb 17, 2023
+			12MG	N216354 002	Feb 17, 2023
+	!		24MG	N216354 003	Feb 17, 2023

DEXAMETHASONE

CONCENTRATE;ORAL

DEXAMETHASONE INTENSOL

!		HIKMA	1MG/ML	A088252 001	Sep 01, 1983
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ELIXIR;ORAL

DEXAMETHASONE

<u>AA</u>	!	ANIMA	<u>0.5MG/5ML</u>	<u>A084754 001</u>	
<u>AA</u>		CHARTWELL MOLECULAR	<u>0.5MG/5ML</u>	<u>A091188 001</u>	May 11, 2011
<u>AA</u>		LYNE	<u>0.5MG/5ML</u>	<u>A090891 001</u>	Jul 12, 2011

IMPLANT; INTRAVITREAL

OZURDEX

+	!	ABBVIE	0.7MG	N022315 001	Jun 17, 2009
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INSERT;OPHTHALMIC

DEXTENZA

+	!	OCULAR THERAPEUTIX	0.4MG	N208742 001	Nov 30, 2018
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SOLUTION;ORAL

DEXAMETHASONE

+	!	HIKMA	0.5MG/5ML	A088248 001	Sep 01, 1983
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SUSPENSION; INTRAOCULAR

DEXYCU KIT

+	!	EYEPOINT PHARMS	9%	N208912 001	Feb 09, 2018
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SUSPENSION/DROPS;OPHTHALMIC

MAXIDEX

+	!	HARROW EYE	0.1%	N013422 001	
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TABLET;ORAL

DEXAMETHASONE

<u>AB</u>		ALVOGEN	<u>4MG</u>	<u>A088481 004</u>	Apr 28, 1983
<u>AB</u>			<u>6MG</u>	<u>A088481 001</u>	Nov 28, 1983
<u>AB</u>		AMNEAL	<u>2MG</u>	<u>A216295 001</u>	Sep 08, 2022
<u>AB</u>			<u>4MG</u>	<u>A215106 001</u>	Oct 14, 2021
<u>AB</u>			<u>6MG</u>	<u>A215106 002</u>	Oct 14, 2021
<u>AB</u>		APOTEX	<u>0.5MG</u>	<u>A217695 001</u>	Aug 23, 2023
<u>AB</u>			<u>0.75MG</u>	<u>A217695 002</u>	Aug 23, 2023
<u>AB</u>			<u>1MG</u>	<u>A217695 003</u>	Aug 23, 2023
<u>AB</u>			<u>1.5MG</u>	<u>A217695 004</u>	Aug 23, 2023
<u>AB</u>			<u>2MG</u>	<u>A217695 005</u>	Aug 23, 2023
<u>AB</u>			<u>4MG</u>	<u>A217695 006</u>	Aug 23, 2023
<u>AB</u>			<u>6MG</u>	<u>A217695 007</u>	Aug 23, 2023
<u>AB</u>		BIONPHARMA	<u>2MG</u>	<u>A217538 001</u>	Apr 26, 2023
<u>AB</u>			<u>4MG</u>	<u>A217001 001</u>	Apr 19, 2023
<u>AB</u>			<u>6MG</u>	<u>A217001 002</u>	Apr 19, 2023
<u>AB</u>	+	HIKMA	<u>0.5MG</u>	<u>A084611 001</u>	
<u>AB</u>	+		<u>0.75MG</u>	<u>A084613 001</u>	
<u>AB</u>	+		<u>1MG</u>	<u>A088306 001</u>	Sep 15, 1983
<u>AB</u>	+		<u>1.5MG</u>	<u>A084610 001</u>	
<u>AB</u>	+		<u>2MG</u>	<u>A087916 001</u>	Aug 26, 1982
<u>AB</u>	+		<u>4MG</u>	<u>A084612 001</u>	
<u>AB</u>	+		<u>6MG</u>	<u>A088316 001</u>	Sep 15, 1983
<u>AB</u>		LARKEN LABS INC	<u>1.5MG</u>	<u>A201270 001</u>	Jul 17, 2017
<u>AB</u>		NOVITIUM PHARMA	<u>0.5MG</u>	<u>A215604 004</u>	Oct 05, 2023

PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE

TABLET; ORAL

DEXAMETHASONE

<u>AB</u>		<u>0.75MG</u>	<u>A215604 005</u>	Oct 05, 2023
<u>AB</u>		<u>1.5MG</u>	<u>A215604 001</u>	Aug 08, 2022
<u>AB</u>		<u>2MG</u>	<u>A217696 001</u>	May 09, 2023
<u>AB</u>		<u>4MG</u>	<u>A215604 002</u>	Aug 08, 2022
<u>AB</u>		<u>6MG</u>	<u>A215604 003</u>	Aug 08, 2022
<u>AB</u>	PRASCO	<u>4MG</u>	<u>A080399 002</u>	Apr 20, 2022
BP	ALVOGEN	0.5MG	A088481 002	Apr 28, 1983
BP		0.75MG	A088481 003	Apr 28, 1983
BP	XSPIRE PHARMA	1.5MG	A088237 001	Apr 28, 1983
	HEMADY			
	+! DEXCEL	20MG	N211379 001	Oct 03, 2019

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE SODIUM PHOSPHATE

<u>AP</u>	AMNEAL	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A208689 001</u>	Aug 22, 2018
<u>AP</u>	EUGIA PHARMA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A206781 001</u>	Dec 01, 2015
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A210966 001</u>	Jun 05, 2020
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A210967 001</u>	Jun 07, 2019
<u>AP</u>	+! FRESENIUS KABI USA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A084916 001</u>	
<u>AP</u>		<u>EQ 4MG PHOSPHATE/ML</u>	<u>A203129 001</u>	Sep 30, 2015
<u>AP</u>	!	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040572 001</u>	Apr 22, 2005
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A209192 001</u>	Jul 06, 2018
<u>AP</u>	GENEYORK PHARMS	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A214891 001</u>	Jul 28, 2023
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A214890 001</u>	Jul 07, 2023
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A215654 001</u>	Aug 04, 2021
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A215654 002</u>	Sep 25, 2023
<u>AP</u>	HIKMA	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A211451 001</u>	Aug 01, 2023
<u>AP</u>	MYLAN LABS LTD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A040803 001</u>	Aug 29, 2008
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040802 001</u>	Aug 29, 2008
<u>AP</u>	SOMERSET	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A207521 001</u>	Jun 08, 2018
<u>AP</u>	SOMERSET THERAPS LLC	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A211036 001</u>	May 10, 2019
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A084282 001</u>	
<u>AP</u>	!	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A087702 001</u>	Sep 07, 1982

DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE

<u>AP</u>	AMNEAL	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A208690 001</u>	Aug 22, 2018
<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040491 001</u>	Apr 11, 2003
<u>AP</u>	SOMERSET THERAPS LLC	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A207442 001</u>	Apr 19, 2018

SOLUTION/DROPS; OPHTHALMIC, OTIC

DEXAMETHASONE SODIUM PHOSPHATE

!	BAUSCH AND LOMB	EQ 0.1% PHOSPHATE	A040069 001	Jul 26, 1996
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DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

MAXITROL

<u>AT</u>	+! SANDOZ	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N050065 002</u>	
		<u>NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE</u>		
<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064063 001</u>	Jul 25, 1994
<u>AT</u>	PADAGIS US	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A062938 001</u>	Jul 31, 1989

SUSPENSION/DROPS; OPHTHALMIC

DEXASPORIN

<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064135 001</u>	Sep 13, 1995
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MAXITROL

<u>AT</u>	+! HARROW EYE	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N050023 002</u>	
<u>AT</u>	SANDOZ	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062341 001</u>	May 22, 1984

DEXAMETHASONE; TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBRADEX

+!	NOVARTIS	0.1%;0.3%	N050616 001	Sep 28, 1988
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SUSPENSION/DROPS; OPHTHALMIC

TOBRADEX

<u>AB</u>	+! SANDOZ	<u>0.1%;0.3%</u>	<u>N050592 001</u>	Aug 18, 1988
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TOBRAMYCIN AND DEXAMETHASONE

<u>AB</u>	BAUSCH AND LOMB	<u>0.1%;0.3%</u>	<u>A064134 001</u>	Oct 27, 1999
<u>AB</u>	PADAGIS US	<u>0.1%;0.3%</u>	<u>A212715 001</u>	Feb 11, 2022
	TOBRADEX ST			
	+! HARROW EYE	0.05%;0.3%	N050818 001	Feb 13, 2009

PRESCRIPTION DRUG PRODUCT LIST

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL

DEXCHLORPHENIRAMINE MALEATE

<u>AA</u>	<u>+!</u>	PAI HOLDINGS PHARM	<u>2MG/5ML</u>	<u>A088251</u>	<u>001</u>	Mar 23, 1984
<u>POLMON</u>						
<u>AA</u>		CAPELLON PHARMS LLC	<u>2MG/5ML</u>	<u>A202520</u>	<u>001</u>	Jul 16, 2018

DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE; ORAL

DEXILANT

<u>AB</u>	<u>+</u>	TAKEDA PHARMS USA	<u>30MG</u>	<u>N022287</u>	<u>001</u>	Jan 30, 2009
<u>AB</u>	<u>+!</u>		<u>60MG</u>	<u>N022287</u>	<u>002</u>	Jan 30, 2009

DEXLANSOPRAZOLE

<u>AB</u>		PAR PHARM INC	<u>30MG</u>	<u>A202294</u>	<u>002</u>	Jun 16, 2022
<u>AB</u>			<u>60MG</u>	<u>A202294</u>	<u>001</u>	Apr 19, 2017
<u>AB</u>		TWI PHARMS	<u>30MG</u>	<u>A202666</u>	<u>001</u>	Sep 16, 2022
<u>AB</u>			<u>60MG</u>	<u>A202666</u>	<u>002</u>	Sep 16, 2022

DEXMEDETOMIDINE HYDROCHLORIDE

FILM; BUCCAL, SUBLINGUAL

IGALMI

<u>+!</u>	BIOXCEL	EQ 0.12MG BASE	N215390	001	Apr 05, 2022
<u>+!</u>		EQ 0.18MG BASE	N215390	002	Apr 05, 2022

INJECTABLE; INJECTION

DEXMEDETOMIDINE HYDROCHLORIDE

<u>AP</u>		ACCORD HLTHCARE	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204023</u>	<u>001</u>	Feb 09, 2016
<u>AP</u>		ACTAVIS INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204686</u>	<u>001</u>	Oct 17, 2016
<u>AP</u>		AMNEAL	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A216604</u>	<u>001</u>	May 15, 2023
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A216604</u>	<u>002</u>	May 15, 2023
<u>AP</u>		AMNEAL PHARMS CO	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A207551</u>	<u>001</u>	May 20, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A207551</u>	<u>002</u>	May 20, 2020
<u>AP</u>		BAXTER HLTHCARE CORP	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208532</u>	<u>001</u>	Aug 21, 2018
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208532</u>	<u>002</u>	Aug 21, 2018
<u>AP</u>		EUGIA PHARMA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205867</u>	<u>001</u>	Mar 17, 2016
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A210321</u>	<u>001</u>	Dec 07, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A210321</u>	<u>002</u>	Dec 07, 2020
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A208129</u>	<u>001</u>	Nov 29, 2018
<u>AP</u>			<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A201072</u>	<u>001</u>	Sep 18, 2015
<u>AP</u>			<u>EQ 400MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208129</u>	<u>002</u>	Nov 29, 2018
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208129</u>	<u>003</u>	Nov 29, 2018
<u>AP</u>		GLAND	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202126</u>	<u>001</u>	Aug 20, 2015
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A209307</u>	<u>001</u>	Jun 03, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A209307</u>	<u>002</u>	Jun 03, 2020
<u>AP</u>		HENGRUI PHARMA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A209065</u>	<u>002</u>	Jun 12, 2020
<u>AP</u>			<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A209065</u>	<u>001</u>	Sep 19, 2017
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A209065</u>	<u>003</u>	Jun 12, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A209065</u>	<u>004</u>	Jun 12, 2020
<u>AP</u>		HIKMA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205046</u>	<u>001</u>	Apr 26, 2017
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A206407</u>	<u>001</u>	Jan 30, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A206407</u>	<u>002</u>	Jan 30, 2020
<u>AP</u>		MEITHEAL	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A204843</u>	<u>001</u>	Jan 18, 2019
<u>AP</u>		MILLA PHARMS	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A217308</u>	<u>001</u>	Jun 07, 2023
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A217308</u>	<u>002</u>	Jun 07, 2023
<u>AP</u>		MYLAN INSTITUTIONAL	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202881</u>	<u>001</u>	Aug 18, 2014
<u>AP</u>		MYLAN LABS LTD	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212571</u>	<u>001</u>	Aug 27, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212571</u>	<u>002</u>	Aug 27, 2020
<u>AP</u>		PAR STERILE PRODUCTS	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A203972</u>	<u>001</u>	Aug 18, 2014
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A208266</u>	<u>001</u>	Sep 15, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A208266</u>	<u>002</u>	Sep 15, 2020
<u>AP</u>		RISING	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A202585</u>	<u>001</u>	Nov 24, 2014
<u>AP</u>		SANDOZ	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A091465</u>	<u>001</u>	Jun 14, 2016
<u>AP</u>		SLAYBACK PHARMA LLC	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212791</u>	<u>001</u>	Dec 04, 2019
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212791</u>	<u>002</u>	Dec 04, 2019
<u>AP</u>		TAGI	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>A212857</u>	<u>001</u>	Nov 23, 2020
<u>AP</u>			<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>A212857</u>	<u>002</u>	Nov 23, 2020
<u>AP</u>			<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>A212857</u>	<u>003</u>	Nov 23, 2020
<u>AP</u>		TEVA PHARMS USA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A205272</u>	<u>001</u>	Nov 28, 2017
<u>AP</u>		ZYDUS PHARMS	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>A206798</u>	<u>001</u>	Feb 27, 2018
<u>PRECEDEX</u>						
<u>AP</u>	<u>+!</u>	HOSPIRA	<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>004</u>	Nov 14, 2014
<u>AP</u>	<u>+!</u>		<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<u>N021038</u>	<u>001</u>	Dec 17, 1999
<u>AP</u>	<u>+!</u>		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>002</u>	Mar 13, 2013

PRESCRIPTION DRUG PRODUCT LIST

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRECEDEX

<u>AP</u>	<u>+</u> !	<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>003</u>	Mar 13, 2013
	<u>+</u> !	<u>EQ 1MG BASE/250ML (EQ 4MCG BASE/ML)</u>	<u>N021038</u>	<u>005</u>	Jan 31, 2020
SOLUTION; INTRAVENOUS					
DEXMEDETOMIDINE HYDROCHLORIDE					
	<u>+</u> !	<u>HQ SPCLT PHARMA EQ 1MG BASE/10ML (EQ 100MCG BASE/ML)</u>	<u>N206628</u>	<u>002</u>	Oct 21, 2015
	<u>+</u>	<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	<u>N206628</u>	<u>003</u>	Jun 22, 2018
	<u>+</u>	<u>EQ 400MCG BASE/4ML (EQ 100MCG BASE/ML)</u>	<u>N206628</u>	<u>001</u>	Oct 21, 2015
	<u>+</u>	<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	<u>N206628</u>	<u>004</u>	Jun 22, 2018

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ADARE PHARMS INC	<u>5MG</u>	<u>A210279</u>	<u>001</u>	Oct 09, 2018
<u>AB</u>		<u>10MG</u>	<u>A210279</u>	<u>002</u>	Oct 09, 2018
<u>AB</u>		<u>15MG</u>	<u>A210279</u>	<u>003</u>	Oct 09, 2018
<u>AB</u>		<u>20MG</u>	<u>A210279</u>	<u>004</u>	Oct 09, 2018
<u>AB</u>		<u>25MG</u>	<u>A210279</u>	<u>005</u>	Oct 09, 2018
<u>AB</u>		<u>30MG</u>	<u>A210279</u>	<u>006</u>	Oct 09, 2018
<u>AB</u>		<u>35MG</u>	<u>A210279</u>	<u>007</u>	Oct 09, 2018
<u>AB</u>		<u>40MG</u>	<u>A210279</u>	<u>008</u>	Oct 09, 2018
<u>AB</u>	ASCENT PHARMS INC	<u>5MG</u>	<u>A215523</u>	<u>001</u>	Dec 08, 2021
<u>AB</u>		<u>10MG</u>	<u>A215523</u>	<u>002</u>	Dec 08, 2021
<u>AB</u>		<u>15MG</u>	<u>A215523</u>	<u>003</u>	Dec 08, 2021
<u>AB</u>		<u>20MG</u>	<u>A215523</u>	<u>004</u>	Dec 08, 2021
<u>AB</u>		<u>25MG</u>	<u>A215523</u>	<u>005</u>	Dec 08, 2021
<u>AB</u>		<u>30MG</u>	<u>A215523</u>	<u>006</u>	Dec 08, 2021
<u>AB</u>		<u>35MG</u>	<u>A215523</u>	<u>007</u>	Dec 08, 2021
<u>AB</u>		<u>40MG</u>	<u>A215523</u>	<u>008</u>	Dec 08, 2021
<u>AB</u>	GRANULES	<u>5MG</u>	<u>A213813</u>	<u>001</u>	Sep 09, 2020
<u>AB</u>		<u>10MG</u>	<u>A213813</u>	<u>002</u>	Sep 09, 2020
<u>AB</u>		<u>15MG</u>	<u>A213813</u>	<u>003</u>	Sep 09, 2020
<u>AB</u>		<u>20MG</u>	<u>A213813</u>	<u>004</u>	Sep 09, 2020
<u>AB</u>		<u>25MG</u>	<u>A213813</u>	<u>005</u>	Sep 09, 2020
<u>AB</u>		<u>30MG</u>	<u>A213813</u>	<u>006</u>	Sep 09, 2020
<u>AB</u>		<u>35MG</u>	<u>A213813</u>	<u>007</u>	Sep 09, 2020
<u>AB</u>		<u>40MG</u>	<u>A213813</u>	<u>008</u>	Sep 09, 2020
<u>AB</u>	IMPAX LABS INC	<u>5MG</u>	<u>A079108</u>	<u>001</u>	Aug 05, 2015
<u>AB</u>		<u>10MG</u>	<u>A079108</u>	<u>002</u>	Aug 05, 2015
<u>AB</u>		<u>15MG</u>	<u>A079108</u>	<u>003</u>	May 19, 2014
<u>AB</u>		<u>20MG</u>	<u>A079108</u>	<u>004</u>	Dec 21, 2015
<u>AB</u>		<u>25MG</u>	<u>A203614</u>	<u>001</u>	Jul 05, 2017
<u>AB</u>		<u>30MG</u>	<u>A079108</u>	<u>005</u>	Nov 21, 2013
<u>AB</u>		<u>35MG</u>	<u>A203614</u>	<u>002</u>	Jul 05, 2017
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>5MG</u>	<u>A078992</u>	<u>001</u>	Nov 23, 2021
<u>AB</u>		<u>10MG</u>	<u>A078992</u>	<u>002</u>	Nov 23, 2021
<u>AB</u>		<u>15MG</u>	<u>A078992</u>	<u>003</u>	Nov 18, 2013
<u>AB</u>		<u>30MG</u>	<u>A078992</u>	<u>004</u>	Nov 18, 2013
<u>AB</u>		<u>20MG</u>	<u>A078992</u>	<u>005</u>	Nov 23, 2021
<u>AB</u>		<u>40MG</u>	<u>A078992</u>	<u>006</u>	Nov 23, 2021
<u>AB</u>	PAR PHARM INC	<u>5MG</u>	<u>A202842</u>	<u>001</u>	Nov 30, 2016
<u>AB</u>		<u>10MG</u>	<u>A202842</u>	<u>002</u>	Nov 30, 2016
<u>AB</u>		<u>15MG</u>	<u>A202842</u>	<u>003</u>	Nov 30, 2016
<u>AB</u>		<u>20MG</u>	<u>A202842</u>	<u>004</u>	Nov 30, 2016
<u>AB</u>		<u>25MG</u>	<u>A202842</u>	<u>005</u>	Nov 30, 2016
<u>AB</u>		<u>30MG</u>	<u>A202842</u>	<u>006</u>	Nov 30, 2016
<u>AB</u>		<u>35MG</u>	<u>A202842</u>	<u>007</u>	Nov 30, 2016
<u>AB</u>		<u>40MG</u>	<u>A202842</u>	<u>008</u>	Nov 30, 2016
<u>AB</u>	SUN PHARM INDS INC	<u>5MG</u>	<u>A206734</u>	<u>001</u>	Nov 05, 2021
<u>AB</u>		<u>10MG</u>	<u>A206734</u>	<u>002</u>	Nov 05, 2021
<u>AB</u>		<u>15MG</u>	<u>A206734</u>	<u>003</u>	Nov 05, 2021
<u>AB</u>		<u>20MG</u>	<u>A206734</u>	<u>004</u>	Nov 05, 2021
<u>AB</u>		<u>25MG</u>	<u>A206734</u>	<u>005</u>	Nov 05, 2021
<u>AB</u>		<u>30MG</u>	<u>A206734</u>	<u>006</u>	Nov 05, 2021
<u>AB</u>		<u>35MG</u>	<u>A206734</u>	<u>007</u>	Nov 05, 2021
<u>AB</u>		<u>40MG</u>	<u>A206734</u>	<u>008</u>	Nov 05, 2021
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A078908</u>	<u>001</u>	Nov 19, 2013
<u>AB</u>		<u>10MG</u>	<u>A078908</u>	<u>002</u>	Nov 19, 2013
<u>AB</u>		<u>15MG</u>	<u>A078908</u>	<u>004</u>	May 19, 2014
<u>AB</u>		<u>20MG</u>	<u>A078908</u>	<u>003</u>	Nov 19, 2013

PRESCRIPTION DRUG PRODUCT LIST

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		<u>25MG</u>	<u>A202731 001</u>	Jul 05, 2017
<u>AB</u>		<u>30MG</u>	<u>A202731 003</u>	May 19, 2014
<u>AB</u>		<u>35MG</u>	<u>A202731 004</u>	Jul 05, 2017
<u>AB</u>		<u>40MG</u>	<u>A202731 002</u>	Nov 19, 2013

FOCALIN XR

<u>AB</u>	+	SANDOZ	<u>5MG</u>	<u>N021802 001</u>	May 26, 2005
<u>AB</u>	+		<u>10MG</u>	<u>N021802 002</u>	May 26, 2005
<u>AB</u>	+		<u>15MG</u>	<u>N021802 004</u>	Aug 01, 2006
<u>AB</u>	+		<u>20MG</u>	<u>N021802 003</u>	May 26, 2005
<u>AB</u>	+		<u>25MG</u>	<u>N021802 008</u>	Apr 21, 2011
<u>AB</u>	+		<u>30MG</u>	<u>N021802 005</u>	Oct 23, 2009
<u>AB</u>	+		<u>35MG</u>	<u>N021802 007</u>	Apr 21, 2011
<u>AB</u>	+	!	<u>40MG</u>	<u>N021802 006</u>	Aug 11, 2010

TABLET;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		ABHAI INC	<u>2.5MG</u>	<u>A206931 001</u>	Dec 04, 2015
<u>AB</u>			<u>5MG</u>	<u>A206931 002</u>	Dec 04, 2015
<u>AB</u>			<u>10MG</u>	<u>A206931 003</u>	Dec 04, 2015
<u>AB</u>		ALKEM LABS LTD	<u>2.5MG</u>	<u>A212631 001</u>	Jul 19, 2019
<u>AB</u>			<u>5MG</u>	<u>A212631 002</u>	Jul 19, 2019
<u>AB</u>			<u>10MG</u>	<u>A212631 003</u>	Jul 19, 2019
<u>AB</u>		CEDIPROF INC	<u>5MG</u>	<u>A209211 001</u>	Sep 19, 2018
<u>AB</u>			<u>10MG</u>	<u>A209211 002</u>	Sep 19, 2018
<u>AB</u>		NOVEL LABS INC	<u>2.5MG</u>	<u>A204534 001</u>	Dec 04, 2015
<u>AB</u>			<u>5MG</u>	<u>A204534 002</u>	Dec 04, 2015
<u>AB</u>			<u>10MG</u>	<u>A204534 003</u>	Dec 04, 2015
<u>AB</u>		RHODES PHARMS	<u>2.5MG</u>	<u>A208756 001</u>	Nov 20, 2017
<u>AB</u>			<u>5MG</u>	<u>A208756 002</u>	Nov 20, 2017
<u>AB</u>			<u>10MG</u>	<u>A208756 003</u>	Nov 20, 2017
<u>AB</u>		SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A201231 001</u>	Sep 24, 2015
<u>AB</u>			<u>5MG</u>	<u>A201231 002</u>	Sep 24, 2015
<u>AB</u>			<u>10MG</u>	<u>A201231 003</u>	Sep 24, 2015
<u>AB</u>		TRIS PHARMA INC	<u>2.5MG</u>	<u>A207901 001</u>	Aug 26, 2016
<u>AB</u>			<u>5MG</u>	<u>A207901 002</u>	Aug 26, 2016
<u>AB</u>			<u>10MG</u>	<u>A207901 003</u>	Aug 26, 2016

FOCALIN

<u>AB</u>	+	SANDOZ	<u>2.5MG</u>	<u>N021278 001</u>	Nov 13, 2001
<u>AB</u>	+		<u>5MG</u>	<u>N021278 002</u>	Nov 13, 2001
<u>AB</u>	+	!	<u>10MG</u>	<u>N021278 003</u>	Nov 13, 2001

DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE

CAPSULE;ORAL

AZSTARYS

+	COMMAVE THERAP	EQ 5.2MG BASE;EQ 26.1MG BASE	N212994 001	May 07, 2021
+		EQ 7.8MG BASE;EQ 39.2MG BASE	N212994 002	May 07, 2021
+	!	EQ 10.4MG BASE;EQ 52.3MG BASE	N212994 003	May 07, 2021

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE;INJECTION

DEXRAZOXANE HYDROCHLORIDE

<u>AP</u>	!	EUGIA PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A200752 001</u>	Oct 19, 2011
<u>AP</u>	!		<u>EQ 500MG BASE/VIAL</u>	<u>A200752 002</u>	Oct 19, 2011
<u>AP</u>		GLAND	<u>EQ 250MG BASE/VIAL</u>	<u>A207321 002</u>	Dec 16, 2019
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A207321 001</u>	Nov 28, 2016
<u>AP</u>		HIKMA	<u>EQ 250MG BASE/VIAL</u>	<u>A076068 001</u>	Sep 28, 2004
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A076068 002</u>	Sep 28, 2004

DEXTROAMPHETAMINE

SYSTEM;TRANSDERMAL

XELSTRYM

+	NOVEN PHARMS INC	4.5MG/9HR	N215401 001	Mar 22, 2022
+		9MG/9HR	N215401 002	Mar 22, 2022
+		13.5MG/9HR	N215401 003	Mar 22, 2022
+	!	18MG/9HR	N215401 004	Mar 22, 2022

PRESCRIPTION DRUG PRODUCT LIST

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXEDRINE

<u>AB</u>	+	IMPAX LABS INC	<u>5MG</u>	<u>N017078</u>	<u>001</u>	
<u>AB</u>	+		<u>10MG</u>	<u>N017078</u>	<u>002</u>	
<u>AB</u>	+		<u>15MG</u>	<u>N017078</u>	<u>003</u>	

DEXTROAMPHETAMINE SULFATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>5MG</u>	<u>A203901</u>	<u>001</u>	Nov 30, 2012
<u>AB</u>			<u>10MG</u>	<u>A203901</u>	<u>002</u>	Nov 30, 2012
<u>AB</u>			<u>15MG</u>	<u>A203901</u>	<u>003</u>	Nov 30, 2012
<u>AB</u>		SPECGX LLC	<u>5MG</u>	<u>A076353</u>	<u>001</u>	May 06, 2003
<u>AB</u>			<u>10MG</u>	<u>A076353</u>	<u>002</u>	May 06, 2003
<u>AB</u>			<u>15MG</u>	<u>A076353</u>	<u>003</u>	May 06, 2003
<u>AB</u>		STRIDES PHARMA	<u>5MG</u>	<u>A205673</u>	<u>001</u>	Oct 31, 2017
<u>AB</u>			<u>10MG</u>	<u>A205673</u>	<u>002</u>	Oct 31, 2017
<u>AB</u>			<u>15MG</u>	<u>A205673</u>	<u>003</u>	Oct 31, 2017

SOLUTION; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	!	PRASCO	<u>5MG/5ML</u>	<u>A040776</u>	<u>001</u>	Jan 29, 2008
<u>AA</u>		TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A203644</u>	<u>001</u>	May 29, 2013

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>		ARBOR PHARMS LLC	<u>2.5MG</u>	<u>A090533</u>	<u>001</u>	Oct 25, 2011
<u>AA</u>			<u>5MG</u>	<u>A090533</u>	<u>002</u>	Oct 25, 2011
<u>AA</u>			<u>7.5MG</u>	<u>A090533</u>	<u>003</u>	Oct 25, 2011
<u>AA</u>			<u>10MG</u>	<u>A090533</u>	<u>004</u>	Oct 25, 2011
<u>AA</u>			<u>15MG</u>	<u>A090533</u>	<u>005</u>	Oct 25, 2011
<u>AA</u>			<u>20MG</u>	<u>A090533</u>	<u>006</u>	Oct 25, 2011
<u>AA</u>			<u>30MG</u>	<u>A090533</u>	<u>007</u>	Oct 25, 2011
<u>AA</u>		AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A202893</u>	<u>001</u>	Jul 31, 2013
<u>AA</u>			<u>10MG</u>	<u>A202893</u>	<u>002</u>	Jul 31, 2013
<u>AA</u>		AVANTHI INC	<u>5MG</u>	<u>A203548</u>	<u>001</u>	Nov 23, 2015
<u>AA</u>			<u>10MG</u>	<u>A203548</u>	<u>002</u>	Nov 23, 2015
<u>AA</u>		BARR	<u>5MG</u>	<u>A040361</u>	<u>001</u>	Jan 31, 2001
<u>AA</u>	!		<u>10MG</u>	<u>A040361</u>	<u>002</u>	Jan 31, 2001
<u>AA</u>		NOVEL LABS INC	<u>5MG</u>	<u>A204330</u>	<u>001</u>	Mar 16, 2016
<u>AA</u>			<u>10MG</u>	<u>A204330</u>	<u>002</u>	Mar 16, 2016
<u>AA</u>		NUVO PHARM	<u>5MG</u>	<u>A210059</u>	<u>001</u>	Oct 18, 2017
<u>AA</u>			<u>10MG</u>	<u>A210059</u>	<u>002</u>	Oct 18, 2017
<u>AA</u>		SPECGX LLC	<u>5MG</u>	<u>A040436</u>	<u>001</u>	Jan 29, 2002
<u>AA</u>			<u>10MG</u>	<u>A040436</u>	<u>002</u>	Jan 29, 2002
<u>AA</u>		TRIS PHARMA INC	<u>5MG</u>	<u>A206095</u>	<u>001</u>	Aug 18, 2022
<u>AA</u>			<u>10MG</u>	<u>A206095</u>	<u>002</u>	Aug 18, 2022
<u>AA</u>		WINDER LABS LLC	<u>2.5MG</u>	<u>A212160</u>	<u>001</u>	Jun 07, 2021
<u>AA</u>			<u>5MG</u>	<u>A212160</u>	<u>002</u>	Jun 07, 2021
<u>AA</u>			<u>7.5MG</u>	<u>A212160</u>	<u>003</u>	Jun 07, 2021
<u>AA</u>			<u>10MG</u>	<u>A212160</u>	<u>004</u>	Jun 07, 2021
<u>AA</u>			<u>15MG</u>	<u>A212160</u>	<u>005</u>	Jun 07, 2021
<u>AA</u>			<u>20MG</u>	<u>A212160</u>	<u>006</u>	Jun 07, 2021
<u>AA</u>			<u>30MG</u>	<u>A212160</u>	<u>007</u>	Jun 07, 2021

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE DM

<u>AA</u>	!	SLATE RUN PHARMA	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A040649</u>	<u>001</u>	Feb 14, 2006
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PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

<u>AA</u>		AMNEAL PHARMS	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A090575</u>	<u>001</u>	Feb 08, 2011
<u>AA</u>		TRIS PHARMA INC	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A091687</u>	<u>001</u>	Jun 28, 2012

PROMETHAZINE W/ DEXTROMETHORPHAN

<u>AA</u>		WOCKHARDT BIO AG	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A088864</u>	<u>001</u>	Jan 04, 1985
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DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE

CAPSULE; ORAL

DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>20MG; 10MG</u>	<u>A202934</u>	<u>001</u>	Oct 10, 2017
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NUDEXTA

<u>AB</u>	+	AVANIR PHARMS	<u>20MG; 10MG</u>	<u>N021879</u>	<u>001</u>	Oct 29, 2010
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PRESCRIPTION DRUG PRODUCT LISTDEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>10GM/100ML</u>	<u>N019626 004</u>	Feb 02, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N016694 001</u>	
<u>AP</u>		FRESENIUS KABI USA	<u>10GM/100ML</u>	<u>A209448 001</u>	Jul 16, 2018
<u>AP</u>	+	ICU MEDICAL INC	<u>10GM/100ML</u>	<u>N018080 001</u>	

DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>50MG/ML</u>	<u>N016730 002</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N016730 001</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N019626 002</u>	Feb 02, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N016673 003</u>	Oct 30, 1985
<u>AP</u>	+		<u>50MG/ML</u>	<u>N020179 002</u>	Dec 07, 1992
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N016673 001</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N020179 001</u>	Dec 07, 1992
<u>AP</u>		FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A207449 001</u>	Oct 21, 2016
<u>AP</u>	+	HOSPIRA	<u>5GM/100ML</u>	<u>N019466 001</u>	Jul 15, 1985
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N019479 001</u>	Sep 17, 1985
<u>AP</u>	+	ICU MEDICAL INC	<u>50MG/ML</u>	<u>N016367 002</u>	

DEXTROSE 50%

<u>AP</u>	+	HOSPIRA	<u>500MG/ML</u>	<u>N019445 001</u>	Jun 03, 1986
<u>AP</u>		INTL MEDICATION SYS	<u>500MG/ML</u>	<u>A203451 001</u>	Mar 26, 2021

DEXTROSE 50% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>50GM/100ML</u>	<u>N020047 001</u>	Jul 02, 1991
<u>AP</u>	+	ICU MEDICAL INC	<u>50GM/100ML</u>	<u>N018563 001</u>	Mar 23, 1982

DEXTROSE 70% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>70GM/100ML</u>	<u>N019626 005</u>	Feb 18, 2015
<u>AP</u>	+	BAXTER HLTHCARE	<u>70GM/100ML</u>	<u>N020047 003</u>	Jul 02, 1991
<u>AP</u>	+	ICU MEDICAL INC	<u>70GM/100ML</u>	<u>N018561 001</u>	Mar 23, 1982
<u>AP</u>	+		<u>70GM/100ML</u>	<u>N019893 001</u>	Dec 26, 1989

DEXTROSE 20% IN PLASTIC CONTAINER

+	ICU MEDICAL INC	20GM/100ML	N018564 001	Mar 23, 1982
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DEXTROSE 25%

+	HOSPIRA	250MG/ML	N019445 002	Nov 23, 1998
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DEXTROSE 30% IN PLASTIC CONTAINER

+	ICU MEDICAL INC	30GM/100ML	N019345 001	Jan 26, 1985
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DEXTROSE 40% IN PLASTIC CONTAINER

+	ICU MEDICAL INC	40GM/100ML	N018562 001	Mar 23, 1982
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DEXTROSE; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC	5GM/100ML; 21MG/100ML; 128MG/100ML; 234MG/100ML	N017610 001
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DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION

ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML; 31MG/100ML; 130MG/100ML; 26MG/100ML; 320MG/100ML	N019873 001	Jun 10, 1993
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DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER

BAXTER HLTHCARE	5GM/100ML; 31MG/100ML; 141MG/100ML; 20MG/100ML; 12MG/100ML; 260MG/100ML	N017484 001
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DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC	5GM/100ML; 30MG/100ML; 141MG/100ML; 15MG/100ML; 260MG/100ML; 25MG/100ML	N019513 001	May 08, 1986
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DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC	5GM/100ML; 30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML	N017609 001
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PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075%

AP	FRESENIUS KABI USA	5GM/100ML;75MG/100ML	A212346 001	Sep 10, 2020
		<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER</u>		
AP	+ BAXTER HLTHCARE	5GM/100ML;75MG/100ML	N017634 004	
		<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15%</u>		
AP	FRESENIUS KABI USA	5GM/100ML;150MG/100ML	A212346 002	Sep 10, 2020
		<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER</u>		
AP	+ BAXTER HLTHCARE	5GM/100ML;150MG/100ML	N017634 001	
		<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>		
AP	+ BAXTER HLTHCARE	5GM/100ML;300MG/100ML	N017634 002	
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML	N019699 004	Sep 29, 1989
		<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;300MG/100ML	N019699 006	Sep 29, 1989
		<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER</u>		
	+ BAXTER HLTHCARE	5GM/100ML;224MG/100ML	N017634 003	
		<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
	ICU MEDICAL INC	5GM/100ML;149MG/100ML	N018371 001	

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ

AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;200MG/100ML	N018037 006	Apr 13, 1982
AP		5GM/100ML;150MG/100ML;200MG/100ML	N018037 007	Apr 13, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)</u>		
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;200MG/100ML	N018037 004	
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ</u>		
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;200MG/100ML	N018037 008	Apr 13, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)</u>		
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;200MG/100ML	N018037 001	
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ</u>		
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;200MG/100ML	N018037 005	Apr 13, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ</u>		
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;200MG/100ML	N018037 009	Apr 13, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ</u>		
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;200MG/100ML	N018037 002	
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K)</u>		
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;200MG/100ML	N018037 003	
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N018629 005	Mar 23, 1982
AP		5GM/100ML;150MG/100ML;330MG/100ML	N018629 002	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N018629 003	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;330MG/100ML	N018629 004	Mar 23, 1982
AP		5GM/100ML;300MG/100ML;330MG/100ML	N018629 006	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N018629 007	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;330MG/100ML	N018629 008	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N018629 001	Mar 23, 1982
		<u>DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER</u>		
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N018008 010	
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;200MG/100ML	N019630 008	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;330MG/100ML	N019630 014	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;450MG/100ML	N019630 020	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;75MG/100ML;900MG/100ML	N019630 026	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;200MG/100ML	N019630 010	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;330MG/100ML	N019630 016	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;450MG/100ML	N019630 022	Feb 17, 1988
		<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>		
AP	B BRAUN	5GM/100ML;150MG/100ML;900MG/100ML	N019630 028	Feb 17, 1988

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	5GM/100ML;300MG/100ML;200MG/100ML	N019630 012	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	5GM/100ML;300MG/100ML;330MG/100ML	N019630 018	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	5GM/100ML;300MG/100ML;450MG/100ML	N019630 024	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	5GM/100ML;300MG/100ML;900MG/100ML	N019630 030	Feb 17, 1988
<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER</u>				
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;225MG/100ML	A212348 001	Jul 30, 2021
<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>				
AP	FRESENIUS KABI USA	5GM/100ML;74.5MG/100ML;450MG/100ML	A213523 001	Mar 09, 2021
AP		5GM/100ML;149MG/100ML;450MG/100ML	A213523 005	Oct 11, 2022
<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;450MG/100ML	N018008 005	Apr 28, 1982
AP		5GM/100ML;150MG/100ML;450MG/100ML	N018008 006	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;450MG/100ML	N018362 009	Jul 05, 1983
AP	+!	5GM/100ML;149MG/100ML;450MG/100ML	N018362 005	Mar 28, 1988
<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER</u>				
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;225MG/100ML	A212348 002	Jul 30, 2021
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;225MG/100ML	N018365 001	
<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>				
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;450MG/100ML	A213523 002	Mar 09, 2021
<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;450MG/100ML	N018008 007	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;450MG/100ML	N018362 010	Jul 05, 1983
<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%</u>				
AP	FRESENIUS KABI USA	5GM/100ML;149MG/100ML;900MG/100ML	A213445 001	Mar 09, 2021
<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;900MG/100ML	N019308 005	Apr 05, 1985
AP	+! ICU MEDICAL INC	5GM/100ML;149MG/100ML;900MG/100ML	N019691 005	Mar 24, 1988
<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>				
AP	FRESENIUS KABI USA	5GM/100ML;224MG/100ML;450MG/100ML	A213523 003	Mar 09, 2021
<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;450MG/100ML	N018008 008	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;224MG/100ML;450MG/100ML	N018362 002	
<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>				
AP	FRESENIUS KABI USA	5GM/100ML;298MG/100ML;450MG/100ML	A213523 004	Mar 09, 2021
<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N018008 009	Apr 28, 1982
AP	+! ICU MEDICAL INC	5GM/100ML;298MG/100ML;450MG/100ML	N018362 003	
<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%</u>				
AP	FRESENIUS KABI USA	5GM/100ML;298MG/100ML;900MG/100ML	A213445 002	Mar 09, 2021
<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;900MG/100ML	N019308 007	Apr 05, 1985
AP	+! ICU MEDICAL INC	5GM/100ML;298MG/100ML;900MG/100ML	N019691 009	Mar 24, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;37MG/100ML;200MG/100ML	N019630 031	Feb 17, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;37MG/100ML;450MG/100ML	N019630 037	Feb 17, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;37MG/100ML;900MG/100ML	N019630 043	Feb 17, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER				
	B BRAUN	5GM/100ML;37MG/100ML;110MG/100ML	N019630 001	Feb 17, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER				
	B BRAUN	5GM/100ML;37MG/100ML;200MG/100ML	N019630 007	Feb 17, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER				
	B BRAUN	5GM/100ML;37MG/100ML;330MG/100ML	N019630 013	Feb 17, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
	B BRAUN	5GM/100ML;37MG/100ML;450MG/100ML	N019630 019	Feb 17, 1988
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	B BRAUN	5GM/100ML;37MG/100ML;900MG/100ML	N019630 025	Feb 17, 1988
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;75MG/100ML;200MG/100ML	N019630 032	Feb 17, 1988
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;75MG/100ML;450MG/100ML	N019630 038	Feb 17, 1988
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;75MG/100ML;900MG/100ML	N019630 044	Feb 17, 1988
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
	B BRAUN	3.3GM/100ML;75MG/100ML;300MG/100ML	N019630 049	May 07, 1992

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;75MG/100ML;110MG/100ML	N019630 002	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;110MG/100ML;200MG/100ML	N019630 033	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;110MG/100ML;450MG/100ML	N019630 039	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;110MG/100ML;900MG/100ML	N019630 045	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	3.3GM/100ML;110MG/100ML;300MG/100ML	N019630 050	May 07, 1992
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML;110MG/100ML	N019630 003	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML;200MG/100ML	N019630 009	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML;330MG/100ML	N019630 015	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML;450MG/100ML	N019630 021	Feb 17, 1988
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML;900MG/100ML	N019630 027	Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;150MG/100ML;200MG/100ML	N019630 034	Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;150MG/100ML;450MG/100ML	N019630 040	Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;150MG/100ML;900MG/100ML	N019630 046	Feb 17, 1988
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	3.3GM/100ML;150MG/100ML;300MG/100ML	N019630 051	May 07, 1992
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;150MG/100ML;110MG/100ML	N019630 004	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;220MG/100ML;200MG/100ML	N019630 035	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;220MG/100ML;450MG/100ML	N019630 041	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;220MG/100ML;900MG/100ML	N019630 047	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	3.3GM/100ML;220MG/100ML;300MG/100ML	N019630 052	May 07, 1992
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;110MG/100ML	N019630 005	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;200MG/100ML	N019630 011	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;330MG/100ML	N019630 017	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;450MG/100ML	N019630 023	Feb 17, 1988
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML;900MG/100ML	N019630 029	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;300MG/100ML;200MG/100ML	N019630 036	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;300MG/100ML;450MG/100ML	N019630 042	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;300MG/100ML;900MG/100ML	N019630 048	Feb 17, 1988
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	3.3GM/100ML;300MG/100ML;300MG/100ML	N019630 053	May 07, 1992
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;300MG/100ML;110MG/100ML	N019630 006	Feb 17, 1988
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;75MG/100ML;900MG/100ML	N019308 004	Apr 05, 1985
	5GM/100ML;150MG/100ML;900MG/100ML	N019308 002	Apr 05, 1985
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;300MG/100ML;900MG/100ML	N019308 003	Apr 05, 1985
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;224MG/100ML;900MG/100ML	N019308 006	Apr 05, 1985
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;150MG/100ML;450MG/100ML	N018008 004	
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;150MG/100ML;900MG/100ML	N019308 001	Apr 05, 1985

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

AP	B BRAUN	<u>2.5GM/100ML;450MG/100ML</u>	<u>N019631 004</u>	Feb 24, 1988
AP	+! BAXTER HLTHCARE	<u>2.5GM/100ML;450MG/100ML</u>	<u>N016697 001</u>	
AP	FRESENIUS KABI USA	<u>2.5GM/100ML;450MG/100ML</u>	<u>A211190 001</u>	Dec 20, 2019
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;200MG/100ML</u>	<u>N019631 007</u>	Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.225%</u>				
AP	FRESENIUS KABI USA	<u>5GM/100ML;225MG/100ML</u>	<u>A211221 001</u>	Sep 15, 2020
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER</u>				
AP	+! ICU MEDICAL INC	<u>5GM/100ML;225MG/100ML</u>	<u>N017606 001</u>	
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>				
AP	FRESENIUS KABI USA	<u>5GM/100ML;300MG/100ML</u>	<u>A211194 001</u>	Aug 26, 2020
AP	+! ICU MEDICAL INC	<u>5GM/100ML;300MG/100ML</u>	<u>N017799 001</u>	
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;330MG/100ML</u>	<u>N019631 008</u>	Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45%</u>				
AP	FRESENIUS KABI USA	<u>5GM/100ML;450MG/100ML</u>	<u>A211276 001</u>	Sep 15, 2020
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;450MG/100ML</u>	<u>N019631 009</u>	Feb 24, 1988
AP	+ ICU MEDICAL INC	<u>5GM/100ML;450MG/100ML</u>	<u>N017607 001</u>	
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9%</u>				
AP	FRESENIUS KABI USA	<u>5GM/100ML;900MG/100ML</u>	<u>A211211 001</u>	Sep 14, 2020
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;900MG/100ML</u>	<u>N019631 010</u>	Feb 24, 1988
AP	+! ICU MEDICAL INC	<u>5GM/100ML;900MG/100ML</u>	<u>N017585 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;200MG/100ML</u>	<u>N016689 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;330MG/100ML</u>	<u>N016687 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;450MG/100ML</u>	<u>N016683 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;900MG/100ML</u>	<u>N016678 001</u>	
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;110MG/100ML	N019631 011	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;200MG/100ML	N019631 012	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;330MG/100ML	N019631 013	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;450MG/100ML	N019631 014	Feb 24, 1988
<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;900MG/100ML	N019631 015	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;110MG/100ML	N019631 001	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;200MG/100ML	N019631 002	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;330MG/100ML	N019631 003	Feb 24, 1988
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
	B BRAUN	2.5GM/100ML;900MG/100ML	N019631 005	Feb 24, 1988
<u>DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>				
	B BRAUN	3.3GM/100ML;300MG/100ML	N019631 016	Jan 19, 1990
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u>				
	B BRAUN	5GM/100ML;110MG/100ML	N019631 006	Feb 24, 1988

DIATRIZOATE MEGLUMINE

SOLUTION; URETHRAL

CYSTOGRAFIN

+ BRACCO 30%

N010040 018

CYSTOGRAFIN DILUTE

+ BRACCO 18%

N010040 022 Nov 09, 1982

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

SOLUTION; ORAL, RECTAL

DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM

AA	ANDA REPOSITORY	<u>66%;10%</u>	<u>A214201 001</u>	Jun 27, 2022
AA	ANNORA PHARMA	<u>66%;10%</u>	<u>A215049 001</u>	Nov 17, 2023
<u>GASTROGRAFIN</u>				
AA	+! BRACCO	<u>66%;10%</u>	<u>N011245 003</u>	

PRESCRIPTION DRUG PRODUCT LIST

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

SOLUTION; ORAL, RECTAL

MD-GASTROVIEW

AA	LIEBEL-FLARSHEIM	66%;10%	A087388	001	
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DIAZEPAM

CONCENTRATE; ORAL

DIAZEPAM

AA	CHARTWELL MOLECULAR	5MG/ML	A204433	001	Apr 14, 2014
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DIAZEPAM INTENSOL

AA	! HIKMA	5MG/ML	A071415	001	Apr 03, 1987
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GEL; RECTAL

DIASTAT ACUDIAL

AB	+! BAUSCH	10MG/2ML (5MG/ML)	N020648	007	Sep 15, 2005
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AB	+!	20MG/4ML (5MG/ML)	N020648	006	Sep 15, 2005
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DIAZEPAM

AB	NOVEL LABS INC	10MG/2ML (5MG/ML)	A091076	001	May 30, 2023
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AB		20MG/4ML (5MG/ML)	A091076	002	May 30, 2023
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DIASTAT

+!	BAUSCH	2.5MG/0.5ML (5MG/ML)	N020648	001	Jul 29, 1997
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INJECTABLE; INJECTION

DIAZEPAM

AP	! BELOTECA	10MG/2ML (5MG/ML)	A210363	001	Mar 18, 2019
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AP		50MG/10ML (5MG/ML)	A211998	001	Dec 26, 2019
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AP	FRESENIUS KABI USA	10MG/2ML (5MG/ML)	A214745	001	Nov 10, 2022
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AP	HIKMA	5MG/ML	A070311	001	Dec 16, 1985
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AP		10MG/2ML (5MG/ML)	A070313	001	Dec 16, 1985
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AP	HOSPIRA	10MG/2ML (5MG/ML)	A072079	001	Dec 20, 1988
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AP	!	50MG/10ML (5MG/ML)	A071583	001	Oct 13, 1987
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SOLUTION; ORAL

DIAZEPAM

AA	CHARTWELL MOLECULAR	5MG/5ML	A206477	001	Jun 24, 2016
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AA	! HIKMA	5MG/5ML	A070928	001	Apr 03, 1987
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SPRAY; NASAL

VALTOCO

+	NEURELIS INC	5MG/SPRAY	N211635	001	Jan 10, 2020
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+		7.5MG/SPRAY	N211635	002	Jan 10, 2020
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+!		10MG/SPRAY	N211635	003	Jan 10, 2020
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TABLET; ORAL

DIAZEPAM

AB	AUROBINDO PHARMA LTD	2MG	A217843	001	Dec 14, 2023
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AB		5MG	A217843	002	Dec 14, 2023
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AB		10MG	A217843	003	Dec 14, 2023
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AB	DR REDDYS LABS SA	2MG	A071134	001	Feb 03, 1987
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AB		5MG	A071135	001	Feb 03, 1987
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AB		10MG	A071136	001	Feb 03, 1987
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AB	IVAX SUB TEVA PHARMS	2MG	A071307	001	Dec 10, 1986
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AB		5MG	A071321	001	Dec 10, 1986
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AB		10MG	A071322	001	Dec 10, 1986
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AB	MYLAN	2MG	A070325	002	Sep 04, 1985
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AB		5MG	A070325	003	Sep 04, 1985
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AB		10MG	A070325	001	Sep 04, 1985
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AB	NUVO PHARM	10MG	A070464	001	Feb 25, 1986
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AB	STRIDES PHARMA	2MG	A077749	001	Mar 31, 2006
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AB		5MG	A077749	002	Mar 31, 2006
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AB		10MG	A077749	003	Mar 31, 2006
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VALIUM

AB	+ WAYLIS THERAP	2MG	N013263	002	
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AB	+	5MG	N013263	004	
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AB	+!	10MG	N013263	006	
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DIAZOXIDE

SUSPENSION; ORAL

DIAZOXIDE

AB	E5 PHARMA INC	50MG/ML	A211050	001	Dec 20, 2019
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AB	NOVITIUM PHARMA	50MG/ML	A210799	001	Jul 08, 2020
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PROGLYCEM

AB	+!	TEVA BRANDED PHARM	50MG/ML	N017453	001
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PRESCRIPTION DRUG PRODUCT LIST

DICHLORPHENAMIDE

TABLET; ORAL

DICHLORPHENAMIDE

AB	TORRENT	50MG	A215924 001	Dec 29, 2022
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KEVEYIS

AB	+! XERIS	50MG	N011366 002	Aug 07, 2015
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DICLOFENAC EPOLAMINE

SYSTEM; TOPICAL

FLECTOR

	+! IBSA	1.3%	N021234 001	Jan 31, 2007
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LICART

	+! IBSA INST BIO	1.3%	N206976 001	Dec 19, 2018
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DICLOFENAC POTASSIUM

CAPSULE; ORAL

DICLOFENAC POTASSIUM

AB	AUROBINDO PHARMA LTD	25MG	A213875 001	Oct 19, 2021
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AB	BIONPHARMA	25MG	A204648 001	Feb 23, 2016
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AB	STRIDES PHARMA	25MG	A210078 001	Dec 03, 2019
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ZIPSOR

AB	+! ASSERTIO	25MG	N022202 001	Jun 16, 2009
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FOR SOLUTION; ORAL

CAMBIA

AB	+! ASSERTIO	50MG	N022165 001	Jun 17, 2009
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DICLOFENAC POTASSIUM

AB	ALKEM LABS LTD	50MG	A216635 001	Jul 20, 2022
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AB	ANNORA PHARMA	50MG	A215375 001	Mar 04, 2022
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AB	PAR FORM	50MG	A202964 001	May 02, 2016
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TABLET; ORAL

CATAFLAM

AB	AMICI	25MG	A076561 002	Jul 21, 2021
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AB		50MG	A076561 001	Mar 18, 2004
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DICLOFENAC POTASSIUM

AB	NOVAST LABS	50MG	A215585 001	Oct 08, 2021
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AB	! RK PHARMA	50MG	A075463 001	Jul 26, 1999
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AB	RUBICON	25MG	A075229 002	Sep 16, 2021
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AB		50MG	A075229 001	Nov 20, 1998
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AB	SENORES PHARMS	25MG	A215787 002	Nov 09, 2023
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AB		50MG	A215787 001	Mar 15, 2023
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AB	TEVA	50MG	A075219 001	Aug 06, 1998
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AB	UMEDICA	50MG	A215750 001	May 11, 2022
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DICLOFENAC SODIUM

GEL; TOPICAL

DICLOFENAC SODIUM

AB	ACTAVIS MID ATLANTIC	3%	A206493 001	Dec 02, 2015
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AB	ALEMBIC	3%	A212351 001	Jul 27, 2022
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AB	AMNEAL	3%	A200936 001	Oct 28, 2013
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AB	GLENMARK PHARMS LTD	3%	A208301 001	Sep 13, 2016
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AB	PADAGIS ISRAEL	3%	A210893 001	Jul 27, 2018
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AB	! TARO	3%	A206298 001	Apr 28, 2016
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	CIPLA	1%	A209903 001	Aug 03, 2018
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SOLUTION; TOPICAL

DICLOFENAC SODIUM

AB	ALEMBIC	2%	A212506 001	Nov 29, 2022
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AB	AMNEAL	2%	A208198 001	Aug 18, 2022
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AB	APOTEX	2%	A207714 001	May 06, 2022
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AB	AUROLIFE PHARMA LLC	2%	A213040 001	Feb 03, 2023
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AB	LUPIN PHARMS	2%	A208021 001	Sep 20, 2022
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AB	TARO	2%	A208098 001	Sep 29, 2022
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PENNSAID

AB	+! HORIZON	2%	N204623 001	Jan 16, 2014
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DICLOFENAC SODIUM

AT	AMNEAL PHARMS	1.5%	A206116 001	Sep 02, 2016
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AT	CADILA	1.5%	A206411 001	Apr 17, 2018
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AT	EPIC PHARMA LLC	1.5%	A206655 001	Jan 28, 2021
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AT	LUPIN LTD	1.5%	A204132 001	Aug 20, 2015
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AT	NOVEL LABS INC	1.5%	A205878 001	Dec 09, 2015
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AT	! TARO	1.5%	A203818 001	Nov 26, 2014
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AT	WATSON LABS INC	1.5%	A202852 001	Nov 24, 2014
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PRESCRIPTION DRUG PRODUCT LIST

DICLOFENAC SODIUM

SOLUTION/DROPS;OPHTHALMIC

DICLOFENAC SODIUM

<u>AT</u>	ALTAIRE PHARMS INC	<u>0.1%</u>	<u>A203383</u>	<u>001</u>	Nov 16, 2015
<u>AT</u>	! BAUSCH AND LOMB	<u>0.1%</u>	<u>A078792</u>	<u>001</u>	Dec 28, 2007
<u>AT</u>	RISING	<u>0.1%</u>	<u>A078553</u>	<u>001</u>	Dec 28, 2007
<u>AT</u>	RUBICON	<u>0.1%</u>	<u>A077600</u>	<u>001</u>	Nov 13, 2008
<u>AT</u>	SANDOZ	<u>0.1%</u>	<u>A078031</u>	<u>001</u>	Feb 06, 2008

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

<u>AB</u>	ACTAVIS ELIZABETH	<u>50MG</u>	<u>A074514</u>	<u>001</u>	Mar 26, 1996
<u>AB</u>		<u>75MG</u>	<u>A074514</u>	<u>002</u>	Mar 26, 1996
<u>AB</u>	CARLSBAD	<u>25MG</u>	<u>A075185</u>	<u>002</u>	Nov 13, 1998
<u>AB</u>		<u>50MG</u>	<u>A075185</u>	<u>003</u>	Nov 13, 1998
<u>AB</u>		<u>75MG</u>	<u>A075185</u>	<u>001</u>	Nov 13, 1998
<u>AB</u>	RUBICON	<u>25MG</u>	<u>A216548</u>	<u>001</u>	May 11, 2023
<u>AB</u>		<u>50MG</u>	<u>A216548</u>	<u>002</u>	May 11, 2023
<u>AB</u>		<u>75MG</u>	<u>A216548</u>	<u>003</u>	May 11, 2023
<u>AB</u>	! UNIQUE	<u>25MG</u>	<u>A090066</u>	<u>001</u>	Dec 01, 2010
<u>AB</u>	!	<u>50MG</u>	<u>A090066</u>	<u>002</u>	Dec 01, 2010
<u>AB</u>	!	<u>75MG</u>	<u>A077863</u>	<u>003</u>	Jun 08, 2007

TABLET, EXTENDED RELEASE;ORAL

DICLOFENAC SODIUM

<u>AB</u>	! DEXCEL LTD	<u>100MG</u>	<u>A076201</u>	<u>001</u>	Nov 06, 2002
<u>AB</u>	RICONPHARMA LLC	<u>100MG</u>	<u>A216275</u>	<u>001</u>	Sep 23, 2022
<u>AB</u>	VPNA	<u>100MG</u>	<u>A075492</u>	<u>001</u>	Feb 11, 2000

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

ARTHROTEC

<u>AB</u>	+ PFIZER	<u>50MG;0.2MG</u>	<u>N020607</u>	<u>001</u>	Dec 24, 1997
<u>AB</u>	+!	<u>75MG;0.2MG</u>	<u>N020607</u>	<u>002</u>	Dec 24, 1997

DICLOFENAC SODIUM AND MISOPROSTOL

<u>AB</u>	ACTAVIS LABS FL INC	<u>50MG;0.2MG</u>	<u>A201089</u>	<u>001</u>	Jul 09, 2012
<u>AB</u>		<u>75MG;0.2MG</u>	<u>A201089</u>	<u>002</u>	Jul 09, 2012
<u>AB</u>	AMNEAL PHARMS	<u>50MG;0.2MG</u>	<u>A203995</u>	<u>001</u>	Nov 25, 2016
<u>AB</u>		<u>75MG;0.2MG</u>	<u>A203995</u>	<u>002</u>	Nov 25, 2016
<u>AB</u>	MICRO LABS	<u>50MG;0.2MG</u>	<u>A204355</u>	<u>001</u>	Jul 15, 2021
<u>AB</u>		<u>75MG;0.2MG</u>	<u>A204355</u>	<u>002</u>	Jul 15, 2021
<u>AB</u>	SANDOZ	<u>50MG;0.2MG</u>	<u>A200158</u>	<u>001</u>	May 09, 2013
<u>AB</u>		<u>75MG;0.2MG</u>	<u>A200158</u>	<u>002</u>	May 09, 2013
<u>AB</u>	YUNG SHIN PHARM	<u>50MG;0.2MG</u>	<u>A205143</u>	<u>001</u>	Feb 19, 2020
<u>AB</u>		<u>75MG;0.2MG</u>	<u>A205143</u>	<u>002</u>	Feb 19, 2020

DICLOXACILLIN SODIUM

CAPSULE;ORAL

DICLOXACILLIN SODIUM

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A216845</u>	<u>001</u>	Sep 23, 2022
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A216845</u>	<u>002</u>	Sep 23, 2022
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A062286</u>	<u>001</u>	Jun 03, 1982
<u>AB</u>	!	<u>EQ 500MG BASE</u>	<u>A062286</u>	<u>002</u>	Jun 03, 1982

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

DICYCLOMINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA USA	<u>10MG</u>	<u>A040319</u>	<u>001</u>	Sep 07, 1999
<u>AB</u>	COREPHARMA	<u>10MG</u>	<u>A216639</u>	<u>001</u>	Mar 24, 2023
<u>AB</u>	! LANNETT	<u>10MG</u>	<u>A084285</u>	<u>001</u>	
<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A217531</u>	<u>001</u>	Aug 30, 2023
<u>AB</u>	TWI PHARMS	<u>10MG</u>	<u>A217054</u>	<u>001</u>	Dec 27, 2022
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A085082</u>	<u>001</u>	Jun 19, 1986
<u>AB</u>	WEST WARD	<u>10MG</u>	<u>A040204</u>	<u>001</u>	Feb 28, 1997

INJECTABLE;INJECTION

BENTYL

<u>AP</u>	+! ABBVIE	<u>10MG/ML</u>	<u>N008370</u>	<u>001</u>	Oct 15, 1984
	<u>BENTYL PRESERVATIVE FREE</u>				
<u>AP</u>	+! ABBVIE	<u>10MG/ML</u>	<u>N008370</u>	<u>002</u>	Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

<u>AP</u>	AM REGENT	<u>10MG/ML</u>	<u>A208353</u>	<u>001</u>	Feb 17, 2017
<u>AP</u>	FOSUN PHARMA	<u>10MG/ML</u>	<u>A210979</u>	<u>001</u>	Jul 02, 2018
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A210257</u>	<u>001</u>	Jan 25, 2019
<u>AP</u>	NEXUS	<u>10MG/ML</u>	<u>A206468</u>	<u>001</u>	Feb 01, 2019
<u>AP</u>	SLATE RUN PHARMA	<u>10MG/ML</u>	<u>A207076</u>	<u>001</u>	Nov 02, 2018

PRESCRIPTION DRUG PRODUCT LIST

DICYCLOMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE)

AP	HIKMA	10MG/ML	A040465 001	Jun 30, 2003
	SYRUP; ORAL			

DICYCLOMINE HYDROCHLORIDE

AA	!	GENERICS	10MG/5ML	A040169 001	Mar 24, 2005
AA		HIKMA	10MG/5ML	A212286 001	May 22, 2020
AA		NOVITIUM PHARMA	10MG/5ML	A214721 001	Apr 23, 2021

TABLET; ORAL

DICYCLOMINE HYDROCHLORIDE

AB		AUROBINDO PHARMA USA	20MG	A040317 001	Sep 07, 1999
AB		BIONPHARMA	20MG	A217916 001	Aug 04, 2023
AB		COREPHARMA	20MG	A216760 001	Nov 28, 2022
AB		HIKMA PHARMS	20MG	A040161 001	Oct 01, 1996
AB		LANNETT	20MG	A040230 001	Feb 26, 1999
AB		RUBICON	20MG	A216736 001	Dec 14, 2022
AB		TWI PHARMS	20MG	A216782 001	Jun 01, 2023
AB	!	WATSON LABS	20MG	A085223 001	Jul 30, 1986

DIENOGEST; ESTRADIOL VALERATE

TABLET; ORAL

NATAZIA

+	!	BAYER HLTHCARE	N/A, 2MG, 3MG, N/A, N/A; 3MG, 2MG, 2MG, 1MG, N/A	N022252 001	May 06, 2010
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DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

DIETHYLPROPION HYDROCHLORIDE

AA		AVANTHI INC	25MG	A201212 001	Dec 22, 2010
AA	!	LANNETT CO INC	25MG	A200177 001	Jul 18, 2011
	TABLET, EXTENDED RELEASE; ORAL				
	DIETHYLPROPION HYDROCHLORIDE				
	!	LANNETT CO INC	75MG	A091680 001	Oct 24, 2011

DIFELIKEFALIN ACETATE

SOLUTION; INTRAVENOUS

KORSUVA

+	!	CARA THERAP	EQ 0.065MG BASE/1.3ML (EQ 0.05MG BASE/ML)	N214916 001	Aug 23, 2021
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DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

BX	!	ANI PHARMS	0.05%	A076263 001	Dec 20, 2002
BX	!	TARO	0.05%	A075508 001	Apr 24, 2000

OINTMENT; TOPICAL

DIFLORASONE DIACETATE

AB		AVONDALE PHARMS	0.05%	A075374 001	Apr 27, 1999
AB		RISING	0.05%	A207440 001	Feb 27, 2017
AB	!	TARO	0.05%	A075331 001	May 14, 1999
AB		THE J MOLNER	0.05%	A210753 001	Jun 12, 2018

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

AB		HERITAGE PHARMA	500MG	A202845 001	Mar 08, 2012
AB	!	TEVA	500MG	A073673 001	Jul 31, 1992
AB		ZYDUS PHARMS	500MG	A203547 001	Jun 16, 2017

DIFLUPREDNATE

EMULSION; OPHTHALMIC

DIFLUPREDNATE

AB		AMNEAL	0.05%	A211526 001	Nov 17, 2021
AB		CIPLA	0.05%	A211776 001	Aug 09, 2021
AB		DR REDDYS	0.05%	A214894 001	Nov 16, 2022

DUREZOL

AB	+	!	SANDOZ	0.05%	N022212 001	Jun 23, 2008
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DIGOXIN

ELIXIR; ORAL

DIGOXIN

AA		AMICI	0.05MG/ML	A215209 001	Mar 11, 2022	
AA	+	!	HIKMA	0.05MG/ML	N021648 001	Aug 26, 2004
AA		NOVITIUM PHARMA	0.05MG/ML	A213000 001	Oct 04, 2019	

PRESCRIPTION DRUG PRODUCT LIST

DIGOXIN

INJECTABLE; INJECTION

DIGOXIN

<u>AP</u>	SANDOZ	<u>0.25MG/ML</u>	<u>A040481</u>	<u>001</u>	Aug 21, 2003
<u>AP</u>	WEST-WARD PHARMS INT	<u>0.25MG/ML</u>	<u>A083391</u>	<u>001</u>	

LANOXIN

<u>AP</u>	+! COVIS	<u>0.25MG/ML</u>	<u>N009330</u>	<u>002</u>	
	LANOXIN PEDIATRIC +! COVIS	0.1MG/ML	N009330	004	

TABLET; ORAL

DIGOXIN

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.0625MG</u>	<u>A214982</u>	<u>001</u>	Feb 08, 2022
<u>AB</u>		<u>0.125MG</u>	<u>A214982</u>	<u>002</u>	Feb 08, 2022
<u>AB</u>		<u>0.25MG</u>	<u>A214982</u>	<u>003</u>	Feb 08, 2022
<u>AB</u>	HIKMA INTL PHARMS	<u>0.125MG</u>	<u>A077002</u>	<u>002</u>	Oct 30, 2007
<u>AB</u>		<u>0.25MG</u>	<u>A077002</u>	<u>001</u>	Oct 30, 2007
<u>AB</u>	IMPAX LABS	<u>0.125MG</u>	<u>A078556</u>	<u>001</u>	Jul 20, 2009
<u>AB</u>		<u>0.25MG</u>	<u>A078556</u>	<u>002</u>	Jul 20, 2009
<u>AB</u>	NOVITIUM PHARMA	<u>0.0625MG</u>	<u>A215307</u>	<u>003</u>	Aug 25, 2022
<u>AB</u>		<u>0.125MG</u>	<u>A215307</u>	<u>001</u>	Nov 22, 2021
<u>AB</u>		<u>0.25MG</u>	<u>A215307</u>	<u>002</u>	Nov 22, 2021
<u>AB</u>	RISING	<u>0.125MG</u>	<u>A040282</u>	<u>001</u>	Dec 23, 1999
<u>AB</u>		<u>0.25MG</u>	<u>A040282</u>	<u>002</u>	Dec 23, 1999
<u>AB</u>	STEVENS J	<u>0.125MG</u>	<u>A076268</u>	<u>001</u>	Jul 26, 2002
<u>AB</u>		<u>0.25MG</u>	<u>A076268</u>	<u>002</u>	Jul 26, 2002
<u>AB</u>	SUN PHARM INDS INC	<u>0.125MG</u>	<u>A076363</u>	<u>001</u>	Jan 31, 2003
<u>AB</u>		<u>0.25MG</u>	<u>A076363</u>	<u>002</u>	Jan 31, 2003

LANOXIN

<u>AB</u>	+ CONCORDIA	<u>0.0625MG</u>	<u>N020405</u>	<u>001</u>	Sep 30, 1997
<u>AB</u>	+	<u>0.125MG</u>	<u>N020405</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+!	<u>0.25MG</u>	<u>N020405</u>	<u>004</u>	Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

DIHYDROERGOTAMINE MESYLATE

<u>AP</u>	GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A215623</u>	<u>001</u>	Aug 18, 2023
<u>AP</u>	HIKMA	<u>1MG/ML</u>	<u>A040453</u>	<u>001</u>	Jun 09, 2003
<u>AP</u>	! HIKMA PHARMS	<u>1MG/ML</u>	<u>A206621</u>	<u>001</u>	Sep 15, 2017
<u>AP</u>	PADAGIS US	<u>1MG/ML</u>	<u>A040475</u>	<u>001</u>	Apr 28, 2003
<u>AP</u>	PROVEPHARM SAS	<u>1MG/ML</u>	<u>A212046</u>	<u>001</u>	Jan 07, 2020
<u>AP</u>	SAGENT PHARMS INC	<u>1MG/ML</u>	<u>A207264</u>	<u>001</u>	Jul 11, 2018

SPRAY, METERED; NASAL

DIHYDROERGOTAMINE MESYLATE

<u>AB</u>	AMNEAL	<u>0.5MG/SPRAY</u>	<u>A214105</u>	<u>001</u>	Jan 04, 2022
<u>AB</u>	CIPLA	<u>0.5MG/SPRAY</u>	<u>A212907</u>	<u>001</u>	May 20, 2020
<u>AB</u>	HIKMA	<u>0.5MG/SPRAY</u>	<u>A211393</u>	<u>001</u>	Feb 28, 2020
<u>AB</u>	RUBICON	<u>0.5MG/SPRAY</u>	<u>A216881</u>	<u>001</u>	Jun 22, 2023

MIGRANAL

<u>AB</u>	+! BAUSCH TRUDHESA	<u>0.5MG/SPRAY</u>	<u>N020148</u>	<u>001</u>	Dec 08, 1997
	+! IMPEL PHARMS	0.725MG/SPRAY	N213436	001	Sep 02, 2021

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB1</u>	DR REDDYS LABS SA	<u>60MG</u>	<u>A215775</u>	<u>001</u>	May 05, 2022
<u>AB1</u>		<u>90MG</u>	<u>A215775</u>	<u>002</u>	May 05, 2022
<u>AB1</u>		<u>120MG</u>	<u>A215775</u>	<u>003</u>	May 05, 2022
<u>AB1</u>	GLENMARK PHARMS LTD	<u>60MG</u>	<u>A212317</u>	<u>001</u>	Mar 22, 2021
<u>AB1</u>		<u>90MG</u>	<u>A212317</u>	<u>002</u>	Mar 22, 2021
<u>AB1</u>		<u>120MG</u>	<u>A212317</u>	<u>003</u>	Mar 22, 2021
<u>AB1</u>	MYLAN	<u>60MG</u>	<u>A074910</u>	<u>001</u>	May 02, 1997
<u>AB1</u>		<u>90MG</u>	<u>A074910</u>	<u>002</u>	May 02, 1997
<u>AB1</u>	!	<u>120MG</u>	<u>A074910</u>	<u>003</u>	May 02, 1997
<u>AB1</u>	TWI PHARMS	<u>60MG</u>	<u>A217377</u>	<u>001</u>	Mar 01, 2023
<u>AB1</u>		<u>90MG</u>	<u>A217377</u>	<u>002</u>	Mar 01, 2023
<u>AB1</u>		<u>120MG</u>	<u>A217377</u>	<u>003</u>	Mar 01, 2023
<u>AB2</u>	ACCORD HLTHCARE	<u>120MG</u>	<u>A206997</u>	<u>001</u>	Apr 28, 2020
<u>AB2</u>		<u>180MG</u>	<u>A206997</u>	<u>002</u>	Apr 28, 2020
<u>AB2</u>		<u>240MG</u>	<u>A206997</u>	<u>003</u>	Apr 28, 2020
<u>AB2</u>	AMTA	<u>120MG</u>	<u>A216304</u>	<u>001</u>	Aug 08, 2022
<u>AB2</u>		<u>180MG</u>	<u>A216304</u>	<u>002</u>	Aug 08, 2022

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB2</u>		<u>240MG</u>	<u>A216304 003</u>	Aug 08, 2022
<u>AB2</u>	APOTEX	<u>120MG</u>	<u>A074943 003</u>	Dec 19, 2000
<u>AB2</u>		<u>180MG</u>	<u>A074943 002</u>	Dec 19, 2000
<u>AB2</u>	!	<u>240MG</u>	<u>A074943 001</u>	Aug 06, 1998

CARDIZEM CD

<u>AB3</u>	+	BAUSCH	<u>120MG</u>	<u>N020062 001</u>	Aug 10, 1992
<u>AB3</u>	+		<u>180MG</u>	<u>N020062 002</u>	Dec 27, 1991
<u>AB3</u>	+		<u>240MG</u>	<u>N020062 003</u>	Dec 27, 1991
<u>AB3</u>	+		<u>300MG</u>	<u>N020062 004</u>	Dec 27, 1991
<u>AB3</u>	+	!	<u>360MG</u>	<u>N020062 005</u>	Aug 24, 1999

CARTIA XT

<u>AB3</u>		ACTAVIS LABS FL INC	<u>120MG</u>	<u>A074752 002</u>	Jul 09, 1998
<u>AB3</u>			<u>180MG</u>	<u>A074752 001</u>	Jul 09, 1998
<u>AB3</u>			<u>240MG</u>	<u>A074752 003</u>	Jul 09, 1998
<u>AB3</u>			<u>300MG</u>	<u>A074752 004</u>	Jul 09, 1998

DILTIAZEM HYDROCHLORIDE

<u>AB3</u>		ACTAVIS ELIZABETH	<u>360MG</u>	<u>A202463 001</u>	Dec 07, 2012
<u>AB3</u>		NOVAST LABS	<u>120MG</u>	<u>A208783 001</u>	Jun 14, 2019
<u>AB3</u>			<u>180MG</u>	<u>A208783 002</u>	Jun 14, 2019
<u>AB3</u>			<u>240MG</u>	<u>A208783 003</u>	Jun 14, 2019
<u>AB3</u>			<u>300MG</u>	<u>A208783 004</u>	Jun 14, 2019
<u>AB3</u>			<u>360MG</u>	<u>A208783 005</u>	Jun 14, 2019
<u>AB3</u>		PAR PHARM	<u>120MG</u>	<u>A074984 001</u>	Dec 20, 1999
<u>AB3</u>			<u>180MG</u>	<u>A074984 002</u>	Dec 20, 1999
<u>AB3</u>			<u>240MG</u>	<u>A074984 003</u>	Dec 20, 1999
<u>AB3</u>			<u>300MG</u>	<u>A074984 004</u>	Dec 20, 1999
<u>AB3</u>		SUN PHARM	<u>120MG</u>	<u>A203023 001</u>	Jun 08, 2017
<u>AB3</u>			<u>180MG</u>	<u>A203023 002</u>	Jun 08, 2017
<u>AB3</u>			<u>240MG</u>	<u>A203023 003</u>	Jun 08, 2017
<u>AB3</u>			<u>300MG</u>	<u>A203023 004</u>	Jun 08, 2017
<u>AB3</u>			<u>360MG</u>	<u>A090492 005</u>	Oct 28, 2011
<u>AB3</u>			<u>360MG</u>	<u>A203023 005</u>	Jun 08, 2017
<u>AB3</u>		TWI PHARMS	<u>120MG</u>	<u>A205231 001</u>	Aug 30, 2018
<u>AB3</u>			<u>180MG</u>	<u>A205231 002</u>	Aug 30, 2018
<u>AB3</u>			<u>240MG</u>	<u>A205231 003</u>	Aug 30, 2018
<u>AB3</u>			<u>300MG</u>	<u>A205231 004</u>	Aug 30, 2018
<u>AB3</u>			<u>360MG</u>	<u>A205231 005</u>	Aug 30, 2018
<u>AB3</u>		VALEANT PHARMS NORTH	<u>120MG</u>	<u>A075116 001</u>	Dec 23, 1999
<u>AB3</u>			<u>180MG</u>	<u>A075116 002</u>	Dec 23, 1999
<u>AB3</u>			<u>240MG</u>	<u>A075116 003</u>	Dec 23, 1999
<u>AB3</u>			<u>300MG</u>	<u>A075116 004</u>	Dec 23, 1999
<u>AB3</u>		ZYDUS PHARMS	<u>120MG</u>	<u>A206534 001</u>	Aug 08, 2017
<u>AB3</u>			<u>180MG</u>	<u>A206534 002</u>	Aug 08, 2017
<u>AB3</u>			<u>240MG</u>	<u>A206534 003</u>	Aug 08, 2017
<u>AB3</u>			<u>300MG</u>	<u>A206534 004</u>	Aug 08, 2017
<u>AB3</u>			<u>360MG</u>	<u>A206534 005</u>	Aug 08, 2017
<u>AB4</u>		SANDOZ	<u>120MG</u>	<u>A091022 001</u>	Sep 28, 2012
<u>AB4</u>			<u>180MG</u>	<u>A091022 002</u>	Sep 28, 2012
<u>AB4</u>			<u>240MG</u>	<u>A091022 003</u>	Sep 28, 2012
<u>AB4</u>			<u>300MG</u>	<u>A091022 004</u>	Sep 28, 2012
<u>AB4</u>			<u>360MG</u>	<u>A091022 005</u>	Sep 28, 2012
<u>AB4</u>			<u>420MG</u>	<u>A091022 006</u>	Sep 28, 2012
<u>AB4</u>		SUN PHARM	<u>120MG</u>	<u>A090421 001</u>	Nov 15, 2010
<u>AB4</u>			<u>180MG</u>	<u>A090421 002</u>	Nov 15, 2010
<u>AB4</u>			<u>240MG</u>	<u>A090421 003</u>	Nov 15, 2010
<u>AB4</u>			<u>300MG</u>	<u>A090421 004</u>	Nov 15, 2010
<u>AB4</u>			<u>360MG</u>	<u>A090421 005</u>	Nov 15, 2010
<u>AB4</u>		ZYDUS PHARMS	<u>120MG</u>	<u>A206641 001</u>	Aug 11, 2017
<u>AB4</u>			<u>180MG</u>	<u>A206641 002</u>	Aug 11, 2017
<u>AB4</u>			<u>240MG</u>	<u>A206641 003</u>	Aug 11, 2017
<u>AB4</u>			<u>300MG</u>	<u>A206641 004</u>	Aug 11, 2017
<u>AB4</u>			<u>360MG</u>	<u>A206641 005</u>	Aug 11, 2017
<u>AB4</u>			<u>420MG</u>	<u>A206641 006</u>	Aug 11, 2017

TAZTIA XT

<u>AB4</u>		ACTAVIS LABS FL INC	<u>120MG</u>	<u>A075401 001</u>	Apr 10, 2003
<u>AB4</u>			<u>180MG</u>	<u>A075401 002</u>	Apr 10, 2003
<u>AB4</u>			<u>240MG</u>	<u>A075401 003</u>	Apr 10, 2003
<u>AB4</u>			<u>300MG</u>	<u>A075401 004</u>	Apr 10, 2003
<u>AB4</u>			<u>360MG</u>	<u>A075401 005</u>	Apr 10, 2003

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

TIAZAC

<u>AB4</u>	+	BAUSCH	<u>120MG</u>	<u>N020401</u>	<u>001</u>	Sep 11, 1995
<u>AB4</u>	+		<u>180MG</u>	<u>N020401</u>	<u>002</u>	Sep 11, 1995
<u>AB4</u>	+		<u>240MG</u>	<u>N020401</u>	<u>003</u>	Sep 11, 1995
<u>AB4</u>	+		<u>300MG</u>	<u>N020401</u>	<u>004</u>	Sep 11, 1995
<u>AB4</u>	+		<u>360MG</u>	<u>N020401</u>	<u>005</u>	Sep 11, 1995
<u>AB4</u>	+	!	<u>420MG</u>	<u>N020401</u>	<u>006</u>	Oct 16, 1998

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

<u>AP</u>		EUGIA PHARMA	<u>5MG/ML</u>	<u>A216552</u>	<u>001</u>	Mar 29, 2023
<u>AP</u>		HIKMA FARMACEUTICA	<u>5MG/ML</u>	<u>A202651</u>	<u>001</u>	Aug 09, 2012
<u>AP</u>		HOSPIRA	<u>5MG/ML</u>	<u>A074941</u>	<u>001</u>	Apr 15, 1998
<u>AP</u>		RISING	<u>5MG/ML</u>	<u>A075086</u>	<u>001</u>	Apr 09, 1998
<u>AP</u>	!	SAGENT	<u>5MG/ML</u>	<u>A074617</u>	<u>001</u>	Feb 28, 1996
<u>AP</u>		WEST-WARD PHARMS	<u>5MG/ML</u>	<u>A078538</u>	<u>001</u>	Dec 17, 2008
		INT				
	!	HOSPIRA	100MG/VIAL	A075853	001	Dec 17, 2002

TABLET; ORAL

CARDIZEM

<u>AB</u>	+	BAUSCH	<u>30MG</u>	<u>N018602</u>	<u>001</u>	Nov 05, 1982
<u>AB</u>	+		<u>60MG</u>	<u>N018602</u>	<u>002</u>	Nov 05, 1982
<u>AB</u>	+		<u>90MG</u>	<u>N018602</u>	<u>003</u>	Dec 08, 1986
<u>AB</u>	+	!	<u>120MG</u>	<u>N018602</u>	<u>004</u>	Dec 08, 1986

DILTIAZEM HYDROCHLORIDE

<u>AB</u>		SCIEGEN PHARMS INC	<u>30MG</u>	<u>A216521</u>	<u>001</u>	Sep 23, 2022
<u>AB</u>			<u>60MG</u>	<u>A216521</u>	<u>002</u>	Sep 23, 2022
<u>AB</u>			<u>90MG</u>	<u>A216521</u>	<u>003</u>	Sep 23, 2022
<u>AB</u>			<u>120MG</u>	<u>A216521</u>	<u>004</u>	Sep 23, 2022
<u>AB</u>		TEVA	<u>30MG</u>	<u>A074185</u>	<u>001</u>	May 31, 1995
<u>AB</u>			<u>60MG</u>	<u>A074185</u>	<u>002</u>	May 31, 1995
<u>AB</u>			<u>90MG</u>	<u>A074185</u>	<u>003</u>	May 31, 1995
<u>AB</u>			<u>120MG</u>	<u>A074185</u>	<u>004</u>	May 31, 1995

TABLET, EXTENDED RELEASE;ORAL

CARDIZEM LA

<u>AB</u>	+	BAUSCH	<u>120MG</u>	<u>N021392</u>	<u>001</u>	Feb 06, 2003
<u>AB</u>	+		<u>180MG</u>	<u>N021392</u>	<u>002</u>	Feb 06, 2003
<u>AB</u>	+		<u>240MG</u>	<u>N021392</u>	<u>003</u>	Feb 06, 2003
<u>AB</u>	+		<u>300MG</u>	<u>N021392</u>	<u>004</u>	Feb 06, 2003
<u>AB</u>	+		<u>360MG</u>	<u>N021392</u>	<u>005</u>	Feb 06, 2003
<u>AB</u>	+	!	<u>420MG</u>	<u>N021392</u>	<u>006</u>	Feb 06, 2003

DILTIAZEM HYDROCHLORIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>120MG</u>	<u>A077686</u>	<u>006</u>	Mar 15, 2010
<u>AB</u>			<u>180MG</u>	<u>A077686</u>	<u>005</u>	Mar 15, 2010
<u>AB</u>			<u>240MG</u>	<u>A077686</u>	<u>004</u>	Mar 15, 2010
<u>AB</u>			<u>300MG</u>	<u>A077686</u>	<u>003</u>	Mar 15, 2010
<u>AB</u>			<u>360MG</u>	<u>A077686</u>	<u>002</u>	Mar 15, 2010
<u>AB</u>			<u>420MG</u>	<u>A077686</u>	<u>001</u>	Mar 15, 2010
<u>AB</u>		AMTA	<u>120MG</u>	<u>A216439</u>	<u>001</u>	Mar 07, 2023
<u>AB</u>			<u>180MG</u>	<u>A216439</u>	<u>002</u>	Mar 07, 2023
<u>AB</u>			<u>240MG</u>	<u>A216439</u>	<u>003</u>	Mar 07, 2023
<u>AB</u>			<u>300MG</u>	<u>A216439</u>	<u>004</u>	Mar 07, 2023
<u>AB</u>			<u>360MG</u>	<u>A216439</u>	<u>005</u>	Mar 07, 2023
<u>AB</u>			<u>420MG</u>	<u>A216439</u>	<u>006</u>	Mar 07, 2023
<u>AB</u>		SCIEGEN PHARMS INC	<u>120MG</u>	<u>A216327</u>	<u>001</u>	Apr 06, 2023
<u>AB</u>			<u>180MG</u>	<u>A216327</u>	<u>002</u>	Apr 06, 2023
<u>AB</u>			<u>240MG</u>	<u>A216327</u>	<u>003</u>	Apr 06, 2023
<u>AB</u>			<u>300MG</u>	<u>A216327</u>	<u>004</u>	Apr 06, 2023
<u>AB</u>			<u>360MG</u>	<u>A216327</u>	<u>005</u>	Apr 06, 2023
<u>AB</u>			<u>420MG</u>	<u>A216327</u>	<u>006</u>	Apr 06, 2023

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

!	FRESENIUS KABI USA	50MG/ML	A040519	001	Jun 23, 2004
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DIMERCAPROL

INJECTABLE; INJECTION

BAL

+	PROVEPHARM SAS	10%	N005939	001	
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PRESCRIPTION DRUG PRODUCT LIST

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

DIMETHYL FUMARATE

<u>AB</u>	ACCORD HLTHCARE	<u>120MG</u>	<u>A210499</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210499</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	ALKEM LABS LTD	<u>120MG</u>	<u>A210440</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210440</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	AMNEAL	<u>120MG</u>	<u>A210402</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210402</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	AUROBINDO PHARMA	<u>120MG</u>	<u>A210385</u>	<u>001</u>	Dec 22, 2022
<u>AB</u>		<u>240MG</u>	<u>A210385</u>	<u>002</u>	Dec 22, 2022
<u>AB</u>	CIPLA	<u>120MG</u>	<u>A210305</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210305</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	GLENMARK PHARMS LTD	<u>120MG</u>	<u>A210309</u>	<u>001</u>	Oct 06, 2020
<u>AB</u>		<u>240MG</u>	<u>A210309</u>	<u>002</u>	Oct 06, 2020
<u>AB</u>	HETERO LABS LTD III	<u>120MG</u>	<u>A210500</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210500</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	LUPIN	<u>120MG</u>	<u>A210226</u>	<u>001</u>	Oct 05, 2020
<u>AB</u>		<u>240MG</u>	<u>A210226</u>	<u>002</u>	Oct 05, 2020
<u>AB</u>	MSN	<u>120MG</u>	<u>A210460</u>	<u>001</u>	Sep 24, 2020
<u>AB</u>		<u>240MG</u>	<u>A210460</u>	<u>002</u>	Sep 24, 2020
<u>AB</u>	MYLAN	<u>120MG</u>	<u>A210531</u>	<u>001</u>	Aug 17, 2020
<u>AB</u>		<u>240MG</u>	<u>A210531</u>	<u>002</u>	Aug 17, 2020
<u>AB</u>	PRINSTON INC	<u>120MG</u>	<u>A210414</u>	<u>001</u>	Oct 18, 2022
<u>AB</u>		<u>240MG</u>	<u>A210414</u>	<u>002</u>	Oct 18, 2022
<u>AB</u>	SOLA PHARMS	<u>120MG</u>	<u>A210436</u>	<u>001</u>	Mar 26, 2021
<u>AB</u>		<u>240MG</u>	<u>A210436</u>	<u>002</u>	Mar 26, 2021
<u>AB</u>	TWI PHARMS	<u>120MG</u>	<u>A210382</u>	<u>001</u>	Oct 14, 2020
<u>AB</u>		<u>240MG</u>	<u>A210382</u>	<u>002</u>	Oct 14, 2020
<u>TECFIDERA</u>					
<u>AB</u>	+ BIOGEN INC	<u>120MG</u>	<u>N204063</u>	<u>001</u>	Mar 27, 2013
<u>AB</u>	+!	<u>240MG</u>	<u>N204063</u>	<u>002</u>	Mar 27, 2013

DIMETHYL SULFOXIDE

SOLUTION;INTRAVESICAL

RIMSO-50

+! MYLAN INSTITUTIONAL 50% N017788 001

DINOPROSTONE

GEL;ENDOCERVICAL

PREPIDIL

+! PFIZER 0.5MG/3GM N019617 001 Dec 09, 1992

INSERT, EXTENDED RELEASE;VAGINAL

CERVIDIL

+! FERRING PHARMS INC 10MG N020411 001 Mar 30, 1995

DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR;ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

+! PHARM ASSOC 12.5MG/5ML A087513 001 Feb 10, 1982

INJECTABLE;INJECTION

DIPHENHYDRAMINE HYDROCHLORIDE

<u>AP</u>	APP PHARMS	<u>50MG/ML</u>	<u>A040466</u>	<u>001</u>	May 28, 2002
<u>AP</u>	MICRO LABS	<u>50MG/ML</u>	<u>A205723</u>	<u>001</u>	Aug 22, 2018
<u>AP</u>	MYLAN INSTITUTIONAL	<u>50MG/ML</u>	<u>A040498</u>	<u>001</u>	Jul 12, 2005
<u>AP</u>	+! WEST-WARD PHARMS	<u>50MG/ML</u>	<u>A080817</u>	<u>002</u>	

INT

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A091526</u>	<u>001</u>	Mar 26, 2013
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DIPYRIDAMOLE

INJECTABLE;INJECTION

DIPYRIDAMOLE

<u>AP</u>	! CHARTWELL	<u>5MG/ML</u>	<u>A074939</u>	<u>001</u>	Apr 13, 1998
	INJECTABLE				
<u>AP</u>	HIKMA	<u>5MG/ML</u>	<u>A074521</u>	<u>001</u>	Oct 18, 1996

TABLET;ORAL

DIPYRIDAMOLE

<u>AB</u>	BARR	<u>25MG</u>	<u>A087184</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>		<u>50MG</u>	<u>A087716</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>		<u>75MG</u>	<u>A087717</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>	IMPAX LABS	<u>25MG</u>	<u>A040782</u>	<u>001</u>	Jul 18, 2007
<u>AB</u>		<u>50MG</u>	<u>A040782</u>	<u>002</u>	Jul 18, 2007
<u>AB</u>		<u>75MG</u>	<u>A040782</u>	<u>003</u>	Jul 18, 2007
<u>AB</u>	OXFORD PHARMS	<u>25MG</u>	<u>A040542</u>	<u>001</u>	Apr 21, 2006

PRESCRIPTION DRUG PRODUCT LIST

DIPYRIDAMOLE

TABLET;ORAL

DIPYRIDAMOLE

<u>AB</u>		<u>50MG</u>	<u>A040542 002</u>	Apr 21, 2006
<u>AB</u>		<u>75MG</u>	<u>A040542 003</u>	Apr 21, 2006
<u>AB</u>	RISING	<u>25MG</u>	<u>A040733 001</u>	Feb 13, 2007
<u>AB</u>		<u>50MG</u>	<u>A040733 002</u>	Feb 13, 2007
<u>AB</u>		<u>75MG</u>	<u>A040733 003</u>	Feb 13, 2007
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A040874 001</u>	Jan 28, 2008
<u>AB</u>		<u>50MG</u>	<u>A040874 002</u>	Jan 28, 2008
<u>AB</u>		<u>75MG</u>	<u>A040874 003</u>	Jan 28, 2008
<u>PERSANTINE</u>				
<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>25MG</u>	<u>N012836 003</u>	Dec 22, 1986
<u>AB</u>	+	<u>50MG</u>	<u>N012836 004</u>	Feb 06, 1987
<u>AB</u>	+	<u>75MG</u>	<u>N012836 005</u>	Feb 06, 1987

DIROXIMEL FUMARATE

CAPSULE, DELAYED RELEASE;ORAL

VUMERITY

+! BIOGEN INC 231MG N211855 001 Oct 29, 2019

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATE

<u>AB</u>	DR REDDYS LABS SA	<u>EQ 100MG BASE</u>	<u>A070173 001</u>	May 31, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A070173 002</u>	May 31, 1985
<u>AB</u>	TEVA	<u>EQ 100MG BASE</u>	<u>A070101 001</u>	Feb 22, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A070102 001</u>	Feb 22, 1985
<u>NORPACE</u>				
<u>AB</u>	+ PFIZER	<u>EQ 100MG BASE</u>	<u>N017447 001</u>	
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N017447 002</u>	

CAPSULE, EXTENDED RELEASE;ORAL

NORPACE CR

+ PFIZER EQ 100MG BASE N018655 001 Jul 20, 1982

+! EQ 150MG BASE N018655 002 Jul 20, 1982

DISULFIRAM

TABLET;ORAL

DISULFIRAM

<u>AB</u>	ALVOGEN	<u>250MG</u>	<u>A091681 001</u>	Aug 08, 2013
<u>AB</u>	CHARTWELL MOLECULES	<u>250MG</u>	<u>A091563 001</u>	Dec 31, 2012
<u>AB</u>	!	<u>500MG</u>	<u>A091563 002</u>	Dec 31, 2012
<u>AB</u>	SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A091619 001</u>	Mar 28, 2011
<u>AB</u>		<u>500MG</u>	<u>A091619 002</u>	Mar 28, 2011

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL

DEPAKOTEAB +! ABBVIE EQ 125MG VALPROIC ACID N019680 001 Sep 12, 1989DIVALPROEX SODIUM

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A213181 001</u>	Mar 02, 2020
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A078979 001</u>	Jan 23, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A078919 001</u>	Jan 27, 2009

TABLET, DELAYED RELEASE;ORAL

DEPAKOTE

<u>AB</u>	+ ABBVIE	<u>EQ 125MG VALPROIC ACID</u>	<u>N018723 003</u>	Oct 26, 1984
<u>AB</u>	+	<u>EQ 250MG VALPROIC ACID</u>	<u>N018723 001</u>	Mar 10, 1983
<u>AB</u>	+	<u>EQ 500MG VALPROIC ACID</u>	<u>N018723 002</u>	Mar 10, 1983

DIVALPROEX SODIUM

<u>AB</u>	APOTEX	<u>EQ 125MG VALPROIC ACID</u>	<u>A077615 003</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077615 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077615 001</u>	Jul 29, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A090554 001</u>	Apr 21, 2011
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A090554 002</u>	Apr 21, 2011
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A090554 003</u>	Apr 21, 2011
<u>AB</u>	CHARTWELL RX	<u>EQ 125MG VALPROIC ACID</u>	<u>A077296 001</u>	Jul 31, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077296 002</u>	Jul 31, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077296 003</u>	Jul 31, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A078755 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078755 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078755 003</u>	Jul 29, 2008

PRESCRIPTION DRUG PRODUCT LIST

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE;ORAL

DIVALPROEX SODIUM

<u>AB</u>	INVATECH	<u>EQ 125MG VALPROIC ACID</u>	<u>A078290 003</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078290 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078290 001</u>	Jul 29, 2008
<u>AB</u>	LUPIN	<u>EQ 125MG VALPROIC ACID</u>	<u>A078790 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078790 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078790 003</u>	Jul 29, 2008
<u>AB</u>	ORBION PHARMS	<u>EQ 125MG VALPROIC ACID</u>	<u>A078853 001</u>	Nov 25, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078853 002</u>	Nov 25, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078853 003</u>	Nov 25, 2008
<u>AB</u>	PRINSTON INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A090210 001</u>	Nov 30, 2009
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A090210 002</u>	Nov 30, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A090210 003</u>	Nov 30, 2009
<u>AB</u>	SUN PHARM INDS	<u>EQ 125MG VALPROIC ACID</u>	<u>A078597 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078597 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078597 003</u>	Jul 29, 2008
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A079163 001</u>	Apr 05, 2011
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A079163 002</u>	Apr 05, 2011
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A079163 003</u>	Apr 05, 2011
<u>AB</u>	UPSHER SMITH LABS	<u>EQ 125MG VALPROIC ACID</u>	<u>A078182 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078182 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078182 003</u>	Jul 29, 2008
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A077100 001</u>	Mar 05, 2009
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077100 002</u>	Mar 05, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077100 003</u>	Mar 05, 2009

TABLET, EXTENDED RELEASE;ORAL

DEPAKOTE ER

<u>AB</u>	+	ABBEVIE	<u>EQ 250MG VALPROIC ACID</u>	<u>N021168 002</u>	May 31, 2002
<u>AB</u>	+		<u>EQ 500MG VALPROIC ACID</u>	<u>N021168 001</u>	Aug 04, 2000

DIVALPROEX SODIUM

<u>AB</u>		AMNEAL PHARMS	<u>EQ 250MG VALPROIC ACID</u>	<u>A203730 001</u>	May 29, 2015
<u>AB</u>	!		<u>EQ 500MG VALPROIC ACID</u>	<u>A203730 002</u>	May 29, 2015
<u>AB</u>		ANNORA PHARMA	<u>EQ 250MG VALPROIC ACID</u>	<u>A215527 001</u>	Sep 26, 2023
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A215527 002</u>	Sep 26, 2023
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A202419 001</u>	Jun 02, 2014
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A202419 002</u>	Jun 02, 2014
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A090161 001</u>	Mar 15, 2012
<u>AB</u>		LUPIN LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A209286 001</u>	Oct 18, 2019
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A209286 002</u>	Oct 18, 2019
<u>AB</u>		MYLAN	<u>EQ 250MG VALPROIC ACID</u>	<u>A077567 001</u>	Jan 29, 2009
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A077567 002</u>	Jan 29, 2009
<u>AB</u>		REDDYS	<u>EQ 500MG VALPROIC ACID</u>	<u>A090070 001</u>	Mar 12, 2012
<u>AB</u>		UNICHEM	<u>EQ 250MG VALPROIC ACID</u>	<u>A214643 001</u>	Feb 25, 2022
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A214643 002</u>	Feb 25, 2022
<u>AB</u>		WOCKHARDT	<u>EQ 250MG VALPROIC ACID</u>	<u>A078705 002</u>	Feb 10, 2009
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A078705 001</u>	Aug 04, 2009
<u>AB</u>		ZYDUS PHARMS USA INC	<u>EQ 250MG VALPROIC ACID</u>	<u>A078239 001</u>	Feb 27, 2009
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A078239 002</u>	Aug 04, 2009

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

<u>AP</u>		HAINAN POLY	<u>EQ 12.5MG BASE/ML</u>	<u>A216131 001</u>	Dec 21, 2022
<u>AP</u>		HIKMA	<u>EQ 12.5MG BASE/ML</u>	<u>A074277 001</u>	Oct 31, 1994
<u>AP</u>	!	HOSPIRA	<u>EQ 12.5MG BASE/ML</u>	<u>A074086 001</u>	Nov 29, 1993

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	+	BAXTER HLTHCARE	<u>EQ 50MG BASE/100ML</u>	<u>N020255 001</u>	Oct 19, 1993
<u>AP</u>	+	+		<u>EQ 100MG BASE/100ML</u>	<u>N020255 003</u>	Oct 19, 1993
<u>AP</u>	+	+		<u>EQ 200MG BASE/100ML</u>	<u>N020255 004</u>	Oct 19, 1993
<u>AP</u>	+	+		<u>EQ 400MG BASE/100ML</u>	<u>N020255 005</u>	Oct 19, 1993
<u>AP</u>	+	+	HOSPIRA	<u>EQ 50MG BASE/100ML</u>	<u>N020201 003</u>	Oct 19, 1993
<u>AP</u>	+	+		<u>EQ 100MG BASE/100ML</u>	<u>N020201 002</u>	Oct 19, 1993
<u>AP</u>	+	+		<u>EQ 200MG BASE/100ML</u>	<u>N020201 001</u>	Oct 19, 1993
<u>AP</u>	+	+		<u>EQ 400MG BASE/100ML</u>	<u>N020201 006</u>	Jul 07, 1994

PRESCRIPTION DRUG PRODUCT LIST

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

<u>AP</u>	+	ACCORD HLTHCARE	<u>20MG/ML (20MG/ML)</u>	<u>N201195 003</u>	Apr 20, 2012
<u>AP</u>	+		<u>80MG/4ML (20MG/ML)</u>	<u>N201195 004</u>	Apr 20, 2012
<u>AP</u>	+		<u>160MG/8ML (20MG/ML)</u>	<u>N201195 005</u>	Apr 20, 2012
<u>AP</u>		ACTAVIS	<u>20MG/ML (20MG/ML)</u>	<u>N203551 001</u>	Apr 12, 2013
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>N203551 002</u>	Apr 12, 2013
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>N203551 004</u>	Sep 21, 2015
<u>AP</u>		ALEMBIC	<u>20MG/2ML (10MG/ML)</u>	<u>A215744 001</u>	Feb 28, 2023
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A215744 002</u>	Feb 28, 2023
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A215744 003</u>	Feb 28, 2023
<u>AP</u>		AMNEAL	<u>20MG/ML (20MG/ML)</u>	<u>A209640 001</u>	Jan 19, 2018
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A209640 002</u>	Jan 19, 2018
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>A209640 003</u>	Jan 19, 2018
<u>AP</u>		DR REDDYS	<u>20MG/ML (20MG/ML)</u>	<u>A204193 001</u>	Nov 05, 2014
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A204193 002</u>	Nov 05, 2014
<u>AP</u>		EUGIA PHARMA	<u>20MG/2ML (10MG/ML)</u>	<u>A214575 001</u>	Jun 25, 2021
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A214575 002</u>	Jun 25, 2021
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A214575 003</u>	Jun 25, 2021
<u>AP</u>		GLAND PHARMA LTD	<u>20MG/2ML (10MG/ML)</u>	<u>A213510 001</u>	Jul 01, 2021
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A213510 002</u>	Jul 01, 2021
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A213510 003</u>	Jul 01, 2021
<u>AP</u>		HENGRUI PHARMA	<u>20MG/ML (20MG/ML)</u>	<u>A207252 001</u>	Aug 09, 2017
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A207252 002</u>	Aug 09, 2017
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>A207252 003</u>	Aug 09, 2017
<u>AP</u>		HIKMA	<u>20MG/ML (20MG/ML)</u>	<u>A204490 001</u>	Jan 14, 2021
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A204490 002</u>	Jan 14, 2021
<u>AP</u>	+	HOSPIRA INC	<u>20MG/2ML (10MG/ML)</u>	<u>N022234 001</u>	Mar 08, 2011
<u>AP</u>	+		<u>80MG/8ML (10MG/ML)</u>	<u>N022234 002</u>	Mar 08, 2011
<u>AP</u>	+		<u>160MG/16ML (10MG/ML)</u>	<u>N022234 003</u>	Mar 08, 2011
<u>AP</u>		MEITHEAL	<u>20MG/2ML (10MG/ML)</u>	<u>A209634 001</u>	Aug 24, 2018
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A209634 002</u>	Aug 24, 2018
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A209634 003</u>	Aug 24, 2018
<u>AP</u>		MYLAN LABS LTD	<u>20MG/2ML (10MG/ML)</u>	<u>A210072 001</u>	Jul 02, 2018
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A210848 001</u>	Jul 06, 2018
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>A208137 001</u>	Apr 01, 2019
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A208859 001</u>	Apr 30, 2018
<u>AP</u>		NOVAST LABS	<u>80MG/8ML (10MG/ML)</u>	<u>A207563 002</u>	Aug 31, 2017
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A207563 003</u>	Aug 31, 2017
<u>AP</u>		SANDOZ	<u>20MG/2ML (10MG/ML)</u>	<u>N201525 001</u>	Jun 29, 2011
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>N201525 002</u>	Jun 29, 2011
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>N201525 003</u>	Jun 29, 2011
<u>AP</u>		SHILPA	<u>20MG/ML (20MG/ML)</u>	<u>A210327 001</u>	May 16, 2019
<u>AP</u>	+		<u>20MG/ML (20MG/ML)</u>	<u>N205934 001</u>	Dec 22, 2015
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A210327 002</u>	May 16, 2019
<u>AP</u>	+		<u>80MG/4ML (20MG/ML)</u>	<u>N205934 002</u>	Dec 22, 2015
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>A210327 003</u>	May 16, 2019
<u>AP</u>	+		<u>160MG/8ML (20MG/ML)</u>	<u>N205934 003</u>	Dec 22, 2015
<u>AP</u>		SUN PHARM	<u>20MG/ML (20MG/ML)</u>	<u>N022534 003</u>	Jan 08, 2019
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>N022534 004</u>	Jan 08, 2019
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>N022534 005</u>	Jan 08, 2019

TAXOTERE

<u>AP</u>	+	SANOFI AVENTIS US	<u>20MG/ML (20MG/ML)</u>	<u>N020449 003</u>	Aug 03, 2010
<u>AP</u>	+		<u>80MG/4ML (20MG/ML)</u>	<u>N020449 004</u>	Aug 02, 2010
<u>AP</u>	+		<u>160MG/8ML (20MG/ML)</u>	<u>N020449 005</u>	Apr 13, 2012

DOCETAXEL

		ACTAVIS	140MG/7ML (20MG/ML)	N203551 003	Apr 12, 2013
	+	HOSPIRA INC	20MG/ML (20MG/ML)	N022234 004	Jun 23, 2016
	+		80MG/4ML (20MG/ML)	N022234 005	Jun 23, 2016
	+		160MG/8ML (20MG/ML)	N022234 007	Jan 24, 2017

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>0.125MG</u>	<u>A210740 001</u>	Jan 22, 2019
<u>AB</u>			<u>0.25MG</u>	<u>A210740 002</u>	Jan 22, 2019
<u>AB</u>			<u>0.5MG</u>	<u>A210740 003</u>	Jan 22, 2019
<u>DOFETILIDE</u>					
<u>AB</u>		ACCORD HLTHCARE	<u>0.125MG</u>	<u>A213338 001</u>	Jun 19, 2020
<u>AB</u>			<u>0.25MG</u>	<u>A213338 002</u>	Jun 19, 2020
<u>AB</u>			<u>0.5MG</u>	<u>A213338 003</u>	Jun 19, 2020

PRESCRIPTION DRUG PRODUCT LIST

DOFETILIDE

CAPSULE;ORAL

DOFETILIDE

<u>AB</u>	BIONPHARMA	<u>0.125MG</u>	<u>A208625</u>	<u>001</u>	Apr 10, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A208625</u>	<u>002</u>	Apr 10, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A208625</u>	<u>003</u>	Apr 10, 2018
<u>AB</u>	DR REDDYS LABS SA	<u>0.125MG</u>	<u>A207058</u>	<u>001</u>	Jun 06, 2016
<u>AB</u>		<u>0.25MG</u>	<u>A207058</u>	<u>002</u>	Jun 06, 2016
<u>AB</u>		<u>0.5MG</u>	<u>A207058</u>	<u>003</u>	Jun 06, 2016
<u>AB</u>	GRANULES	<u>0.125MG</u>	<u>A212750</u>	<u>001</u>	Oct 14, 2021
<u>AB</u>		<u>0.25MG</u>	<u>A212750</u>	<u>002</u>	Oct 14, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A212750</u>	<u>003</u>	Oct 14, 2021
<u>AB</u>	MSN	<u>0.125MG</u>	<u>A213220</u>	<u>001</u>	Jan 29, 2020
<u>AB</u>		<u>0.25MG</u>	<u>A213220</u>	<u>002</u>	Jan 29, 2020
<u>AB</u>		<u>0.5MG</u>	<u>A213220</u>	<u>003</u>	Jan 29, 2020
<u>AB</u>	SIGMAPHARM LABS LLC	<u>0.125MG</u>	<u>A207746</u>	<u>001</u>	Mar 26, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A207746</u>	<u>002</u>	Mar 26, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A207746</u>	<u>003</u>	Mar 26, 2018
<u>AB</u>	STRIDES PHARMA	<u>0.125MG</u>	<u>A208519</u>	<u>001</u>	Oct 09, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A208519</u>	<u>002</u>	Oct 09, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A208519</u>	<u>003</u>	Oct 09, 2018
<u>AB</u>	SUN PHARM	<u>0.125MG</u>	<u>A210466</u>	<u>001</u>	Oct 09, 2018
<u>AB</u>		<u>0.25MG</u>	<u>A210466</u>	<u>002</u>	Oct 09, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A210466</u>	<u>003</u>	Oct 09, 2018
<u>TIKOSYN</u>					
<u>AB</u>	+ PFIZER	<u>0.125MG</u>	<u>N020931</u>	<u>001</u>	Oct 01, 1999
<u>AB</u>	+	<u>0.25MG</u>	<u>N020931</u>	<u>002</u>	Oct 01, 1999
<u>AB</u>	+!	<u>0.5MG</u>	<u>N020931</u>	<u>003</u>	Oct 01, 1999

DOLASETRON MESYLATE

TABLET;ORAL

ANZEMET

+! VALIDUS PHARMS 50MG N020623 001 Sep 11, 1997

DOLUTEGRAVIR SODIUM

TABLET;ORAL

TIVICAY

+ VIIV HLTHCARE EQ 10MG BASE N204790 002 Jun 09, 2016

+ EQ 25MG BASE N204790 003 Jun 09, 2016

+! EQ 50MG BASE N204790 001 Aug 12, 2013

TABLET, FOR SUSPENSION;ORAL

TIVICAY PD

+! VIIV HLTHCARE EQ 5MG BASE N213983 001 Jun 12, 2020

DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

DOVATO

+! VIIV HLTHCARE EQ 50MG BASE;300MG N211994 001 Apr 08, 2019

DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

JULUCA

+! VIIV HLTHCARE EQ 50MG BASE;EQ 25MG BASE N210192 001 Nov 21, 2017

DONEPEZIL HYDROCHLORIDE

SYSTEM;TRANSDERMAL

ADLARITY

+ CORIUM 5MG/DAY N212304 001 Mar 11, 2022

+! 10MG/DAY N212304 002 Mar 11, 2022

TABLET;ORAL

ARICEPT

<u>AB</u>	+ EISAI INC	<u>5MG</u>	<u>N020690</u>	<u>002</u>	Nov 25, 1996
<u>AB</u>	+!	<u>10MG</u>	<u>N020690</u>	<u>001</u>	Nov 25, 1996
<u>AB</u>	+!	<u>23MG</u>	<u>N022568</u>	<u>001</u>	Jul 23, 2010

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	ACI	<u>5MG</u>	<u>A078662</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A078662</u>	<u>002</u>	May 31, 2011
<u>AB</u>	ALEMBIC PHARMS LTD	<u>5MG</u>	<u>A201724</u>	<u>001</u>	Feb 25, 2013
<u>AB</u>		<u>10MG</u>	<u>A201724</u>	<u>002</u>	Feb 25, 2013
<u>AB</u>	AUROBINDO	<u>5MG</u>	<u>A090056</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090056</u>	<u>002</u>	May 31, 2011
<u>AB</u>	CADILA	<u>5MG</u>	<u>A090100</u>	<u>001</u>	Oct 24, 2012
<u>AB</u>		<u>10MG</u>	<u>A090100</u>	<u>002</u>	Oct 24, 2012
<u>AB</u>	CADILA PHARMS LTD	<u>5MG</u>	<u>A204609</u>	<u>001</u>	Sep 19, 2017
<u>AB</u>		<u>10MG</u>	<u>A204609</u>	<u>002</u>	Sep 19, 2017

PRESCRIPTION DRUG PRODUCT LIST

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	CHARTWELL RX	<u>5MG</u>	<u>A090425</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090425</u>	<u>002</u>	May 31, 2011
<u>AB</u>	CIPLA LTD	<u>5MG</u>	<u>A077518</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A077518</u>	<u>002</u>	May 31, 2011
<u>AB</u>	DEXCEL	<u>23MG</u>	<u>A203713</u>	<u>001</u>	Feb 19, 2016
<u>AB</u>	DR REDDYS	<u>23MG</u>	<u>A202723</u>	<u>001</u>	Jul 24, 2013
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A201001</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A201001</u>	<u>002</u>	May 31, 2011
<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A203034</u>	<u>001</u>	Jan 30, 2015
<u>AB</u>		<u>10MG</u>	<u>A203034</u>	<u>002</u>	Jan 30, 2015
<u>AB</u>	INDICUS PHARMA	<u>5MG</u>	<u>A201634</u>	<u>001</u>	Jun 13, 2012
<u>AB</u>		<u>10MG</u>	<u>A201634</u>	<u>002</u>	Jun 13, 2012
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A090768</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090768</u>	<u>002</u>	May 31, 2011
<u>AB</u>	LUPIN LTD	<u>23MG</u>	<u>A202782</u>	<u>001</u>	Oct 30, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201146</u>	<u>001</u>	Aug 17, 2012
<u>AB</u>		<u>10MG</u>	<u>A201146</u>	<u>002</u>	Aug 17, 2012
<u>AB</u>		<u>23MG</u>	<u>A202631</u>	<u>001</u>	Jan 22, 2014
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A200292</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A200292</u>	<u>002</u>	May 31, 2011
<u>AB</u>	RISING	<u>5MG</u>	<u>A202114</u>	<u>001</u>	Jul 05, 2013
<u>AB</u>		<u>10MG</u>	<u>A202114</u>	<u>002</u>	Jul 05, 2013
<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A203907</u>	<u>001</u>	Oct 29, 2014
<u>AB</u>		<u>10MG</u>	<u>A203907</u>	<u>002</u>	Oct 29, 2014
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A090551</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090551</u>	<u>002</u>	May 31, 2011
<u>AB</u>	SUN PHARM INDS	<u>5MG</u>	<u>A090493</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090493</u>	<u>002</u>	May 31, 2011
<u>AB</u>	TORRENT PHARMS	<u>5MG</u>	<u>A090686</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090686</u>	<u>002</u>	May 31, 2011
<u>AB</u>	TWI PHARMS	<u>23MG</u>	<u>A203104</u>	<u>001</u>	Oct 29, 2014
<u>AB</u>	ZYDUS PHARMS	<u>23MG</u>	<u>A203162</u>	<u>001</u>	Aug 31, 2017

TABLET, ORALLY DISINTEGRATING; ORAL

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	HISUN PHARM HANGZHOU	<u>5MG</u>	<u>A205269</u>	<u>001</u>	Jul 27, 2018
<u>AB</u>		<u>10MG</u>	<u>A205269</u>	<u>002</u>	Jul 27, 2018
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201787</u>	<u>001</u>	Dec 14, 2012
<u>AB</u>		<u>10MG</u>	<u>A201787</u>	<u>002</u>	Dec 14, 2012
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A091198</u>	<u>001</u>	May 10, 2011
<u>AB</u>	!	<u>10MG</u>	<u>A091198</u>	<u>002</u>	May 10, 2011

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>10MG; 21MG</u>	<u>A208237</u>	<u>001</u>	Dec 15, 2023
<u>AB</u>	<u>NAMZARIC</u>				
<u>AB</u>	+ ABBVIE	<u>10MG; 21MG</u>	<u>N206439</u>	<u>004</u>	Jul 18, 2016
	+	10MG; 7MG	N206439	003	Jul 18, 2016
	+	10MG; 14MG	N206439	001	Dec 23, 2014
	+!	10MG; 28MG	N206439	002	Dec 23, 2014

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

<u>AP</u>	HIKMA INTL PHARMS	<u>40MG/ML</u>	<u>A207707</u>	<u>001</u>	Apr 11, 2018
<u>AP</u>		<u>80MG/ML</u>	<u>A207707</u>	<u>002</u>	Apr 11, 2018
<u>AP</u>	+! HOSPIRA	<u>40MG/ML</u>	<u>N018132</u>	<u>001</u>	
<u>AP</u>	+!	<u>80MG/100ML</u>	<u>N018132</u>	<u>002</u>	Feb 04, 1982
<u>AP</u>	+!	<u>80MG/ML</u>	<u>N018132</u>	<u>004</u>	Jul 09, 1982
<u>AP</u>	+!	<u>160MG/100ML</u>	<u>N018132</u>	<u>003</u>	Feb 04, 1982

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+! BAXTER HLTHCARE	<u>80MG/100ML</u>	<u>N019615</u>	<u>001</u>	Mar 27, 1987
<u>AP</u>	+!	<u>160MG/100ML</u>	<u>N019615</u>	<u>002</u>	Mar 27, 1987
<u>AP</u>	+!	<u>320MG/100ML</u>	<u>N019615</u>	<u>003</u>	Mar 27, 1987
<u>AP</u>	+! HOSPIRA	<u>80MG/100ML</u>	<u>N018826</u>	<u>001</u>	Sep 30, 1983
<u>AP</u>	+!	<u>160MG/100ML</u>	<u>N018826</u>	<u>002</u>	Sep 30, 1983
<u>AP</u>	+!	<u>320MG/100ML</u>	<u>N018826</u>	<u>003</u>	Sep 30, 1983
	+! BAXTER HLTHCARE	640MG/100ML	N019615	004	Mar 27, 1987

PRESCRIPTION DRUG PRODUCT LIST

DORAVIRINE

TABLET; ORAL

PIFELTRO

+! MSD MERCK CO 100MG

N210806 001 Aug 30, 2018

DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

DELSTRIGO

+! MSD MERCK CO 100MG; 300MG; 300MG

N210807 001 Aug 30, 2018

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

<u>AT</u>	ALEMBIC	<u>EQ 2% BASE</u>	<u>A212639 001</u>	Aug 09, 2019
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 2% BASE</u>	<u>A090143 001</u>	Jun 25, 2009
<u>AT</u>	FDC LTD	<u>EQ 2% BASE</u>	<u>A205294 001</u>	Jan 24, 2019
<u>AT</u>	GLAND PHARMA LTD	<u>EQ 2% BASE</u>	<u>A215660 001</u>	Jan 27, 2022
<u>AT</u>	HIKMA	<u>EQ 2% BASE</u>	<u>A077846 001</u>	Oct 28, 2008
<u>AT</u>	INDOCO	<u>EQ 2% BASE</u>	<u>A202053 001</u>	Sep 11, 2014
<u>AT</u>	! MICRO LABS	<u>EQ 2% BASE</u>	<u>A204778 001</u>	Nov 08, 2019
<u>AT</u>	RUBICON	<u>EQ 2% BASE</u>	<u>A078395 001</u>	Oct 28, 2008
<u>AT</u>	SANDOZ	<u>EQ 2% BASE</u>	<u>A078748 001</u>	Nov 06, 2008
<u>AT</u>		<u>EQ 2% BASE</u>	<u>A078981 001</u>	Apr 13, 2009

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COSOPT

<u>AT1</u>	+! THEA PHARMA	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>N020869 001</u>	Apr 07, 1998
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DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

<u>AT1</u>	ALEMBIC	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A213099 001</u>	May 04, 2021
<u>AT1</u>	BAUSCH AND LOMB	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A090037 001</u>	Jul 14, 2009
<u>AT1</u>	EPIC PHARMA LLC	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A203058 001</u>	Sep 22, 2014
<u>AT1</u>	EUGIA PHARMA	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A207629 001</u>	May 14, 2021
<u>AT1</u>	FDC LTD	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A205295 001</u>	Jun 13, 2019
<u>AT1</u>	GLAND PHARMA LTD	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A215520 001</u>	Sep 19, 2022
<u>AT1</u>	HIKMA	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A077847 001</u>	Oct 28, 2008
<u>AT1</u>	INDOCO	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A202054 001</u>	Sep 03, 2014
<u>AT1</u>	MICRO LABS	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A204777 001</u>	May 28, 2020
<u>AT1</u>	SANDOZ	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A078749 001</u>	Nov 06, 2008
<u>AT1</u>		<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A090604 001</u>	Nov 18, 2009
<u>AT1</u>	SOMERSET	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A207523 001</u>	Jun 25, 2019

COSOPT PF

<u>AT2</u>	+! THEA PHARMA	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>N202667 001</u>	Feb 01, 2012
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DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

<u>AT2</u>	EUGIA PHARMA	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A207630 001</u>	Jul 24, 2018
<u>AT2</u>	INGENUS PHARMS LLC	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A217260 001</u>	May 01, 2023
<u>AT2</u>	MICRO LABS	<u>EQ 2% BASE; EQ 0.5% BASE</u>	<u>A215936 001</u>	Jan 25, 2022

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOPRAM

<u>AP</u>	+! HIKMA	<u>20MG/ML</u>	<u>N014879 001</u>	
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DOXAPRAM HYDROCHLORIDE

<u>AP</u>	CHARTWELL	<u>20MG/ML</u>	<u>A076266 001</u>	Jan 10, 2003
	INJECTABLE			

DOXAZOSIN MESYLATE

TABLET; ORAL

CARDURA

<u>AB</u>	+! VIATRIS	<u>EQ 1MG BASE</u>	<u>N019668 001</u>	Nov 02, 1990
<u>AB</u>	+	<u>EQ 2MG BASE</u>	<u>N019668 002</u>	Nov 02, 1990
<u>AB</u>	+	<u>EQ 4MG BASE</u>	<u>N019668 003</u>	Nov 02, 1990
<u>AB</u>	+	<u>EQ 8MG BASE</u>	<u>N019668 004</u>	Nov 02, 1990

DOXAZOSIN MESYLATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 1MG BASE</u>	<u>A202824 001</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A202824 002</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A202824 003</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A202824 004</u>	Jun 11, 2014
<u>AB</u>	APOTEX	<u>EQ 1MG BASE</u>	<u>A075580 001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075580 002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A075580 003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A075580 004</u>	Oct 18, 2000
<u>AB</u>	HERITAGE PHARMA	<u>EQ 1MG BASE</u>	<u>A205210 001</u>	Feb 13, 2018
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205210 002</u>	Feb 13, 2018
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205210 003</u>	Feb 13, 2018

PRESCRIPTION DRUG PRODUCT LIST

DOXAZOSIN MESYLATE

TABLET;ORAL

DOXAZOSIN MESYLATE

<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A205210 004</u>	Feb 13, 2018
<u>AB</u>	RISING	<u>EQ 1MG BASE</u>	<u>A212727 001</u>	Mar 15, 2022
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A212727 002</u>	Mar 15, 2022
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A212727 003</u>	Mar 15, 2022
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A212727 004</u>	Mar 15, 2022
<u>AB</u>	TEVA	<u>EQ 1MG BASE</u>	<u>A075536 001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075536 002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A075536 003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A075536 004</u>	Oct 18, 2000
<u>AB</u>	UNICHEM	<u>EQ 1MG BASE</u>	<u>A212329 001</u>	Jan 10, 2024
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A212329 002</u>	Jan 10, 2024
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A212329 003</u>	Jan 10, 2024
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A212329 004</u>	Jan 10, 2024
<u>AB</u>	UPSHER SMITH LABS	<u>EQ 1MG BASE</u>	<u>A209013 001</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A209013 002</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A209013 003</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A209013 004</u>	Apr 17, 2018
<u>AB</u>	ZYDUS PHARMS	<u>EQ 1MG BASE</u>	<u>A208719 001</u>	Jul 07, 2017
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A208719 002</u>	Jul 07, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A208719 003</u>	Jul 07, 2017
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A208719 004</u>	Jul 07, 2017

TABLET, EXTENDED RELEASE;ORAL

CARDURA XL

+ VIATRIS

+!

EQ 4MG BASE

EQ 8MG BASE

N021269 001 Feb 22, 2005

N021269 002 Feb 22, 2005

DOXEPIN HYDROCHLORIDE

CAPSULE;ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A212624 001</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A212624 002</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A212624 003</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A212624 004</u>	Sep 13, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A212624 005</u>	Sep 13, 2019
<u>AB</u>	ALEMBIC	<u>EQ 10MG BASE</u>	<u>A215076 001</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215076 002</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215076 003</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215076 004</u>	Apr 21, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215076 005</u>	Apr 21, 2021
<u>AB</u>	AMNEAL PHARMS CO	<u>EQ 10MG BASE</u>	<u>A207482 001</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A207482 002</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A207482 003</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A207482 004</u>	Jun 28, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A207482 005</u>	Jun 28, 2017
<u>AB</u>	APPCO	<u>EQ 10MG BASE</u>	<u>A214908 001</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A214908 002</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A214908 003</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214908 004</u>	Apr 01, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A214908 005</u>	Apr 01, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A211603 001</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211603 002</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A211603 003</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211603 004</u>	Mar 27, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211603 005</u>	Mar 27, 2019
<u>AB</u>	CHARTWELL RX	<u>EQ 10MG BASE</u>	<u>A210268 001</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A210268 002</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210268 003</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A210268 004</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210268 005</u>	Sep 04, 2020
<u>AB</u>	CHEMISTRY HLTH	<u>EQ 10MG BASE</u>	<u>A211619 001</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211619 002</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A211619 003</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211619 004</u>	Mar 09, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211619 005</u>	Mar 09, 2021
<u>AB</u>	CONTRACT PHARMACAL	<u>EQ 10MG BASE</u>	<u>A213474 001</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213474 002</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213474 003</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A213474 004</u>	Jul 28, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213474 005</u>	Jul 28, 2020

PRESCRIPTION DRUG PRODUCT LIST

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	EDENBRIDGE PHARMS	<u>EQ 150MG BASE</u>	<u>A213796 001</u>	Apr 19, 2022
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 10MG BASE</u>	<u>A210675 001</u>	Oct 16, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A210675 002</u>	Oct 16, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210675 004</u>	Mar 03, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A210675 005</u>	Mar 03, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210675 003</u>	Oct 16, 2020
<u>AB</u>	JUBILANT CADISTA	<u>EQ 10MG BASE</u>	<u>A215483 001</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215483 002</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215483 003</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215483 004</u>	Mar 14, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215483 005</u>	Mar 14, 2022
<u>AB</u>	LEADING	<u>EQ 150MG BASE</u>	<u>A211618 001</u>	Mar 01, 2021
<u>AB</u>	MANKIND PHARMA	<u>EQ 10MG BASE</u>	<u>A215710 001</u>	Feb 09, 2022
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215710 002</u>	Feb 09, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215710 003</u>	Feb 09, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215710 004</u>	Feb 09, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215710 005</u>	Feb 09, 2022
<u>AB</u>	MICRO LABS	<u>EQ 10MG BASE</u>	<u>A217688 001</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A217688 002</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A217688 003</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A217688 004</u>	Jun 27, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A217688 005</u>	Jun 27, 2023
<u>AB</u>	MSN	<u>EQ 10MG BASE</u>	<u>A215113 001</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A215113 002</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A215113 003</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215113 004</u>	Jun 24, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A215113 005</u>	Jun 24, 2022
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A070791 002</u>	May 13, 1986
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A070791 003</u>	May 13, 1986
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A070791 001</u>	May 13, 1986
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A070791 004</u>	May 13, 1986
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A070791 005</u>	May 13, 1986
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A071422 006</u>	Nov 09, 1987
<u>AB</u>	PAR PHARM	<u>EQ 10MG BASE</u>	<u>A213063 001</u>	Jul 01, 2020
<u>AB</u>	TARO	<u>EQ 25MG BASE</u>	<u>A213063 002</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213063 003</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A213063 004</u>	Jul 01, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213063 005</u>	Jul 01, 2020
<u>AB</u>	UNIQUE PHARM	<u>EQ 10MG BASE</u>	<u>A217975 001</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A217975 002</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A217975 003</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A217975 004</u>	Aug 21, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A217975 005</u>	Aug 21, 2023
<u>AB</u>	ZYDUS LIFESCIENCES	<u>EQ 10MG BASE</u>	<u>A210700 001</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A210700 002</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210700 003</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A210700 004</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210700 005</u>	Mar 23, 2023
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A210140 001</u>	Mar 21, 2023

CONCENTRATE; ORAL

DOXEPIN HYDROCHLORIDE

! LANNETT CO INC

EQ 10MG BASE/ML

A074721 001 Dec 29, 1998

CREAM; TOPICAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	AMNEAL	<u>5%</u>	<u>A212357 001</u>	Aug 16, 2023
<u>AB</u>	TEVA PHARMS	<u>5%</u>	<u>A215408 001</u>	Feb 17, 2023
<u>ZONALON</u>				
<u>AB</u>	+! MYLAN	<u>5%</u>	<u>N020126 001</u>	Apr 01, 1994

TABLET; ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 3MG BASE</u>	<u>A201951 001</u>	Jul 26, 2013
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201951 002</u>	Jul 26, 2013
<u>AB</u>	MSN	<u>EQ 3MG BASE</u>	<u>A214823 001</u>	Apr 03, 2023
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A214823 002</u>	Apr 03, 2023
<u>AB</u>	RK PHARMA	<u>EQ 3MG BASE</u>	<u>A202337 001</u>	Jan 20, 2016
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202337 002</u>	Jan 20, 2016
<u>AB</u>	STRIDES PHARMA	<u>EQ 3MG BASE</u>	<u>A202510 001</u>	Jul 24, 2020
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202510 002</u>	Jul 24, 2020

PRESCRIPTION DRUG PRODUCT LIST

DOXEPIN HYDROCHLORIDE

TABLET; ORAL

SILENOR

<u>AB</u>	+	CURRAX	<u>EQ 3MG BASE</u>	<u>N022036 001</u>	Mar 17, 2010
<u>AB</u>	+	!	<u>EQ 6MG BASE</u>	<u>N022036 002</u>	Mar 17, 2010

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

<u>AB</u>		AVET	<u>0.5MCG</u>	<u>A205360 001</u>	Sep 15, 2020
<u>AB</u>			<u>1MCG</u>	<u>A205360 002</u>	Sep 15, 2020
<u>AB</u>			<u>2.5MCG</u>	<u>A205360 003</u>	Sep 15, 2020
<u>AB</u>		RISING	<u>0.5MCG</u>	<u>A201518 001</u>	Sep 09, 2016
<u>AB</u>			<u>1MCG</u>	<u>A201518 002</u>	Sep 09, 2016
<u>AB</u>			<u>2.5MCG</u>	<u>A201518 003</u>	Sep 09, 2016

HECTOROL

<u>AB</u>	+	SANOFI	<u>0.5MCG</u>	<u>N020862 002</u>	Apr 23, 2004
<u>AB</u>	+		<u>1MCG</u>	<u>N020862 003</u>	Jul 13, 2009
<u>AB</u>	+	!	<u>2.5MCG</u>	<u>N020862 001</u>	Jun 09, 1999

INJECTABLE; INJECTION

DOXERCALCIFEROL

<u>AP</u>		ALEMBIC	<u>4MCG/2ML (2MCG/ML)</u>	<u>A215810 001</u>	Jun 15, 2023
<u>AP</u>		AMNEAL	<u>2MCG/ML (2MCG/ML)</u>	<u>A208974 001</u>	May 24, 2017
<u>AP</u>			<u>4MCG/2ML (2MCG/ML)</u>	<u>A208974 002</u>	May 24, 2017
<u>AP</u>		EUGIA PHARMA	<u>2MCG/ML (2MCG/ML)</u>	<u>A213717 001</u>	Jan 24, 2022
<u>AP</u>			<u>4MCG/2ML (2MCG/ML)</u>	<u>A213717 002</u>	Jan 24, 2022
<u>AP</u>		GLAND PHARMA LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210452 001</u>	Sep 26, 2019
<u>AP</u>		HIKMA	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091101 001</u>	Aug 30, 2013
<u>AP</u>	+	HOSPIRA	<u>4MCG/2ML (2MCG/ML)</u>	<u>N208614 001</u>	Jul 24, 2018
<u>AP</u>		LUPIN LTD	<u>4MCG/2ML (2MCG/ML)</u>	<u>A210801 001</u>	Nov 01, 2018
<u>AP</u>		MEITHEAL	<u>4MCG/2ML (2MCG/ML)</u>	<u>A211670 001</u>	Feb 07, 2020
<u>AP</u>		SANDOZ	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091333 001</u>	May 05, 2014
<u>AP</u>			<u>4MCG/2ML (2MCG/ML)</u>	<u>A200926 001</u>	Feb 04, 2014

HECTOROL

<u>AP</u>	+	SANOFI	<u>2MCG/ML (2MCG/ML)</u>	<u>N021027 002</u>	Apr 06, 2000
<u>AP</u>	+	!	<u>4MCG/2ML (2MCG/ML)</u>	<u>N021027 001</u>	Apr 06, 2000

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>		ACTAVIS INC	<u>2MG/ML</u>	<u>A203622 001</u>	Jun 27, 2014
<u>AP</u>			<u>200MG/100ML</u>	<u>A203622 002</u>	Jun 27, 2014
<u>AP</u>		AMNEAL	<u>20MG/VIAL</u>	<u>A208888 001</u>	Feb 17, 2017
<u>AP</u>			<u>50MG/VIAL</u>	<u>A208888 002</u>	Feb 17, 2017
<u>AP</u>		FRESENIUS KABI USA	<u>2MG/ML</u>	<u>A063277 001</u>	Oct 26, 1995
<u>AP</u>		GLAND PHARMA LTD	<u>2MG/ML</u>	<u>A209825 001</u>	Aug 11, 2017
<u>AP</u>		HIKMA	<u>2MG/ML</u>	<u>A062975 001</u>	Mar 17, 1989
<u>AP</u>	!		<u>20MG/VIAL</u>	<u>A062921 002</u>	Mar 17, 1989
<u>AP</u>	!		<u>50MG/VIAL</u>	<u>A062921 003</u>	Mar 17, 1989
<u>AP</u>			<u>200MG/100ML</u>	<u>A064097 001</u>	Sep 13, 1994
<u>AP</u>		MYLAN LABS LTD	<u>50MG/VIAL</u>	<u>A200170 002</u>	Oct 28, 2011
<u>AP</u>	+	PFIZER	<u>2MG/ML</u>	<u>N050629 001</u>	Dec 23, 1987
<u>AP</u>	+		<u>200MG/100ML</u>	<u>N050629 002</u>	May 03, 1988
<u>AP</u>		SAGENT PHARMS	<u>2MG/ML</u>	<u>A091495 001</u>	Mar 18, 2013
<u>AP</u>		SUN PHARM INDS	<u>2MG/ML</u>	<u>A091418 001</u>	Feb 15, 2012
<u>AP</u>	!	HIKMA	<u>10MG/VIAL</u>	<u>A062921 001</u>	Mar 17, 1989
<u>AP</u>	+	PFIZER	<u>150MG/75ML</u>	<u>N050629 003</u>	Mar 28, 2011

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL (LIPOSOMAL)

<u>AB</u>	+	BAXTER HLTHCARE CORP	<u>20MG/10ML (2MG/ML)</u>	<u>N050718 001</u>	Nov 17, 1995
<u>AB</u>	+		<u>50MG/25ML (2MG/ML)</u>	<u>N050718 002</u>	Jun 13, 2000

DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL)

<u>AB</u>		AYANA PHARMA LTD	<u>20MG/10ML (2MG/ML)</u>	<u>A207228 001</u>	Oct 12, 2021
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A207228 002</u>	Oct 12, 2021
<u>AB</u>		BAXTER HLTHCARE CORP	<u>20MG/10ML (2MG/ML)</u>	<u>A212219 001</u>	Oct 19, 2022
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A212219 002</u>	Oct 19, 2022
<u>AB</u>		DR REDDYS	<u>20MG/10ML (2MG/ML)</u>	<u>A208657 001</u>	May 15, 2017
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A208657 002</u>	May 15, 2017
<u>AB</u>	!	SUN PHARM	<u>20MG/10ML (2MG/ML)</u>	<u>A203263 001</u>	Feb 04, 2013
<u>AB</u>	!		<u>50MG/25ML (2MG/ML)</u>	<u>A203263 002</u>	Feb 04, 2013
<u>AB</u>		ZYDUS	<u>20MG/10ML (2MG/ML)</u>	<u>A212299 001</u>	Sep 10, 2020
<u>AB</u>			<u>50MG/25ML (2MG/ML)</u>	<u>A212299 002</u>	Sep 10, 2020

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE

CAPSULE;ORAL

DOXYCYCLINE

<u>AB</u>	ALEMBIC	<u>EQ 75MG BASE</u>	<u>A209165 001</u>	Jul 28, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209165 002</u>	Jul 28, 2017
<u>AB</u>	COSETTE	<u>EQ 50MG BASE</u>	<u>A204446 001</u>	May 28, 2015
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A204446 002</u>	May 28, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204446 003</u>	May 28, 2015
<u>AB</u>	DR REDDYS LABS SA	<u>EQ 50MG BASE</u>	<u>A209396 001</u>	Sep 29, 2017
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A209396 002</u>	Sep 29, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209396 003</u>	Sep 29, 2017
<u>AB</u>	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204234 001</u>	Mar 05, 2014
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A204234 002</u>	Mar 05, 2014
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204234 003</u>	Mar 05, 2014
<u>AB</u>	STRIDES PHARMA	<u>EQ 50MG BASE</u>	<u>A065055 001</u>	Dec 01, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065055 002</u>	Dec 01, 2000
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A065055 003</u>	Jul 15, 2005
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 50MG BASE</u>	<u>A065053 001</u>	Nov 22, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065053 003</u>	Sep 10, 2003
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A065053 002</u>	Nov 22, 2000
<u>AB</u>	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A205115 001</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A205115 002</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A205115 003</u>	Feb 18, 2016

MONODOX

<u>AB</u>	+	CHARTWELL RX	<u>EQ 50MG BASE</u>	<u>N050641 002</u>	Feb 10, 1992
<u>AB</u>	+		<u>EQ 75MG BASE</u>	<u>N050641 003</u>	Oct 18, 2006
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N050641 001</u>	Dec 29, 1989
<u>AB</u>	+		<u>EQ 150MG BASE</u>	<u>N050641 004</u>	Feb 17, 2022
		ORACEA			
	+	!	GALDERMA LABS LP	40MG	N050805 001 May 26, 2006
					FOR SUSPENSION;ORAL

DOXYCYCLINE

<u>AB</u>	CHARTWELL	<u>EQ 25MG BASE/5ML</u>	<u>A065454 001</u>	Jul 16, 2008
<u>AB</u>	LUPIN LTD	<u>EQ 25MG BASE/5ML</u>	<u>A201678 001</u>	Mar 18, 2013

VIBRAMYCIN

<u>AB</u>	+	!	PFIZER	<u>EQ 25MG BASE/5ML</u>	<u>N050006 001</u>
					TABLET;ORAL

DOXYCYCLINE

<u>AB</u>	HERITAGE PHARMS	<u>EQ 50MG BASE</u>	<u>A091605 001</u>	Dec 20, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A091605 002</u>	Dec 20, 2011
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091605 003</u>	Dec 20, 2011
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A091605 004</u>	Dec 20, 2011
<u>AB</u>	LANNETT CO INC	<u>EQ 50MG BASE</u>	<u>A065285 001</u>	Dec 08, 2005
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065285 003</u>	Jul 30, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065285 002</u>	Dec 08, 2005
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065285 004</u>	Jul 30, 2008
<u>AB</u>	STRIDES PHARMA	<u>EQ 50MG BASE</u>	<u>A065070 001</u>	Dec 15, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065070 003</u>	Dec 30, 2002
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065070 002</u>	Dec 15, 2000
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065070 004</u>	Jul 14, 2005
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 50MG BASE</u>	<u>A065356 001</u>	May 31, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065356 002</u>	May 31, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065356 003</u>	May 31, 2006
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065356 004</u>	Jul 29, 2010
<u>AB</u>	ZYDUS PHARMS	<u>EQ 50MG BASE</u>	<u>A209582 001</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A209582 002</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209582 003</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209582 004</u>	Sep 28, 2017

DOXYCYCLINE CALCIUM

SUSPENSION;ORAL

VIBRAMYCIN

	+	!	PFIZER	EQ 50MG BASE/5ML	N050480 001
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DOXYCYCLINE HYCLATE

CAPSULE;ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 50MG BASE</u>	<u>A062031 002</u>	Oct 13, 1982
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062031 001</u>	
<u>AB</u>	ALEMBIC	<u>EQ 50MG BASE</u>	<u>A210527 001</u>	Jun 13, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210527 002</u>	Jun 13, 2018
<u>AB</u>	AMNEAL PHARMS	<u>EQ 100MG BASE</u>	<u>A207289 001</u>	Jun 27, 2016
<u>AB</u>	CADILA	<u>EQ 50MG BASE</u>	<u>A207774 001</u>	May 31, 2018

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A207774 002</u>	May 31, 2018
<u>AB</u>	CHANGZHOU PHARM	<u>EQ 100MG BASE</u>	<u>A209402 001</u>	Oct 07, 2019
<u>AB</u>	CHARTWELL	<u>EQ 50MG BASE</u>	<u>A062500 001</u>	Sep 11, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062500 002</u>	Sep 11, 1984
<u>AB</u>	HIKMA INTL PHARMS	<u>EQ 50MG BASE</u>	<u>A062396 002</u>	Nov 07, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062396 001</u>	May 07, 1984
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 50MG BASE</u>	<u>A062676 002</u>	Jul 10, 1986
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062676 001</u>	Jul 10, 1986

VIBRAMYCIN

<u>AB</u>	+! PFIZER	<u>EQ 100MG BASE</u>	<u>N050007 002</u>	
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INJECTABLE; INJECTION

DOXY 100

<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 100MG BASE/VIAL</u>	<u>A062475 001</u>	Dec 09, 1983
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DOXY 200

<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A062475 002</u>	Dec 09, 1983
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DOXYCYCLINE

<u>AP</u>	MYLAN LABS LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A091406 001</u>	Aug 21, 2012
<u>AP</u>	! ZYDUS PHARMS	<u>EQ 100MG BASE/VIAL</u>	<u>A207757 001</u>	Sep 28, 2017
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A207757 002</u>	Sep 28, 2017

DOXYCYCLINE HYCLATE

<u>AP</u>	GLAND PHARMA LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A215583 001</u>	Apr 12, 2023
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 100MG BASE/VIAL</u>	<u>A062992 001</u>	Feb 16, 1989
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A062992 002</u>	Feb 16, 1989

TABLET; ORAL

ACTICLATE

<u>AB</u>	+ ALMIRALL	<u>EQ 75MG BASE</u>	<u>N205931 001</u>	Jul 25, 2014
<u>AB</u>	+!	<u>EQ 150MG BASE</u>	<u>N205931 002</u>	Jul 25, 2014

DOXYCYCLINE HYCLATE

<u>AB</u>	ACELLA	<u>EQ 100MG BASE</u>	<u>A210664 001</u>	Mar 16, 2020
<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 100MG BASE</u>	<u>A062421 001</u>	Feb 02, 1983
<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 75MG BASE</u>	<u>A211584 001</u>	Jun 01, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211584 002</u>	Jun 01, 2020
<u>AB</u>	ALEMBIC	<u>EQ 20MG BASE</u>	<u>A210537 001</u>	Mar 03, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A210536 002</u>	Sep 14, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A211744 001</u>	Jun 30, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211744 002</u>	Jun 30, 2020
<u>AB</u>	APOTEX	<u>EQ 75MG BASE</u>	<u>A209243 001</u>	Apr 15, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209243 002</u>	Apr 15, 2019
<u>AB</u>	CADILA	<u>EQ 100MG BASE</u>	<u>A207773 001</u>	Oct 30, 2017
<u>AB</u>	CARIBE HOLDINGS	<u>EQ 50MG BASE</u>	<u>A062269 003</u>	Oct 05, 1983
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A062269 002</u>	Nov 08, 1982
<u>AB</u>	CHANGZHOU PHARM	<u>EQ 100MG BASE</u>	<u>A211343 001</u>	Oct 09, 2019
<u>AB</u>	CHARTWELL	<u>EQ 50MG BASE</u>	<u>A062505 002</u>	Jul 27, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062505 001</u>	Sep 11, 1984
<u>AB</u>	! CHARTWELL MOLECULAR	<u>EQ 20MG BASE</u>	<u>A065277 001</u>	Nov 10, 2005
<u>AB</u>	DR REDDYS LABS SA	<u>EQ 75MG BASE</u>	<u>A208765 001</u>	Jun 14, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208765 002</u>	Jun 14, 2017
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 20MG BASE</u>	<u>A065182 001</u>	May 13, 2005
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214207 001</u>	Dec 16, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214207 002</u>	Dec 16, 2020
<u>AB</u>	HIKMA INTL PHARMS	<u>EQ 100MG BASE</u>	<u>A065095 001</u>	Jul 02, 2003
<u>AB</u>	LARKEN LABS	<u>EQ 20MG BASE</u>	<u>A065287 001</u>	Feb 28, 2006
<u>AB</u>	LUPIN LTD	<u>EQ 75MG BASE</u>	<u>A208818 001</u>	Sep 27, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208818 002</u>	Sep 27, 2017
<u>AB</u>	MYLAN	<u>EQ 100MG BASE</u>	<u>A062432 001</u>	Feb 15, 1983
<u>AB</u>	NOVEL LABS INC	<u>EQ 100MG BASE</u>	<u>A207558 001</u>	Sep 06, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 20MG BASE</u>	<u>A065134 001</u>	May 13, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062677 001</u>	Jul 10, 1986

TABLET, DELAYED RELEASE; ORAL

DORYX

<u>AB</u>	+ MAYNE PHARMA	<u>EQ 75MG BASE</u>	<u>N050795 001</u>	May 06, 2005
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N050795 003</u>	Jun 20, 2008
<u>AB</u>	+	<u>EQ 200MG BASE</u>	<u>N050795 005</u>	Apr 11, 2013

DOXYCYCLINE HYCLATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 50MG BASE</u>	<u>A090134 003</u>	May 22, 2018
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090134 001</u>	Dec 14, 2011
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090134 002</u>	Dec 14, 2011

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

TABLET, DELAYED RELEASE;ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090134 004</u>	May 22, 2018
<u>AB</u>	ALEMBIC	<u>EQ 75MG BASE</u>	<u>A213075 001</u>	Jan 03, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213075 002</u>	Jan 03, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A213075 003</u>	Jan 03, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A213075 004</u>	Jan 03, 2022
<u>AB</u>	HERITAGE PHARMS	<u>EQ 75MG BASE</u>	<u>A200856 001</u>	Apr 30, 2013
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A200856 002</u>	Apr 30, 2013
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A200856 003</u>	Apr 30, 2013
<u>AB</u>	!	<u>EQ 200MG BASE</u>	<u>A200856 004</u>	Nov 13, 2018
<u>AB</u>	PRINSTON INC	<u>EQ 50MG BASE</u>	<u>A207494 003</u>	Feb 19, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A207494 001</u>	Nov 15, 2016
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A207494 002</u>	Nov 15, 2016
	DORYX			
	+ MAYNE PHARMA	EQ 80MG BASE	N050795 004	Apr 11, 2013
	DORYX MPC			
	+ MAYNE PHARMA	EQ 60MG BASE	N050795 007	May 20, 2016

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, DELAYED RELEASE;ORAL

DICLEGIS

<u>AB</u>	+! DUCHESNAY	<u>10MG;10MG</u>	<u>N021876 001</u>	Apr 08, 2013
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DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>10MG;10MG</u>	<u>A205811 001</u>	Aug 19, 2016
<u>AB</u>	BIONPHARMA	<u>10MG;10MG</u>	<u>A217000 001</u>	Aug 04, 2023
<u>AB</u>	MYLAN PHARMS INC	<u>10MG;10MG</u>	<u>A207825 001</u>	Jul 06, 2020
<u>AB</u>	PAR PHARM INC	<u>10MG;10MG</u>	<u>A208518 001</u>	Dec 06, 2017

TABLET, EXTENDED RELEASE;ORAL

BONJESTA

	+! DUCHESNAY	20MG;20MG	N209661 001	Nov 07, 2016
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DRONABINOL

CAPSULE;ORAL

DRONABINOL

<u>AB</u>	ASCENT PHARMS INC	<u>2.5MG</u>	<u>A207421 001</u>	Feb 07, 2020
<u>AB</u>		<u>5MG</u>	<u>A207421 002</u>	Feb 07, 2020
<u>AB</u>		<u>10MG</u>	<u>A207421 003</u>	Feb 07, 2020
<u>AB</u>	HIKMA	<u>2.5MG</u>	<u>A079217 001</u>	Jun 20, 2014
<u>AB</u>		<u>5MG</u>	<u>A079217 002</u>	Jun 20, 2014
<u>AB</u>		<u>10MG</u>	<u>A079217 003</u>	Jun 20, 2014
<u>AB</u>	SVC PHARMA	<u>2.5MG</u>	<u>A078292 001</u>	Jun 27, 2008
<u>AB</u>		<u>5MG</u>	<u>A078292 002</u>	Jun 27, 2008
<u>AB</u>		<u>10MG</u>	<u>A078292 003</u>	Jun 27, 2008

MARINOL

<u>AB</u>	+ ALKEM LABS LTD	<u>2.5MG</u>	<u>N018651 001</u>	May 31, 1985
<u>AB</u>	+!	<u>5MG</u>	<u>N018651 002</u>	May 31, 1985
<u>AB</u>	+	<u>10MG</u>	<u>N018651 003</u>	May 31, 1985

SOLUTION;ORAL

SYNDROS

	+! BENUVIA OPERATIONS	5MG/ML	N205525 001	Mar 23, 2017
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DRONEDARONE HYDROCHLORIDE

TABLET;ORAL

MULTAQ

	+! SANOFI AVENTIS US	EQ 400MG BASE	N022425 001	Jul 01, 2009
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DROPERIDOL

INJECTABLE;INJECTION

DROPERIDOL

	! AM REGENT	2.5MG/ML	A072123 001	Oct 24, 1988
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DROSPIRENONE

TABLET;ORAL

SLYND

	+! EXELTIS USA INC	4MG	N211367 001	May 23, 2019
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DROSPIRENONE; ESTETROL

TABLET;ORAL

NEXTSTELLIS

	+! MAYNE PHARMA	3MG;14.2MG	N214154 001	Apr 15, 2021
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PRESCRIPTION DRUG PRODUCT LIST

DROSPIRENONE; ESTRADIOL

TABLET; ORAL

ANGELIQ

+	BAYER HLTHCARE	0.25MG; 0.5MG	N021355 001	Feb 29, 2012
+	!	0.5MG; 1MG	N021355 002	Sep 28, 2005

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

AB	GLENMARK PHARMS LTD	3MG;0.02MG	A204296 001	Aug 17, 2015
AB	HETERO LABS	3MG;0.02MG	A211944 001	Mar 22, 2019
AB	HLTHCARE	3MG;0.02MG	A203291 001	Jul 18, 2017
AB	MYLAN LABS LTD	3MG;0.02MG	A202594 001	Oct 22, 2015
AB	WATSON LABS	3MG;0.02MG	A078833 001	Nov 28, 2011

LO-ZUMANDIMINE

AB	AUROBINDO PHARMA LTD	3MG;0.02MG	A209632 001	Feb 27, 2018
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LORYNA

AB	XIROMED	3MG;0.02MG	A079221 001	Mar 28, 2011
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MELAMISA

AB	NOVAST LABS	3MG;0.02MG	A202016 001	Jan 26, 2016
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NIKKI

AB	LUPIN LTD	3MG;0.02MG	A201661 001	May 27, 2014
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YAZ

AB	+	BAYER HLTHCARE	3MG;0.02MG	N021676 001	Mar 16, 2006
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TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL

AB	DR REDDYS LABS SA	3MG;0.03MG	A090081 001	Sep 07, 2010
AB	GLENMARK PHARMS LTD	3MG;0.03MG	A204848 001	Mar 25, 2016
AB	HETERO LABS	3MG;0.03MG	A213034 001	Jan 24, 2020
AB	LUPIN LTD	3MG;0.03MG	A201663 001	Dec 18, 2012
AB	MYLAN LABS LTD	3MG;0.03MG	A202131 001	May 04, 2015
AB	NAARI PTE LTD	3MG;0.03MG	A207245 001	Nov 22, 2016

SYEDA

AB	XIROMED	3MG;0.03MG	A090114 001	Mar 28, 2011
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YAELA

AB	NOVAST LABS	3MG;0.03MG	A202015 001	Nov 19, 2014
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YASMIN

AB	+	BAYER HLTHCARE	3MG;0.03MG	N021098 001	May 11, 2001
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ZUMANDIMINE

AB	AUROBINDO PHARMA LTD	3MG;0.03MG	A209407 001	Mar 26, 2018
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DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM

TABLET; ORAL

BEYAZ

AB	+	BAYER HLTHCARE	3MG,N/A:0.02MG,N/A:0.451MG,0.451MG	N022532 001	Sep 24, 2010
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DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM

AB	LUPIN LTD	3MG,N/A:0.02MG,N/A:0.451MG,0.451MG	A205947 001	Jun 13, 2018
AB	WATSON LABS INC	3MG,N/A:0.02MG,N/A:0.451MG,0.451MG	A203593 001	Oct 11, 2016
AB		3MG,N/A:0.03MG,N/A:0.451MG,0.451MG	A203594 001	Oct 11, 2016

SAFYRAL

AB	+	BAYER HLTHCARE	3MG,N/A:0.03MG,N/A:0.451MG,0.451MG	N022574 001	Dec 16, 2010
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TYDEMY

AB	LUPIN LTD	3MG,N/A:0.03MG,N/A:0.451MG,0.451MG	A205948 001	Dec 12, 2017
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DROXIDOPA

CAPSULE; ORAL

DROXIDOPA

AB	AJANTA PHARMA LTD	100MG	A214391 001	Feb 18, 2021
AB		200MG	A214391 002	Feb 18, 2021
AB		300MG	A214391 003	Feb 18, 2021
AB	ALKEM LABS LTD	100MG	A213911 001	Feb 18, 2021
AB		200MG	A213911 002	Feb 18, 2021
AB		300MG	A213911 003	Feb 18, 2021
AB	ANNORA	100MG	A211726 002	Aug 09, 2021
AB		200MG	A211726 003	Aug 09, 2021
AB		300MG	A211726 001	Feb 18, 2021
AB	AUROBINDO PHARMA LTD	100MG	A214387 001	Feb 18, 2021
AB		200MG	A214387 002	Feb 18, 2021
AB		300MG	A214387 003	Feb 18, 2021
AB	BIONPHARMA	100MG	A213033 001	Apr 28, 2021
AB		200MG	A213033 002	Apr 28, 2021

PRESCRIPTION DRUG PRODUCT LIST

DROXIDOPA

CAPSULE; ORAL

DROXIDOPA

<u>AB</u>		<u>300MG</u>	<u>A213033</u>	<u>003</u>	Apr 28, 2021
<u>AB</u>	BLUEPHARMA INDUSTRIA	<u>100MG</u>	<u>A214543</u>	<u>001</u>	May 05, 2021
<u>AB</u>		<u>200MG</u>	<u>A214543</u>	<u>002</u>	May 05, 2021
<u>AB</u>		<u>300MG</u>	<u>A214543</u>	<u>003</u>	May 05, 2021
<u>AB</u>	LUPIN PHARMS	<u>100MG</u>	<u>A211652</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A211652</u>	<u>002</u>	Feb 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A211652</u>	<u>003</u>	Feb 18, 2021
<u>AB</u>	MSN PHARMS INC	<u>100MG</u>	<u>A211741</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A211741</u>	<u>002</u>	Feb 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A211741</u>	<u>003</u>	Feb 18, 2021
<u>AB</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A214017</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A214017</u>	<u>002</u>	Feb 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A214017</u>	<u>003</u>	Feb 18, 2021
<u>AB</u>	SLATE RUN PHARMA	<u>100MG</u>	<u>A215265</u>	<u>001</u>	Nov 01, 2021
<u>AB</u>		<u>200MG</u>	<u>A215265</u>	<u>002</u>	Nov 01, 2021
<u>AB</u>		<u>300MG</u>	<u>A215265</u>	<u>003</u>	Nov 01, 2021
<u>AB</u>	SUN PHARM	<u>100MG</u>	<u>A214384</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A214384</u>	<u>002</u>	Feb 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A214384</u>	<u>003</u>	Feb 18, 2021
<u>AB</u>	UPSHER SMITH LABS	<u>100MG</u>	<u>A213661</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A213661</u>	<u>002</u>	Feb 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A213661</u>	<u>003</u>	Feb 18, 2021
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A211818</u>	<u>001</u>	Feb 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A211818</u>	<u>002</u>	Feb 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A211818</u>	<u>003</u>	Feb 18, 2021
<u>NORTHERA</u>					
<u>AB</u>	+ LUNDBECK NA LTD	<u>100MG</u>	<u>N203202</u>	<u>001</u>	Feb 18, 2014
<u>AB</u>	+	<u>200MG</u>	<u>N203202</u>	<u>002</u>	Feb 18, 2014
<u>AB</u>	+!	<u>300MG</u>	<u>N203202</u>	<u>003</u>	Feb 18, 2014

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

CYMBALTA

<u>AB</u>	+ LILLY	<u>EQ 20MG BASE</u>	<u>N021427</u>	<u>001</u>	Aug 03, 2004
<u>AB</u>	+	<u>EQ 30MG BASE</u>	<u>N021427</u>	<u>002</u>	Aug 03, 2004
<u>AB</u>	+!	<u>EQ 60MG BASE</u>	<u>N021427</u>	<u>004</u>	Aug 03, 2004

DULOXETINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 20MG BASE</u>	<u>A090776</u>	<u>001</u>	Dec 17, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090776</u>	<u>002</u>	Dec 17, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090776</u>	<u>003</u>	Dec 17, 2013
<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A208706</u>	<u>001</u>	Jan 06, 2017
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A208706</u>	<u>002</u>	Jan 06, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208706</u>	<u>004</u>	Mar 11, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A208706</u>	<u>003</u>	Jan 06, 2017
<u>AB</u>	ALEMBIC	<u>EQ 20MG BASE</u>	<u>A202949</u>	<u>001</u>	Jun 09, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202949</u>	<u>002</u>	Jun 09, 2014
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A202949</u>	<u>003</u>	Jun 09, 2014
<u>AB</u>	ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A203197</u>	<u>001</u>	Aug 26, 2015
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A203197</u>	<u>002</u>	Aug 26, 2015
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A203197</u>	<u>003</u>	Aug 26, 2015
<u>AB</u>	ANCHEN PHARMS	<u>EQ 20MG BASE</u>	<u>A090780</u>	<u>001</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090780</u>	<u>002</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090780</u>	<u>003</u>	Oct 28, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A090778</u>	<u>001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090778</u>	<u>002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090778</u>	<u>003</u>	Dec 11, 2013
<u>AB</u>	BRECKENRIDGE	<u>EQ 20MG BASE</u>	<u>A203088</u>	<u>001</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A203088</u>	<u>002</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A203088</u>	<u>004</u>	May 18, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A203088</u>	<u>003</u>	Jun 11, 2014
<u>AB</u>	CSEPC OUYI	<u>EQ 20MG BASE</u>	<u>A211310</u>	<u>001</u>	Oct 16, 2018
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A211310</u>	<u>002</u>	Oct 16, 2018
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A211310</u>	<u>003</u>	Oct 16, 2018
<u>AB</u>	HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A204343</u>	<u>001</u>	Aug 03, 2016
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A204343</u>	<u>002</u>	Aug 03, 2016
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204343</u>	<u>003</u>	Aug 03, 2016
<u>AB</u>	INVENTIA	<u>EQ 20MG BASE</u>	<u>A202336</u>	<u>001</u>	Oct 28, 2015
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202336</u>	<u>002</u>	Oct 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

DULOXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A202336 003</u>	Oct 28, 2015
<u>AB</u>	LUPIN LTD	<u>EQ 20MG BASE</u>	<u>A090694 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090694 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090694 003</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090694 004</u>	Dec 11, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A204815 001</u>	Mar 23, 2017
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A204815 002</u>	Mar 23, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204815 003</u>	Mar 23, 2017
<u>AB</u>	MARKSANS PHARMA	<u>EQ 20MG BASE</u>	<u>A090723 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090723 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090723 003</u>	Dec 11, 2013
<u>AB</u>	PRINSTON INC	<u>EQ 20MG BASE</u>	<u>A206653 001</u>	May 18, 2017
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A206653 002</u>	May 18, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A206653 003</u>	May 18, 2017
<u>AB</u>	QINGDAO BAHEAL PHARM	<u>EQ 20MG BASE</u>	<u>A210599 001</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A210599 002</u>	Apr 17, 2019
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A210599 003</u>	Apr 17, 2019
<u>AB</u>	SUN PHARM	<u>EQ 20MG BASE</u>	<u>A090745 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090745 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090745 003</u>	Dec 11, 2013
<u>AB</u>	SUNSHINE	<u>EQ 20MG BASE</u>	<u>A212328 001</u>	Feb 11, 2021
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A212328 002</u>	Feb 11, 2021
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A212328 003</u>	Feb 11, 2021
<u>AB</u>	TORRENT	<u>EQ 20MG BASE</u>	<u>A090774 001</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090774 002</u>	Dec 11, 2013
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090774 003</u>	Dec 11, 2013
<u>AB</u>	ZYDUS HLTHCARE	<u>EQ 20MG BASE</u>	<u>A090739 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090739 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090739 004</u>	Apr 18, 2023
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090739 003</u>	Jan 08, 2014
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE</u>	<u>A090728 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A090728 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090728 003</u>	Jan 08, 2014

DURLOBACTAM SODIUM; DURLOBACTAM SODIUM; SULBACTAM SODIUM

POWDER; INTRAVENOUS

XACDURO (COPACKAGED)

+	!	ENTASIS THERAP	EQ 500MG BASE/VIAL; EQ 500MG BASE/VIAL;EQ 1GM BASE/VIAL	N216974 001	May 23, 2023
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DUTASTERIDE

CAPSULE;ORAL

AVODART

<u>AB</u>	+	!	WOODWARD	<u>0.5MG</u>	<u>N021319 001</u>	Nov 20, 2001
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DUTASTERIDE

<u>AB</u>			ACELLA	<u>0.5MG</u>	<u>A206373 001</u>	Mar 17, 2016
<u>AB</u>			ADAPTIS	<u>0.5MG</u>	<u>A204376 001</u>	Apr 07, 2017
<u>AB</u>			AMNEAL PHARMS	<u>0.5MG</u>	<u>A203118 001</u>	Nov 20, 2015
<u>AB</u>			ASCENT PHARMS INC	<u>0.5MG</u>	<u>A206574 001</u>	Oct 21, 2016
<u>AB</u>			AUROBINDO PHARMA LTD	<u>0.5MG</u>	<u>A202660 001</u>	Nov 20, 2015
<u>AB</u>			BARR	<u>0.5MG</u>	<u>A090095 001</u>	Dec 21, 2010
<u>AB</u>			BIONPHARMA	<u>0.5MG</u>	<u>A200899 001</u>	Nov 20, 2015
<u>AB</u>			CADILA	<u>0.5MG</u>	<u>A204373 001</u>	Oct 04, 2017
<u>AB</u>			HUMANWELL PURACAP	<u>0.5MG</u>	<u>A209909 001</u>	Nov 21, 2017
<u>AB</u>			STRIDES PHARMA	<u>0.5MG</u>	<u>A204262 001</u>	Nov 20, 2015
<u>AB</u>			VINTAGE	<u>0.5MG</u>	<u>A202421 001</u>	Nov 20, 2015

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE;ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

<u>AB</u>			ANCHEN PHARMS	<u>0.5MG;0.4MG</u>	<u>A202509 001</u>	Feb 26, 2014
<u>AB</u>			ZYDUS PHARMS	<u>0.5MG;0.4MG</u>	<u>A207769 001</u>	May 24, 2018

JALYN

<u>AB</u>	+	!	WOODWARD	<u>0.5MG;0.4MG</u>	<u>N022460 001</u>	Jun 14, 2010
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PRESCRIPTION DRUG PRODUCT LIST

DUVELISIB

CAPSULE; ORAL

COPIKTRA

+ SECURA

15MG

N211155 001 Sep 24, 2018

+

25MG

N211155 002 Sep 24, 2018

DYCLONINE HYDROCHLORIDE

SOLUTION; TOPICAL

DYCLOPRO

! SEPTODONT

0.5%

A200480 001 Nov 20, 2018

!

1%

A200480 002 Nov 20, 2018

ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

+! FERA PHARMS LLC

0.125%

N011963 001

ECONAZOLE NITRATE

AEROSOL, FOAM; TOPICAL

ECOZA

+! RESILIA PHARMS

1%

N205175 001 Oct 24, 2013

CREAM; TOPICAL

ECONAZOLE NITRATEAB ! PADAGIS ISRAEL1%A076479 001 Jun 23, 2004AB TARO1%A076005 001 Nov 26, 2002EDARAVONE

SOLUTION; INTRAVENOUS

RADICAVA

+! MITSUBISHI TANABE

30MG/100ML (0.3MG/ML)

N209176 001 May 05, 2017

+!

60MG/100ML (0.6MG/ML)

N209176 002 Nov 15, 2018

SUSPENSION; ORAL

RADICAVA ORS

+! MITSUBISHI TANABE

105MG/5ML

N215446 001 May 12, 2022

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

EDETATE CALCIUM DISODIUM

! CASPER PHARMA LLC

200MG/ML

A216435 001 May 03, 2023

EDOXYBAN TOSYLATE

TABLET; ORAL

SAVAYSA

+ DAIICHI SANKYO INC

EQ 15MG BASE

N206316 001 Jan 08, 2015

+

EQ 30MG BASE

N206316 002 Jan 08, 2015

+!

EQ 60MG BASE

N206316 003 Jan 08, 2015

EFAVIRENZ

CAPSULE; ORAL

EFAVIRENZ

AUROBINDO PHARMA

50MG

A078064 001 Dec 15, 2017

100MG

A078064 002 Dec 15, 2017

!

200MG

A078064 003 Dec 15, 2017

TABLET; ORAL

EFAVIRENZAB AUROBINDO PHARMA600MGA077673 001 Sep 21, 2018

LTD

AB CIPLA600MGA204766 001 Jun 15, 2018AB ! HETERO LABS LTD III600MGA078886 001 Apr 27, 2018AB STRIDES PHARMA600MGA204869 001 Mar 12, 2018EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

ATRIPLAAB +! GILEAD SCIENCES600MG; 200MG; 300MGN021937 001 Jul 12, 2006EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATEAB AUROBINDO PHARMA600MG; 200MG; 300MGA203041 001 Sep 04, 2018AB CIPLA600MG; 200MG; 300MGA206894 001 Jun 03, 2019AB HETERO LABS LTD V600MG; 200MG; 300MGA203053 001 Jan 24, 2022AB LAURUS600MG; 200MG; 300MGA213541 001 Dec 22, 2021AB MACLEODS PHARMS LTD600MG; 200MG; 300MGA204287 001 Sep 13, 2021AB TEVA PHARMS USA600MG; 200MG; 300MGA091215 001 Nov 09, 2018EFAVIRENZ; EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATEAB STRIDES PHARMA600MG; 200MG; 300MGA201802 001 Oct 03, 2023

PRESCRIPTION DRUG PRODUCT LIST

EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

AB	LAURUS	400MG;300MG;300MG	A213038 001	May 14, 2020
AB		600MG;300MG;300MG	A212786 001	May 14, 2020
SYMFI				
AB	+! MYLAN LABS LTD	600MG;300MG;300MG	N022142 001	Mar 22, 2018
SYMFI LO				
AB	+! MYLAN	400MG;300MG;300MG	N208255 001	Feb 05, 2018

EFINACONAZOLE

SOLUTION;TOPICAL

EFINACONAZOLE

AB	NIVAGEN PHARMS INC	10%	A211969 001	Jun 21, 2021
AB	TEVA PHARMS USA	10%	A211827 001	Dec 16, 2020

JUBLIA

AB	+! BAUSCH	10%	N203567 001	Jun 06, 2014
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EFLORNITHINE HYDROCHLORIDE

TABLET;ORAL

IWILFIN

+!	USWM	EQ 192MG BASE	N215500 001	Dec 13, 2023
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ELACESTRANT DIHYDROCHLORIDE

TABLET;ORAL

ORSERDU

+	STEMLINE THERAP	EQ 86MG BASE	N217639 001	Jan 27, 2023
+!		EQ 345MG BASE	N217639 002	Jan 27, 2023

ELAGOLIX SODIUM

TABLET;ORAL

ORILISSA

+	ABBVIE	EQ 150MG BASE	N210450 001	Jul 23, 2018
+!		EQ 200MG BASE	N210450 002	Jul 23, 2018

ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE; ELAGOLIX SODIUM

CAPSULE;ORAL

ORIAHNN (COPACKAGED)

+!	ABBVIE	EQ 300MG BASE,1MG,0.5MG; EQ 300MG BASE	N213388 001	May 29, 2020
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ELBASVIR; GRAZOPREVIR

TABLET;ORAL

ZEPATIER

+!	MSD SUB MERCK	50MG;100MG	N208261 001	Jan 28, 2016
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ELETRIPTAN HYDROBROMIDE

TABLET;ORAL

ELETRIPTAN HYDROBROMIDE

AB	AJANTA PHARMA LTD	EQ 20MG BASE	A205186 001	Aug 29, 2017
AB		EQ 40MG BASE	A205186 002	Aug 29, 2017
AB	AUROBINDO PHARMA	EQ 20MG BASE	A210708 001	Jan 15, 2019
AB		EQ 40MG BASE	A210708 002	Jan 15, 2019
AB	BEXIMCO PHARMS USA	EQ 20MG BASE	A215467 001	Jul 13, 2022
AB		EQ 40MG BASE	A215467 002	Jul 13, 2022
AB	MYLAN	EQ 20MG BASE	A205152 001	Aug 11, 2017
AB		EQ 40MG BASE	A205152 002	Aug 11, 2017
AB	TEVA PHARMS USA	EQ 20MG BASE	A202040 001	Jun 27, 2017
AB		EQ 40MG BASE	A202040 002	Jun 27, 2017
AB	ZYDUS PHARMS	EQ 20MG BASE	A206409 001	Jun 16, 2017
AB		EQ 40MG BASE	A206409 002	Jun 16, 2017

RELPAK

AB	+ UPJOHN	EQ 20MG BASE	N021016 001	Dec 26, 2002
AB	+!	EQ 40MG BASE	N021016 002	Dec 26, 2002

ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR

GRANULES;ORAL

TRIKAFTA (COPACKAGED)

+	VERTEX PHARMS INC	80MG, 60MG, 40MG;59.5MG	N217660 001	Apr 26, 2023
+!		100MG, 75MG, 50MG;75MG	N217660 002	Apr 26, 2023

TABLET;ORAL

TRIKAFTA (COPACKAGED)

+	VERTEX PHARMS INC	50MG, 37.5MG, 25MG; 75MG	N212273 002	Jun 08, 2021
+!		100MG, 75MG, 50MG; 150MG	N212273 001	Oct 21, 2019

PRESCRIPTION DRUG PRODUCT LIST

ELIGLUSTAT TARTRATE

CAPSULE; ORAL

CERDELGA

AB	+!	GENZYME CORP	<u>EQ 84MG BASE</u>	<u>N205494 001</u>	Aug 19, 2014
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ELIGLUSTAT TARTRATE

AB		ATZANT	<u>EQ 84MG BASE</u>	<u>A212463 001</u>	Sep 08, 2021
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AB		UPSHER SMITH LABS	<u>EQ 84MG BASE</u>	<u>A212420 001</u>	May 02, 2023
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ELTROMBOPAG CHOLINE

TABLET; ORAL

ALVAIZ

	+	TEVA PHARMS INC	EQ 9MG BASE	N216774 001	Nov 29, 2023
	+		EQ 18MG BASE	N216774 002	Nov 29, 2023
	+		EQ 36MG BASE	N216774 003	Nov 29, 2023
	+!		EQ 54MG BASE	N216774 004	Nov 29, 2023

ELTROMBOPAG OLAMINE

FOR SUSPENSION; ORAL

PROMACTA KIT

	+	NOVARTIS	EQ 12.5MG ACID/PACKET	N207027 002	Sep 27, 2018
	+!		EQ 25MG ACID/PACKET	N207027 001	Aug 24, 2015

TABLET; ORAL

PROMACTA

	+	NOVARTIS	EQ 12.5MG ACID	N022291 004	Oct 20, 2011
	+		EQ 25MG ACID	N022291 001	Nov 20, 2008
	+		EQ 50MG ACID	N022291 002	Nov 20, 2008
	+!		EQ 75MG ACID	N022291 003	Sep 08, 2009

ELUXADOLINE

TABLET; ORAL

VIBERZI

	+	ABBVIE	75MG	N206940 001	May 27, 2015
	+!		100MG	N206940 002	May 27, 2015

EMPAGLIFLOZIN

TABLET; ORAL

JARDIANCE

	+	BOEHRINGER INGELHEIM	10MG	N204629 001	Aug 01, 2014
	+!		25MG	N204629 002	Aug 01, 2014

EMPAGLIFLOZIN; LINAGLIPTIN

TABLET; ORAL

GLYXAMBI

	+	BOEHRINGER INGELHEIM	10MG; 5MG	N206073 001	Jan 30, 2015
	+!		25MG; 5MG	N206073 002	Jan 30, 2015

EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TRIJARDY XR

	+	BOEHRINGER INGELHEIM	5MG; 2.5MG; 1GM	N212614 001	Jan 27, 2020
	+		10MG; 5MG; 1GM	N212614 002	Jan 27, 2020
	+		12.5MG; 2.5MG; 1GM	N212614 003	Jan 27, 2020
	+!		25MG; 5MG; 1GM	N212614 004	Jan 27, 2020

EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

SYNJARDY

	+	BOEHRINGER INGELHEIM	5MG; 500MG	N206111 001	Aug 26, 2015
	+		5MG; 1GM	N206111 002	Aug 26, 2015
	+		12.5MG; 500MG	N206111 003	Aug 26, 2015
	+!		12.5MG; 1GM	N206111 004	Aug 26, 2015

TABLET, EXTENDED RELEASE; ORAL

SYNJARDY XR

	+	BOEHRINGER INGELHEIM	5MG; 1GM	N208658 001	Dec 09, 2016
	+		10MG; 1GM	N208658 002	Dec 09, 2016
	+		12.5MG; 1GM	N208658 003	Dec 09, 2016
	+!		25MG; 1GM	N208658 004	Dec 09, 2016

PRESCRIPTION DRUG PRODUCT LIST

EMTRICITABINE

CAPSULE; ORAL

EMTRICITABINE

AB	AUROBINDO PHARMA LTD	200MG	A079188 001	Mar 15, 2023
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AB	CIPLA	200MG	A091168 001	Jul 02, 2018
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EMTRIVA

AB	+! GILEAD	200MG	N021500 001	Jul 02, 2003
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SOLUTION; ORAL

EMTRIVA

+!	GILEAD	10MG/ML	N021896 001	Sep 28, 2005
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EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

ODEFSEY

+!	GILEAD SCIENCES INC	200MG;EQ 25MG BASE;EQ 25MG BASE	N208351 001	Mar 01, 2016
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EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

COMPLERA

+!	GILEAD SCIENCES INC	200MG;EQ 25MG BASE;300MG	N202123 001	Aug 10, 2011
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EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

DESCOVY

+	GILEAD SCIENCES INC	120MG;EQ 15MG BASE	N208215 002	Jan 07, 2022
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+!		200MG;EQ 25MG BASE	N208215 001	Apr 04, 2016
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EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

AB	HETERO LABS LTD III	200MG;300MG	A201806 001	Oct 07, 2021
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EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

AB	AMNEAL PHARMS CO	100MG;150MG	A209721 001	Aug 22, 2018
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AB		133MG;200MG	A209721 002	Aug 22, 2018
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AB		167MG;250MG	A209721 003	Aug 22, 2018
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AB		200MG;300MG	A209721 004	Aug 22, 2018
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AB	AUROBINDO PHARMA	200MG;300MG	A090513 001	Jan 26, 2018
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AB	AUROBINDO PHARMA LTD	100MG;150MG	A211640 001	Mar 09, 2023
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AB		133MG;200MG	A211640 002	Mar 09, 2023
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AB		167MG;250MG	A211640 003	Mar 09, 2023
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AB	LAURUS	200MG;300MG	A212114 001	Jul 26, 2019
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AB	LUPIN LTD	200MG;300MG	A204131 001	Jun 04, 2021
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AB	MACLEODS PHARMS LTD	200MG;300MG	A203442 001	May 15, 2020
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AB	MYLAN	200MG;300MG	A206436 001	Apr 09, 2018
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AB	STRIDES PHARMA	200MG;300MG	A091055 001	Jan 13, 2021
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AB	TEVA PHARMS USA	200MG;300MG	A090894 001	Jun 08, 2017
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AB	ZYDUS PHARMS	100MG;150MG	A212689 002	Jul 01, 2021
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AB		133MG;200MG	A212689 003	Jul 01, 2021
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AB		167MG;250MG	A212689 004	Jul 01, 2021
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AB		200MG;300MG	A212689 001	Feb 28, 2020
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TRUVADA

AB	+ GILEAD	100MG;150MG	N021752 002	Mar 10, 2016
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AB	+	133MG;200MG	N021752 003	Mar 10, 2016
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AB	+	167MG;250MG	N021752 004	Mar 10, 2016
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AB	+!	200MG;300MG	N021752 001	Aug 02, 2004
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ENALAPRIL MALEATE

SOLUTION; ORAL

ENALAPRIL MALEATE

AB	ALKEM LABS LTD	1MG/ML	A213714 001	Mar 30, 2022
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AB	ANNORA PHARMA	1MG/ML	A214467 001	Feb 24, 2022
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AB	AUROBINDO PHARMA	1MG/ML	A216458 001	Jan 08, 2024
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AB	BIONPHARMA	1MG/ML	A212408 001	Aug 10, 2021
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EPANED

AB	+! AZURITY	1MG/ML	N208686 001	Sep 20, 2016
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TABLET; ORAL

ENALAPRIL MALEATE

AB	HERITAGE PHARMA	2.5MG	A075479 001	Aug 22, 2000
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AB		5MG	A075479 002	Aug 22, 2000
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AB		10MG	A075479 003	Aug 22, 2000
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AB		20MG	A075479 004	Aug 22, 2000
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AB	PRINSTON INC	2.5MG	A213273 001	Jul 07, 2022
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AB		5MG	A213273 002	Jul 07, 2022
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PRESCRIPTION DRUG PRODUCT LIST

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

<u>AB</u>		<u>10MG</u>	<u>A213273 003</u>	Jul 07, 2022
<u>AB</u>		<u>20MG</u>	<u>A213273 004</u>	Jul 07, 2022
<u>AB</u>	SANDOZ INC	<u>2.5MG</u>	<u>A075496 001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075496 002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075459 001</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075459 002</u>	Aug 22, 2000
<u>AB</u>	TARO	<u>2.5MG</u>	<u>A075657 001</u>	Jan 23, 2001
<u>AB</u>		<u>5MG</u>	<u>A075657 002</u>	Jan 23, 2001
<u>AB</u>		<u>10MG</u>	<u>A075657 003</u>	Jan 23, 2001
<u>AB</u>		<u>20MG</u>	<u>A075657 004</u>	Jan 23, 2001
<u>AB</u>	WOCKHARDT LTD	<u>2.5MG</u>	<u>A075483 001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075483 002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075483 003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075483 004</u>	Aug 22, 2000
<u>VASOTEC</u>				
<u>AB</u>	+ BAUSCH	<u>2.5MG</u>	<u>N018998 005</u>	Jul 26, 1988
<u>AB</u>	+	<u>5MG</u>	<u>N018998 001</u>	Dec 24, 1985
<u>AB</u>	+	<u>10MG</u>	<u>N018998 002</u>	Dec 24, 1985
<u>AB</u>	+!	<u>20MG</u>	<u>N018998 003</u>	Dec 24, 1985

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	COSETTE	<u>5MG;12.5MG</u>	<u>A075727 001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075727 002</u>	Sep 18, 2001
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG;12.5MG</u>	<u>A075909 001</u>	Oct 15, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075909 002</u>	Oct 15, 2001
<u>AB</u>	TARO PHARM INDS	<u>5MG;12.5MG</u>	<u>A075788 001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075788 002</u>	Sep 18, 2001
<u>VASERETIC</u>				
<u>AB</u>	+ BAUSCH	<u>5MG;12.5MG</u>	<u>N019221 003</u>	Jul 12, 1995
<u>AB</u>	+!	<u>10MG;25MG</u>	<u>N019221 001</u>	Oct 31, 1986

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

<u>AP</u>	CHARTWELL INJECTABLE	<u>1.25MG/ML</u>	<u>A075634 001</u>	Aug 22, 2000
<u>AP</u>	DR REDDYS	<u>1.25MG/ML</u>	<u>A075578 001</u>	Aug 22, 2000
<u>AP</u>	! HIKMA FARMACEUTICA	<u>1.25MG/ML</u>	<u>A078687 001</u>	Dec 23, 2008
<u>AP</u>	HOSPIRA	<u>1.25MG/ML</u>	<u>A075458 001</u>	Aug 22, 2000

ENASIDENIB MESYLATE

TABLET; ORAL

IDHIFA

+	BRISTOL MYERS SQUIBB	EQ 50MG BASE	N209606 001	Aug 01, 2017
+	!	EQ 100MG BASE	N209606 002	Aug 01, 2017

ENCORAFENIB

CAPSULE; ORAL

BRAFTOVI

+	! ARRAY BIOPHARMA INC	75MG	N210496 002	Jun 27, 2018
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ENFUVRTIDE

INJECTABLE; SUBCUTANEOUS

FUZEON

+	! ROCHE	90MG/VIAL	N021481 001	Mar 13, 2003
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ENOXAPARIN SODIUM

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

ENOXAPARIN SODIUM

<u>AB</u>	AMPHASTAR PHARMS INC	<u>300MG/3ML (100MG/ML)</u>	<u>A208600 001</u>	Mar 14, 2019
<u>AB</u>	NANJING KING-FRIEND	<u>300MG/3ML (100MG/ML)</u>	<u>A214856 001</u>	Jun 14, 2022
<u>AB</u>	SANDOZ INC	<u>300MG/3ML (100MG/ML)</u>	<u>A078660 001</u>	Nov 28, 2011

LOVENOX

<u>AB</u>	+ SANOFI AVENTIS US	<u>300MG/3ML (100MG/ML)</u>	<u>N020164 009</u>	Jan 23, 2003
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INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

<u>AP</u>	AMPHASTAR PHARM	<u>30MG/0.3ML (100MG/ML)</u>	<u>A076684 001</u>	Sep 19, 2011
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A076684 002</u>	Sep 19, 2011
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A076684 003</u>	Sep 19, 2011

PRESCRIPTION DRUG PRODUCT LIST

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A076684 004</u>	Sep 19, 2011
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A076684 005</u>	Sep 19, 2011
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A076684 006</u>	Sep 19, 2011
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A076684 007</u>	Sep 19, 2011
<u>AP</u>	BE PHARMS	<u>30MG/0.3ML (100MG/ML)</u>	<u>A214646 001</u>	Jun 06, 2023
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A214646 002</u>	Jun 06, 2023
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A214646 003</u>	Jun 06, 2023
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A214646 004</u>	Jun 06, 2023
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A214646 005</u>	Jun 06, 2023
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A214646 006</u>	Jun 06, 2023
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A214646 007</u>	Jun 06, 2023
<u>AP</u>	GLAND	<u>30MG/0.3ML (100MG/ML)</u>	<u>A078990 001</u>	Sep 28, 2018
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A078990 002</u>	Sep 28, 2018
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A078990 003</u>	Sep 28, 2018
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A078990 004</u>	Sep 28, 2018
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A078990 005</u>	Sep 28, 2018
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A078990 006</u>	Sep 28, 2018
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A078990 007</u>	Sep 28, 2018
<u>AP</u>	NANJING KING-FRIEND	<u>30MG/0.3ML (100MG/ML)</u>	<u>A206834 001</u>	Nov 29, 2019
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A206834 002</u>	Nov 29, 2019
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A206834 003</u>	Nov 29, 2019
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A206834 004</u>	Nov 29, 2019
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A206834 005</u>	Nov 29, 2019
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A206834 006</u>	Nov 29, 2019
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A206834 007</u>	Nov 29, 2019
<u>AP</u>	SANDOZ	<u>30MG/0.3ML (100MG/ML)</u>	<u>A077857 002</u>	Jul 23, 2010
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A077857 003</u>	Jul 23, 2010
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A077857 004</u>	Jul 23, 2010
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A077857 005</u>	Jul 23, 2010
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A077857 001</u>	Jul 23, 2010
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A077857 006</u>	Jul 23, 2010
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A077857 007</u>	Jul 23, 2010
<u>AP</u>	SHENZHEN TECHDOW	<u>30MG/0.3ML (100MG/ML)</u>	<u>A205660 001</u>	Mar 15, 2023
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A205660 002</u>	Mar 15, 2023
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A205660 003</u>	Mar 15, 2023
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A205660 004</u>	Mar 15, 2023
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A205660 005</u>	Mar 15, 2023
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A205660 006</u>	Mar 15, 2023
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A205660 007</u>	Mar 15, 2023
<u>AP</u>	ZYDUS PHARMS	<u>30MG/0.3ML (100MG/ML)</u>	<u>A076726 001</u>	Jun 23, 2014
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A076726 002</u>	Jun 23, 2014
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A076726 003</u>	Jun 23, 2014
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A076726 004</u>	Jun 23, 2014
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A076726 005</u>	Jun 23, 2014
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A076726 006</u>	Jun 23, 2014
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A076726 007</u>	Jun 23, 2014
<u>LOVENOX (PRESERVATIVE FREE)</u>				
<u>AP</u>	+	SANOFI AVENTIS US	<u>30MG/0.3ML (100MG/ML)</u>	<u>N020164 001</u> Mar 29, 1993
<u>AP</u>	+		<u>40MG/0.4ML (100MG/ML)</u>	<u>N020164 002</u> Jan 30, 1998
<u>AP</u>	+		<u>60MG/0.6ML (100MG/ML)</u>	<u>N020164 003</u> Mar 27, 1998
<u>AP</u>	+		<u>80MG/0.8ML (100MG/ML)</u>	<u>N020164 004</u> Mar 27, 1998
<u>AP</u>	+		<u>100MG/ML (100MG/ML)</u>	<u>N020164 005</u> Mar 27, 1998
<u>AP</u>	+		<u>120MG/0.8ML (150MG/ML)</u>	<u>N020164 007</u> Jun 02, 2000
<u>AP</u>	+		<u>150MG/ML (150MG/ML)</u>	<u>N020164 008</u> Jun 02, 2000

ENTACAPONE

TABLET; ORAL

COMTAN

<u>AB</u>	+	ORION PHARMA	<u>200MG</u>	<u>N020796 001</u> Oct 19, 1999
<u>ENTACAPONE</u>				
<u>AB</u>		AJANTA PHARMA LTD	<u>200MG</u>	<u>A205792 001</u> Aug 31, 2017
<u>AB</u>		ALEMBIC	<u>200MG</u>	<u>A212601 001</u> Jan 04, 2022
<u>AB</u>		AUROBINDO PHARMA LTD	<u>200MG</u>	<u>A203437 001</u> Jun 19, 2015
<u>AB</u>		MACLEODS PHARMS LTD	<u>200MG</u>	<u>A207210 001</u> Jun 05, 2017
<u>AB</u>		SUN PHARM	<u>200MG</u>	<u>A090690 001</u> Jul 16, 2012
<u>AB</u>		SUNSHINE	<u>200MG</u>	<u>A206669 001</u> Oct 03, 2018
<u>AB</u>		WOCKHARDT BIO AG	<u>200MG</u>	<u>A078941 001</u> Aug 16, 2012

PRESCRIPTION DRUG PRODUCT LIST

ENTECAVIR

SOLUTION; ORAL

BARACLUDE

+	BRISTOL MYERS SQUIBB	0.05MG/ML	N021798	001	Mar 29, 2005
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TABLET; ORAL

BARACLUDE

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>0.5MG</u>	<u>N021797</u>	<u>001</u>	Mar 29, 2005
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<u>AB</u>	+		<u>1MG</u>	<u>N021797</u>	<u>002</u>	Mar 29, 2005
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ENTECAVIR

<u>AB</u>		AMNEAL PHARMS	<u>0.5MG</u>	<u>A206652</u>	<u>001</u>	Nov 12, 2015
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<u>AB</u>			<u>1MG</u>	<u>A206652</u>	<u>002</u>	Nov 12, 2015
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<u>AB</u>		AUROBINDO PHARMA	<u>0.5MG</u>	<u>A206217</u>	<u>001</u>	Aug 26, 2015
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<u>AB</u>			<u>1MG</u>	<u>A206217</u>	<u>002</u>	Aug 26, 2015
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<u>AB</u>		BRECKENRIDGE	<u>0.5MG</u>	<u>A208721</u>	<u>001</u>	Mar 15, 2018
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<u>AB</u>			<u>1MG</u>	<u>A208721</u>	<u>002</u>	Mar 15, 2018
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<u>AB</u>		BRIGHTGENE	<u>0.5MG</u>	<u>A212126</u>	<u>001</u>	Sep 25, 2019
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<u>AB</u>			<u>1MG</u>	<u>A212126</u>	<u>002</u>	Sep 25, 2019
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<u>AB</u>		CIPLA	<u>0.5MG</u>	<u>A206872</u>	<u>001</u>	Dec 06, 2016
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<u>AB</u>			<u>1MG</u>	<u>A206872</u>	<u>002</u>	Dec 06, 2016
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<u>AB</u>		HETERO LABS LTD V	<u>0.5MG</u>	<u>A205740</u>	<u>001</u>	Aug 21, 2015
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<u>AB</u>			<u>1MG</u>	<u>A205740</u>	<u>002</u>	Aug 21, 2015
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<u>AB</u>		PRINSTON INC	<u>0.5MG</u>	<u>A208782</u>	<u>001</u>	Oct 10, 2017
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<u>AB</u>			<u>1MG</u>	<u>A208782</u>	<u>002</u>	Oct 10, 2017
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<u>AB</u>		VITRUVIAS	<u>0.5MG</u>	<u>A212106</u>	<u>001</u>	Aug 10, 2020
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<u>AB</u>			<u>1MG</u>	<u>A212106</u>	<u>002</u>	Aug 10, 2020
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<u>AB</u>		ZYDUS PHARMS	<u>0.5MG</u>	<u>A206745</u>	<u>001</u>	Jun 23, 2017
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<u>AB</u>			<u>1MG</u>	<u>A206745</u>	<u>002</u>	Jun 23, 2017
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ENTRECTINIB

CAPSULE; ORAL

ROZLYTREK

+	GENENTECH INC	100MG	N212725	001	Aug 15, 2019
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+		200MG	N212725	002	Aug 15, 2019
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PELLETS; ORAL

ROZLYTREK

+	GENENTECH INC	50MG/PACKET	N218550	001	Oct 20, 2023
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ENZALUTAMIDE

CAPSULE; ORAL

XTANDI

+	ASTELLAS	40MG	N203415	001	Aug 31, 2012
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TABLET; ORAL

XTANDI

+	ASTELLAS	40MG	N213674	001	Aug 04, 2020
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+		80MG	N213674	002	Aug 04, 2020
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EPHEDRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

REZIPRES

+	DR REDDYS LABS SA	23.5MG/5ML (4.7MG/ML)	N213536	001	Jun 14, 2021
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+		47MG/10ML (4.7MG/ML)	N213536	004	Dec 07, 2023
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EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

AKOVAZ

<u>AP</u>	+	EXELA PHARMA	<u>50MG/ML (50MG/ML)</u>	<u>N208289</u>	<u>001</u>	Apr 29, 2016
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CORPHEDRA

<u>AP</u>		PAR STERILE PRODUCTS	<u>50MG/ML (50MG/ML)</u>	<u>N208943</u>	<u>001</u>	Jan 27, 2017
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EPHEDRINE SULFATE

<u>AP</u>		AMNEAL	<u>50MG/ML (50MG/ML)</u>	<u>A212932</u>	<u>001</u>	Oct 23, 2019
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<u>AP</u>		DR REDDYS	<u>50MG/ML (50MG/ML)</u>	<u>A212649</u>	<u>001</u>	Oct 03, 2020
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<u>AP</u>		EUGIA PHARMA	<u>50MG/ML (50MG/ML)</u>	<u>A214579</u>	<u>001</u>	Jun 14, 2021
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<u>AP</u>		FRESENIUS KABI USA	<u>50MG/ML (50MG/ML)</u>	<u>A209646</u>	<u>001</u>	Aug 04, 2020
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<u>AP</u>		GLAND PHARMA LTD	<u>50MG/ML (50MG/ML)</u>	<u>A216146</u>	<u>001</u>	Feb 25, 2022
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<u>AP</u>		HIKMA	<u>50MG/ML (50MG/ML)</u>	<u>A214334</u>	<u>001</u>	Dec 15, 2020
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<u>AP</u>		MANKIND PHARMA	<u>50MG/ML (50MG/ML)</u>	<u>A216129</u>	<u>001</u>	Apr 14, 2022
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<u>AP</u>		SAGENT PHARMS INC	<u>50MG/ML (50MG/ML)</u>	<u>A214528</u>	<u>001</u>	Mar 09, 2023
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<u>AP</u>		SANDOZ	<u>50MG/ML (50MG/ML)</u>	<u>A209784</u>	<u>001</u>	Aug 23, 2017
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<u>AP</u>		XIROMED	<u>50MG/ML (50MG/ML)</u>	<u>A215825</u>	<u>001</u>	Apr 21, 2022
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AKOVAZ

+	EXELA PHARMA	25MG/5ML (5MG/ML)	N208289	002	Aug 02, 2021
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PRESCRIPTION DRUG PRODUCT LIST

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

EMERPHED

+!	NEXUS	25MG/5ML (5MG/ML)	N213407 002	Feb 28, 2023
+!		50MG/10ML (5MG/ML)	N213407 001	Apr 17, 2020

EPHEDRINE SULFATE

+!	OPERAND PHARMS	25MG/5ML (5MG/ML)	N213994 002	Apr 22, 2022
+!		50MG/10ML (5MG/ML)	N213994 001	Oct 16, 2020

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

EPINASTINE HYDROCHLORIDE

AT	APOTEX	0.05%	A090919 001	Oct 31, 2011
AT	BRECKENRIDGE	0.05%	A090870 001	Mar 14, 2011
AT	! SOMERSET THERAPS LLC	0.05%	A090951 001	Oct 31, 2011

EPINEPHRINE

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE (AUTOINJECTOR)

AB	TEVA PHARMS USA	0.15MG/DELIVERY	A090589 002	Aug 16, 2018
AB		0.3MG/DELIVERY	A090589 001	Aug 16, 2018

EPIPEN

AB	+! MYLAN SPECIALITY LP	0.3MG/DELIVERY	N019430 001	Dec 22, 1987
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EPIPEN JR.

AB	+! MYLAN SPECIALITY LP	0.15MG/DELIVERY	N019430 002	Dec 22, 1987
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ADRENACLICK

BX	+! IMPAX	EQ 0.15MG/DELIVERY	N020800 003	Nov 25, 2009
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BX	+!	EQ 0.3MG/DELIVERY	N020800 004	Nov 25, 2009
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SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

ADRENALIN

AP	+! PAR STERILE PRODUCTS	EQ 30MG BASE/30ML (EQ 1MG BASE/ML)	N204640 001	Dec 18, 2013
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EPINEPHRINE

AP	INTL MEDICATION SYS	EQ 30MG BASE/30ML (EQ 1MG BASE/ML)	A211880 001	Apr 24, 2020
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ADRENALIN

+!	PAR STERILE PRODUCTS	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N204200 001	Dec 07, 2012
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EPINEPHRINE

+!	BPI LABS	10MG/10ML (1MG/ML)	N205029 002	Feb 04, 2022
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SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

AUVI-Q

BX	+! KALEO INC	EQ 0.15MG/DELIVERY	N201739 002	Aug 10, 2012
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BX	+	EQ 0.3MG/DELIVERY	N201739 001	Aug 10, 2012
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+		EQ 0.1MG/DELIVERY	N201739 003	Nov 17, 2017
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SYMJEPI

+!	ADAMIS PHARMS CORP	0.3MG/0.3ML (0.3MG/0.3ML)	N207534 001	Jun 15, 2017
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SOLUTION; INTRAVENOUS

ADRENALIN

+!	PAR STERILE PRODUCTS	2MG/250ML (8MCG/ML)	N215875 001	Apr 21, 2023
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+!		4MG/250ML (16MCG/ML)	N215875 002	Apr 21, 2023
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+!		5MG/250ML (20MCG/ML)	N215875 003	Apr 21, 2023
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+!		8MG/250ML (32MCG/ML)	N215875 004	Apr 21, 2023
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+!		10MG/250ML (40MCG/ML)	N215875 005	Apr 21, 2023
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EPINEPHRINE

+!	BPI LABS	1MG/ML (1MG/ML)	N205029 003	May 12, 2023
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+!	HOSPIRA	1MG/10ML (0.1MG/ML)	N209359 001	Nov 05, 2019
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+!	INTL MEDICATION SYS	1MG/10ML (0.1MG/ML)	N211363 001	Aug 15, 2022
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SOLUTION; INTRAVENOUS, INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE

+!	BPI LABS	1MG/ML (1MG/ML)	N205029 001	Jul 29, 2014
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EPINEPHRINE BITARTRATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIGNOSPAN FORTE

+!	DEPROCO	EQ 0.02MG BASE/ML; 2%	A088389 001	Jan 22, 1985
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LIGNOSPAN STANDARD

+!	DEPROCO	EQ 0.01MG BASE/ML; 2%	A088390 001	Jan 22, 1985
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PRESCRIPTION DRUG PRODUCT LIST

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE DENTAL

<u>AP</u>	<u>+!</u>	<u>DENTSPLY PHARM</u>	<u>0.005MG/ML;4%</u>	<u>N021383</u>	<u>001</u>	
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PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

<u>AP</u>		<u>SEPTODONT INC</u>	<u>0.005MG/ML;4%</u>	<u>A078959</u>	<u>001</u>	<u>Aug 30, 2011</u>
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EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

<u>AP</u>		<u>B BRAUN MEDICAL INC</u>	<u>0.005MG/ML;1.5%</u>	<u>A208475</u>	<u>001</u>	<u>Sep 08, 2021</u>
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<u>AP</u>		<u>HOSPIRA</u>	<u>0.005MG/ML;0.5%</u>	<u>A089635</u>	<u>001</u>	<u>Jun 21, 1988</u>
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<u>AP</u>			<u>0.005MG/ML;1.5%</u>	<u>A088571</u>	<u>001</u>	<u>Sep 13, 1985</u>
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<u>AP</u>			<u>0.005MG/ML;1.5%</u>	<u>A089645</u>	<u>001</u>	<u>Jun 21, 1988</u>
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<u>AP</u>			<u>0.005MG/ML;2%</u>	<u>A089651</u>	<u>001</u>	<u>Jun 21, 1988</u>
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<u>AP</u>			<u>0.01MG/ML;1%</u>	<u>A089644</u>	<u>001</u>	<u>Jun 21, 1988</u>
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XYLOCAINE W/ EPINEPHRINE

<u>AP</u>	<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>0.005MG/ML;0.5%</u>	<u>N006488</u>	<u>012</u>	
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<u>AP</u>	<u>+!</u>		<u>0.005MG/ML;1.5%</u>	<u>N006488</u>	<u>017</u>	
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<u>AP</u>	<u>+!</u>		<u>0.005MG/ML;2%</u>	<u>N006488</u>	<u>019</u>	<u>Nov 13, 1986</u>
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<u>AP</u>	<u>+!</u>		<u>0.01MG/ML;1%</u>	<u>N006488</u>	<u>004</u>	
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LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

<u>!</u>	<u>HOSPIRA</u>	<u>0.01MG/ML;2%</u>	<u>A089646</u>	<u>001</u>	<u>Jun 21, 1988</u>
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XYLOCAINE W/ EPINEPHRINE

<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>0.005MG/ML;1%</u>	<u>N006488</u>	<u>018</u>	<u>Nov 13, 1986</u>
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<u>+!</u>		<u>0.02MG/ML;2%</u>	<u>N006488</u>	<u>005</u>	
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EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ELLEENCE

<u>AP</u>	<u>+!</u>	<u>PFIZER INC</u>	<u>200MG/100ML (2MG/ML)</u>	<u>N050778</u>	<u>001</u>	<u>Sep 15, 1999</u>
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<u>AP</u>	<u>+</u>		<u>50MG/25ML (2MG/ML)</u>	<u>N050778</u>	<u>002</u>	<u>Sep 15, 1999</u>
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EPIRUBICIN HYDROCHLORIDE

<u>AP</u>		<u>AKORN</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A090163</u>	<u>001</u>	<u>Jun 24, 2009</u>
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<u>AP</u>		<u>CIPLA LTD</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A065361</u>	<u>001</u>	<u>Oct 22, 2007</u>
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<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065361</u>	<u>002</u>	<u>Oct 22, 2007</u>
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<u>AP</u>		<u>HIKMA</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A065289</u>	<u>001</u>	<u>Jun 27, 2007</u>
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<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065289</u>	<u>002</u>	<u>Jun 27, 2007</u>
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<u>AP</u>		<u>HISUN PHARM</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A090075</u>	<u>001</u>	<u>Mar 25, 2010</u>
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<u>AP</u>		<u>HANGZHOU</u>				
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<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A090075</u>	<u>002</u>	<u>Mar 25, 2010</u>
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<u>AP</u>		<u>IMPAX LABS INC</u>	<u>50MG/25ML (2MG/ML)</u>	<u>A065331</u>	<u>001</u>	<u>Aug 09, 2007</u>
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<u>AP</u>			<u>200MG/100ML (2MG/ML)</u>	<u>A065331</u>	<u>002</u>	<u>Aug 09, 2007</u>
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EPLERENONE

TABLET; ORAL

EPLERENONE

<u>AB</u>		<u>ACCORD HLTHCARE</u>	<u>25MG</u>	<u>A206922</u>	<u>001</u>	<u>Jul 13, 2017</u>
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<u>AB</u>			<u>50MG</u>	<u>A206922</u>	<u>002</u>	<u>Jul 13, 2017</u>
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<u>AB</u>		<u>ANNORA PHARMA</u>	<u>25MG</u>	<u>A213812</u>	<u>001</u>	<u>Jun 02, 2023</u>
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<u>AB</u>			<u>50MG</u>	<u>A213812</u>	<u>002</u>	<u>Jun 02, 2023</u>
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<u>AB</u>		<u>BRECKENRIDGE</u>	<u>25MG</u>	<u>A208283</u>	<u>001</u>	<u>Sep 14, 2018</u>
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<u>AB</u>			<u>50MG</u>	<u>A208283</u>	<u>002</u>	<u>Sep 14, 2018</u>
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<u>AB</u>		<u>CHARTWELL RX</u>	<u>25MG</u>	<u>A078482</u>	<u>001</u>	<u>Jul 30, 2008</u>
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<u>AB</u>			<u>50MG</u>	<u>A078482</u>	<u>002</u>	<u>Jul 30, 2008</u>
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<u>AB</u>		<u>PIRAMAL HLTHCARE UK</u>	<u>25MG</u>	<u>A212765</u>	<u>001</u>	<u>Aug 10, 2020</u>
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<u>AB</u>			<u>50MG</u>	<u>A212765</u>	<u>002</u>	<u>Aug 10, 2020</u>
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<u>AB</u>		<u>PRASCO</u>	<u>25MG</u>	<u>A203896</u>	<u>001</u>	<u>Feb 02, 2017</u>
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<u>AB</u>			<u>50MG</u>	<u>A203896</u>	<u>002</u>	<u>Feb 02, 2017</u>
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<u>AB</u>		<u>RISING</u>	<u>25MG</u>	<u>A214663</u>	<u>001</u>	<u>Mar 23, 2022</u>
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<u>AB</u>			<u>50MG</u>	<u>A214663</u>	<u>002</u>	<u>Mar 23, 2022</u>
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<u>AB</u>		<u>SANDOZ</u>	<u>25MG</u>	<u>A078510</u>	<u>001</u>	<u>Aug 01, 2008</u>
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<u>AB</u>			<u>50MG</u>	<u>A078510</u>	<u>002</u>	<u>Aug 01, 2008</u>
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<u>AB</u>		<u>WESTMINSTER PHARMS</u>	<u>25MG</u>	<u>A207842</u>	<u>001</u>	<u>Oct 25, 2021</u>
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<u>AB</u>			<u>50MG</u>	<u>A207842</u>	<u>002</u>	<u>Oct 25, 2021</u>
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INSPRA

<u>AB</u>	<u>+</u>	<u>UPJOHN</u>	<u>25MG</u>	<u>N021437</u>	<u>001</u>	<u>Sep 27, 2002</u>
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<u>AB</u>	<u>+!</u>		<u>50MG</u>	<u>N021437</u>	<u>002</u>	<u>Sep 27, 2002</u>
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PRESCRIPTION DRUG PRODUCT LIST

EPLONTERSEN SODIUM

SOLUTION; SUBCUTANEOUS

WAINUA

+! IONIS PHARMS INC EQ 45MG BASE/0.8ML (EQ 45MG BASE/0.8ML) N217388 001 Dec 21, 2023

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

EPOPROSTENOL SODIUMAP1 HONG KONG EQ 0.5MG BASE/VIAL A078396 001 Apr 23, 2008AP1 EQ 1.5MG BASE/VIAL A078396 002 Apr 23, 2008FLOLANAP1 +! GLAXOSMITHKLINE LLC EQ 0.5MG BASE/VIAL N020444 001 Sep 20, 1995AP1 +! EQ 1.5MG BASE/VIAL N020444 002 Sep 20, 1995EPOPROSTENOL SODIUMAP2 ! SUN PHARM EQ 0.5MG BASE/VIAL A210473 001 Jan 15, 2021AP2 EQ 1.5MG BASE/VIAL A210473 002 Jan 15, 2021VELETRIAP2 + ACTELION EQ 0.5MG BASE/VIAL N022260 002 Jun 28, 2012AP2 +! EQ 1.5MG BASE/VIAL N022260 001 Jun 27, 2008EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDEAP ACCORD HLTHCARE 2MG/ML A205557 001 Nov 06, 2017AP 75MG/100ML A205557 002 Nov 06, 2017AP BAXTER HLTHCARE 75MG/100ML A208554 002 Nov 23, 2018

CORP

AP EUGIA PHARMA 2MG/ML A206127 001 Dec 08, 2015AP 75MG/100ML A206127 002 Dec 08, 2015AP ! MYLAN LABS LTD 2MG/ML A203258 001 Jul 20, 2018AP ! 75MG/100ML A203258 002 Jul 20, 2018AP SAGENT PHARMS INC 2MG/ML A204693 001 Mar 07, 2018AP 75MG/100ML A204693 002 Mar 07, 2018AP SHUANGCHENG 2MG/ML A213081 001 Sep 07, 2021AP 75MG/100ML A213081 002 Apr 20, 2022AP SLATE RUN PHARMA 2MG/ML A209864 001 Jan 25, 2019AP 75MG/100ML A209864 002 Jan 25, 2019AP TEVA PHARMS USA 2MG/ML A090854 001 Jun 12, 2015ERAVACYCLINE DIHYDROCHLORIDE

POWDER; INTRAVENOUS

KERAVA

+! TETRAPHASE PHARMS EQ 50MG BASE/VIAL N211109 001 Aug 27, 2018

+! EQ 100MG BASE/VIAL N211109 002 Jun 03, 2020

ERDAFITINIB

TABLET; ORAL

BALVERSA

+ JANSSEN BIOTECH 3MG N212018 001 Apr 12, 2019

+ 4MG N212018 002 Apr 12, 2019

+! 5MG N212018 003 Apr 12, 2019

ERGOCALCIFEROL

CAPSULE; ORAL

DRISDOLAA + SS PHARMA 50,000 IU N003444 001ERGOCALCIFEROLAA CHARTWELL RX 50,000 IU A040833 001 May 20, 2009AA PURACAP PHARM LLC 50,000 IU A204276 001 Dec 07, 2018AA ! STRIDES PHARMA 50,000 IU A090455 001 Aug 03, 2010VITAMIN DAA BIONPHARMA 50,000 IU A080704 001ERGOLOID MESYLATES

TABLET; ORAL

ERGOLOID MESYLATES

! SUN PHARM 1MG A081113 001 Oct 31, 1991

INDUSTRIES

ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL

ERGOMAR

! TERSERA 2MG A087693 001 Feb 24, 1983

PRESCRIPTION DRUG PRODUCT LIST

ERIBULIN MESYLATE

SOLUTION; INTRAVENOUS

HALAVEN

+! EISAI INC

1MG/2ML (0.5MG/ML)

N201532 001 Nov 15, 2010

ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

ERLOTINIB HYDROCHLORIDE

<u>AB</u>	ALEMBIC	<u>EQ 25MG BASE</u>	<u>A214719 001</u>	Jul 08, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A214719 002</u>	Jul 08, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214719 003</u>	Jul 08, 2021
<u>AB</u>	APOTEX	<u>EQ 25MG BASE</u>	<u>A208396 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A208396 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208396 003</u>	Nov 05, 2019
<u>AB</u>	MSN	<u>EQ 25MG BASE</u>	<u>A214366 001</u>	May 10, 2021
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A214366 002</u>	May 10, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214366 003</u>	May 10, 2021
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 25MG BASE</u>	<u>A208488 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A208488 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208488 003</u>	Nov 05, 2019
<u>AB</u>	RISING	<u>EQ 25MG BASE</u>	<u>A091002 001</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091002 002</u>	Jun 11, 2014
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091002 003</u>	Jun 11, 2014
<u>AB</u>	SHILPA	<u>EQ 25MG BASE</u>	<u>A211960 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A211960 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211960 003</u>	Nov 05, 2019
<u>AB</u>	SUN PHARM	<u>EQ 25MG BASE</u>	<u>A210300 001</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A210300 002</u>	Nov 05, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A210300 003</u>	Nov 05, 2019
<u>AB</u>	TEVA PHARMS USA INC	<u>EQ 25MG BASE</u>	<u>A091059 001</u>	Nov 09, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091059 002</u>	Aug 28, 2015
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091059 003</u>	Aug 28, 2015
<u>AB</u>	ZYDUS PHARMS	<u>EQ 25MG BASE</u>	<u>A213065 001</u>	Apr 16, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213065 002</u>	Apr 16, 2020
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A213065 003</u>	Apr 16, 2020

ERTAPENEM SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

ERTAPENEM SODIUM

<u>AP</u>	ACS DOBFAR SPA	<u>EQ 1GM BASE/VIAL</u>	<u>A208790 001</u>	Apr 16, 2018
<u>AP</u>	EUGIA PHARMA	<u>EQ 1GM BASE/VIAL</u>	<u>A209133 001</u>	Jun 25, 2018
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A212040 001</u>	Mar 26, 2021
<u>AP</u>	SAVIOR LIFETEC CORP	<u>EQ 1GM BASE/VIAL</u>	<u>A207647 001</u>	Mar 19, 2019
<u>AP</u>	SUN PHARM	<u>EQ 1GM BASE/VIAL</u>	<u>A209145 001</u>	May 02, 2023

INVANZ

<u>AP</u>	+! MSD SUB MERCK	<u>EQ 1GM BASE/VIAL</u>	<u>N021337 001</u>	Nov 21, 2001
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ERTUGLIFLOZIN

TABLET; ORAL

STEGLATRO

+ MSD SUB MERCK

5MG

N209803 001 Dec 19, 2017

+!

15MG

N209803 002 Dec 19, 2017

ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

SEGLUROMET

+ MSD SUB MERCK

2.5MG; 500MG

N209806 001 Dec 19, 2017

+

2.5MG; 1GM

N209806 002 Dec 19, 2017

+

7.5MG; 500MG

N209806 003 Dec 19, 2017

+!

7.5MG; 1GM

N209806 004 Dec 19, 2017

ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

STEGLUJAN

+ MSD SUB MERCK

5MG; EQ 100MG BASE

N209805 001 Dec 19, 2017

+!

15MG; EQ 100MG BASE

N209805 002 Dec 19, 2017

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYC

<u>AB</u>	+! DR REDDYS LABS SA	<u>250MG</u>	<u>N050536 001</u>	
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ERYTHROMYCIN

<u>AB</u>	ARBOR PHARMS LLC	<u>250MG</u>	<u>A062746 001</u>	Dec 22, 1986
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PRESCRIPTION DRUG PRODUCT LIST

ERYTHROMYCIN

GEL; TOPICAL

ERYGEL

AT	+!	MYLAN	2%	N050617	001	Oct 21, 1987
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ERYTHROMYCIN

AT		FOUGERA PHARMS	2%	A064184	001	Sep 30, 1997
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AT		PADAGIS US	2%	A063211	001	Jan 29, 1993
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OINTMENT; OPHTHALMIC

ERYTHROMYCIN

AT		BAUSCH AND LOMB	0.5%	A064067	001	Jul 29, 1994
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AT	!	PADAGIS US	0.5%	A062447	001	Sep 26, 1983
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AT		SENTISS	0.5%	A064030	001	Jul 18, 1996
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SOLUTION; TOPICAL

ERYTHRA-DETM

AT		MICRO LABS	2%	A062687	001	Feb 05, 1988
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ERYTHROMYCIN

AT	!	PADAGIS US	2%	A063038	001	Jan 11, 1991
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SWAB; TOPICAL

ERYTHROMYCIN

AT		EPIC PHARMA LLC	2%	A090215	001	May 12, 2010
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AT	!	PADAGIS US	2%	A064126	001	Jul 03, 1996
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TABLET; ORAL

ERYTHROMYCIN

AB		ALEMBIC	250MG	A215661	001	Aug 24, 2023
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AB			500MG	A215661	002	Aug 24, 2023
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AB		ALKEM LABS LTD	250MG	A216066	001	Jul 13, 2022
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AB			500MG	A216066	002	Jul 13, 2022
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AB		AMNEAL PHARMS CO	250MG	A209720	001	Mar 09, 2018
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AB			500MG	A209720	002	Mar 09, 2018
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AB		ARBOR PHARMS LLC	250MG	A061621	001	
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AB			500MG	A061621	002	
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AB		CADILA PHARMS LTD	250MG	A213628	001	Jun 28, 2021
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AB	!		500MG	A213628	002	Jun 28, 2021
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AB		TEVA PHARMS USA INC	250MG	A214549	001	Feb 11, 2021
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AB			500MG	A214549	002	Feb 11, 2021
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AB		TORRENT	250MG	A212015	001	Jul 06, 2020
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AB			500MG	A212015	002	Jul 06, 2020
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AB		ZYDUS	250MG	A212693	001	Mar 07, 2023
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AB			500MG	A212693	002	Mar 07, 2023
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TABLET, DELAYED RELEASE; ORAL

ERY-TAB

AB		ARBOR PHARMS LLC	250MG	A062298	001	
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AB			333MG	A062298	003	Mar 29, 1982
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AB	!		500MG	A062298	002	
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ERYTHROMYCIN

AB		AMNEAL PHARMS CO	250MG	A210954	001	Jul 02, 2019
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AB			333MG	A210954	002	Jul 02, 2019
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AB			500MG	A210954	003	Jul 02, 2019
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AB		TORRENT	250MG	A211975	001	Jul 26, 2021
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AB			500MG	A211975	002	Jul 26, 2021
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ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

E.E.S.

AB	+	AZURITY	EQ 200MG BASE/5ML	N050207	001	
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ERYPED

AB	+	AZURITY	EQ 200MG BASE/5ML	N050207	003	Mar 30, 1987
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AB	+!		EQ 400MG BASE/5ML	N050207	002	
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ERYTHROMYCIN ETHYLSUCCINATE

AB		AMNEAL PHARMS	EQ 200MG BASE/5ML	A211204	001	Nov 01, 2019
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AB			EQ 400MG BASE/5ML	A211204	002	Nov 01, 2019
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AB		ANI PHARMS	EQ 200MG BASE/5ML	A062055	001	
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AB			EQ 200MG BASE/5ML	A062055	003	Nov 02, 2018
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AB			EQ 400MG BASE/5ML	A062055	002	Nov 02, 2018
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AB		CADILA PHARMS LTD	EQ 200MG BASE/5ML	A216212	001	Nov 21, 2022
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AB			EQ 400MG BASE/5ML	A216212	002	Nov 21, 2022
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TABLET; ORAL

E.E.S. 400

BX	!	ARBOR PHARMS LLC	EQ 400MG BASE	A061905	002	Aug 12, 1982
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ERYTHROMYCIN ETHYLSUCCINATE

BX	!	ARBOR PHARMS LLC	EQ 400MG BASE	A061904	001	
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PRESCRIPTION DRUG PRODUCT LIST

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062638 001</u>	Oct 31, 1986
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N050609 001</u>	Sep 24, 1986

ERYTHROMYCIN LACTOBIONATE

<u>AP</u>	NEXUS	<u>EQ 500MG BASE/VIAL</u>	<u>A215290 001</u>	Feb 14, 2022
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ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROCIN STEARATE

!	ARBOR PHARMS LLC	EQ 250MG BASE	A060359 001	
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ESCITALOPRAM OXALATE

SOLUTION; ORAL

ESCITALOPRAM OXALATE

<u>AA</u>	AMNEAL PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A202227 001</u>	Mar 14, 2012	
<u>AA</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE/5ML</u>	<u>A079062 001</u>	Apr 02, 2012	
<u>AA</u>	CHARTWELL MOLECULAR	<u>EQ 5MG BASE/5ML</u>	<u>A090477 001</u>	Jun 12, 2013	
<u>AA</u>	!	HETERO LABS LTD III	<u>EQ 5MG BASE/5ML</u>	<u>A202221 001</u>	Jun 12, 2012
<u>AA</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE/5ML</u>	<u>A202754 001</u>	Mar 31, 2016	
<u>AA</u>	TARO	<u>EQ 5MG BASE/5ML</u>	<u>A079121 001</u>	May 03, 2012	

TABLET; ORAL

ESCITALOPRAM OXALATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A202389 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202389 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A202389 003</u>	Sep 11, 2012
<u>AB</u>	AMNEAL PHARMS	<u>EQ 5MG BASE</u>	<u>A205619 001</u>	May 17, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205619 002</u>	May 17, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205619 003</u>	May 17, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A090432 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090432 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090432 003</u>	Sep 11, 2012
<u>AB</u>	CADILA	<u>EQ 5MG BASE</u>	<u>A077734 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077734 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077734 003</u>	Sep 11, 2012
<u>AB</u>	GRAVITI PHARMS	<u>EQ 5MG BASE</u>	<u>A078777 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078777 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078777 003</u>	Sep 11, 2012
<u>AB</u>	INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A078604 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078604 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078604 003</u>	Sep 11, 2012
<u>AB</u>	JUBILANT CADISTA	<u>EQ 5MG BASE</u>	<u>A202280 001</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202280 002</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A202280 003</u>	Sep 12, 2012
<u>AB</u>	LUPIN LTD	<u>EQ 5MG BASE</u>	<u>A078169 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078169 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078169 003</u>	Sep 11, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A202210 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202210 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A202210 003</u>	Sep 11, 2012
<u>AB</u>	PHARM ASSOC	<u>EQ 5MG BASE</u>	<u>A077512 001</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077512 002</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077512 003</u>	Sep 12, 2012
<u>AB</u>	PRINSTON INC	<u>EQ 5MG BASE</u>	<u>A078032 001</u>	Aug 28, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078032 002</u>	Aug 28, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078032 003</u>	Aug 28, 2015
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A090939 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090939 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090939 003</u>	Sep 11, 2012

LEXAPRO

<u>AB</u>	+	ABBVIE	<u>EQ 5MG BASE</u>	<u>N021323 001</u>	Aug 14, 2002
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N021323 002</u>	Aug 14, 2002
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N021323 003</u>	Aug 14, 2002

ESKETAMINE HYDROCHLORIDE

SPRAY; NASAL

SPRAVATO

!	JANSSEN PHARMS	EQ 28MG BASE	N211243 001	Mar 05, 2019
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PRESCRIPTION DRUG PRODUCT LIST

ESLICARBAZEPINE ACETATE

TABLET; ORAL

APTIOM

<u>AB</u>	+	SUMITOMO PHARMA AM	<u>200MG</u>	<u>N022416 001</u>	Nov 08, 2013
<u>AB</u>	+		<u>400MG</u>	<u>N022416 002</u>	Nov 08, 2013
<u>AB</u>	+		<u>600MG</u>	<u>N022416 003</u>	Nov 08, 2013
<u>AB</u>	+		<u>800MG</u>	<u>N022416 004</u>	Nov 08, 2013

ESLICARBAZEPINE ACETATE

<u>AB</u>		ALKEM LABS LTD	<u>200MG</u>	<u>A211199 001</u>	Oct 06, 2023
<u>AB</u>			<u>400MG</u>	<u>A211199 002</u>	Oct 06, 2023
<u>AB</u>			<u>600MG</u>	<u>A211199 003</u>	Oct 06, 2023
<u>AB</u>			<u>800MG</u>	<u>A211199 004</u>	Oct 06, 2023
<u>AB</u>		APOTEX	<u>200MG</u>	<u>A211236 001</u>	Dec 07, 2023
<u>AB</u>			<u>400MG</u>	<u>A211236 002</u>	Dec 07, 2023
<u>AB</u>			<u>600MG</u>	<u>A211236 003</u>	Dec 07, 2023
<u>AB</u>			<u>800MG</u>	<u>A211236 004</u>	Dec 07, 2023
<u>AB</u>		DR REDDYS	<u>200MG</u>	<u>A211238 001</u>	Jun 29, 2021
<u>AB</u>			<u>400MG</u>	<u>A211238 002</u>	Jun 29, 2021
<u>AB</u>			<u>600MG</u>	<u>A211238 003</u>	Jun 29, 2021
<u>AB</u>			<u>800MG</u>	<u>A211238 004</u>	Jun 29, 2021
<u>AB</u>		HETERO LABS LTD V	<u>200MG</u>	<u>A211186 001</u>	Aug 03, 2023
<u>AB</u>			<u>400MG</u>	<u>A211186 002</u>	Aug 03, 2023
<u>AB</u>			<u>600MG</u>	<u>A211186 003</u>	Aug 03, 2023
<u>AB</u>			<u>800MG</u>	<u>A211186 004</u>	Aug 03, 2023

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

<u>AP</u>	+	BAXTER HLTHCARE	<u>10MG/ML</u>	<u>N019386 006</u>	Feb 25, 2003
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BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>2GM/100ML</u>	<u>N019386 005</u>	Jan 27, 2003
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BREVIBLOC IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>1GM/100ML</u>	<u>N019386 004</u>	Feb 16, 2001
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ESMOLOL HYDROCHLORIDE

<u>AP</u>		AMNEAL	<u>1GM/100ML</u>	<u>A216603 001</u>	Dec 13, 2022
<u>AP</u>			<u>2GM/100ML</u>	<u>A216603 002</u>	Dec 13, 2022
<u>AP</u>		EUGIA PHARMA	<u>10MG/ML</u>	<u>A205520 001</u>	Jul 23, 2015
<u>AP</u>			<u>1GM/100ML</u>	<u>A216244 001</u>	Mar 21, 2022
<u>AP</u>			<u>2GM/100ML</u>	<u>A216244 002</u>	Mar 21, 2022
<u>AP</u>		GLAND PHARMA LTD	<u>10MG/ML</u>	<u>A208538 001</u>	Aug 14, 2019
<u>AP</u>		HIKMA	<u>10MG/ML</u>	<u>A076323 001</u>	Aug 10, 2004
<u>AP</u>		HQ SPCLT PHARMA	<u>1GM/100ML</u>	<u>A214172 001</u>	Dec 02, 2022
<u>AP</u>			<u>2GM/100ML</u>	<u>A214172 002</u>	Dec 02, 2022
<u>AP</u>		MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>A076474 001</u>	May 02, 2005
<u>AP</u>		MYLAN LABS LTD	<u>1GM/100ML</u>	<u>A206608 001</u>	Jun 08, 2018
<u>AP</u>			<u>2GM/100ML</u>	<u>A206608 002</u>	Jun 08, 2018
<u>AP</u>		SAGENT PHARMS INC	<u>1GM/100ML</u>	<u>A207107 001</u>	Jun 08, 2018
<u>AP</u>			<u>2GM/100ML</u>	<u>A207107 002</u>	Jun 08, 2018

SOLUTION; INTRAVENOUS

ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER

+	HQ SPCLT PHARMA	<u>2GM/100ML (20MG/ML)</u>	<u>N205703 002</u>	Apr 07, 2016
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ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER

+	HQ SPCLT PHARMA	<u>2.5GM/250ML (10MG/ML)</u>	<u>N205703 001</u>	Apr 07, 2016
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ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

ESOMEPRAZOLE MAGNESIUM

<u>AB</u>		ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A208333 001</u>	Oct 20, 2017
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A208333 002</u>	Oct 20, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A205606 001</u>	Apr 21, 2016
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A205606 002</u>	Apr 21, 2016
<u>AB</u>		CISEN	<u>EQ 20MG BASE</u>	<u>A213158 001</u>	Sep 22, 2020
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A213158 002</u>	Sep 22, 2020
<u>AB</u>		CSPC OUYI	<u>EQ 20MG BASE</u>	<u>A212949 001</u>	Oct 02, 2020
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A212949 002</u>	Oct 02, 2020
<u>AB</u>		DR REDDYS	<u>EQ 20MG BASE</u>	<u>A078279 001</u>	Sep 25, 2015
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A078279 002</u>	Sep 25, 2015
<u>AB</u>		ETHYPHARM	<u>EQ 20MG BASE</u>	<u>A090841 001</u>	Mar 31, 2021
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A090841 002</u>	Mar 31, 2021
<u>AB</u>		GLENMARK PHARMS	<u>EQ 20MG BASE</u>	<u>A209495 001</u>	May 10, 2019
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A209495 002</u>	May 10, 2019
<u>AB</u>		GRANULES	<u>EQ 20MG BASE</u>	<u>A217427 001</u>	Oct 18, 2023
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A217427 002</u>	Oct 18, 2023

PRESCRIPTION DRUG PRODUCT LIST

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS;ORAL

ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	GRAVITI PHARMS	<u>EQ 20MG BASE</u>	<u>A213486 001</u>	Mar 19, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213486 002</u>	Mar 19, 2021
<u>AB</u>	GUANGZHOU NOVAKEN	<u>EQ 20MG BASE</u>	<u>A213859 001</u>	Nov 18, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A213859 002</u>	Nov 18, 2020
<u>AB</u>	HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A211977 001</u>	Jun 02, 2020
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211977 002</u>	Jun 02, 2020
<u>AB</u>	INDCHEMIE HEALTH	<u>EQ 20MG BASE</u>	<u>A210559 001</u>	Feb 26, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A210559 002</u>	Feb 26, 2021
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078003 001</u>	Jan 26, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078003 002</u>	Jan 26, 2015
<u>AB</u>	LANNETT CO INC	<u>EQ 20MG BASE</u>	<u>A205563 001</u>	Sep 01, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205563 002</u>	Sep 01, 2017
<u>AB</u>	MYLAN	<u>EQ 20MG BASE</u>	<u>A078936 001</u>	Aug 02, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078936 002</u>	Aug 03, 2015
<u>AB</u>	PRINSTON INC	<u>EQ 20MG BASE</u>	<u>A214920 001</u>	Mar 28, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A214920 002</u>	Mar 28, 2023
<u>AB</u>	SUN PHARM	<u>EQ 20MG BASE</u>	<u>A209735 001</u>	Apr 30, 2018
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A209735 002</u>	Apr 30, 2018
<u>AB</u>	ZHEJIANG YONGTAI	<u>EQ 20MG BASE</u>	<u>A217022 001</u>	Dec 27, 2023
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A217022 002</u>	Dec 27, 2023
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE</u>	<u>A206296 001</u>	May 22, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206296 002</u>	May 22, 2019

NEXIUM

<u>AB</u>	+ ASTRAZENECA	<u>EQ 20MG BASE</u>	<u>N021153 001</u>	Feb 20, 2001
<u>AB</u>	+!	<u>EQ 40MG BASE</u>	<u>N021153 002</u>	Feb 20, 2001

FOR SUSPENSION, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	CIPLA	<u>EQ 10MG BASE/PACKET</u>	<u>A211752 001</u>	Mar 23, 2020
<u>AB</u>		<u>EQ 20MG BASE/PACKET</u>	<u>A211751 001</u>	Mar 23, 2020
<u>AB</u>		<u>EQ 40MG BASE/PACKET</u>	<u>A211751 002</u>	Mar 23, 2020
<u>AB</u>	ZYDUS PHARMS	<u>EQ 20MG BASE/PACKET</u>	<u>A206055 001</u>	Jun 07, 2023
<u>AB</u>		<u>EQ 40MG BASE/PACKET</u>	<u>A206055 002</u>	Jun 07, 2023

NEXIUM

<u>AB</u>	+ ASTRAZENECA	<u>EQ 10MG BASE/PACKET</u>	<u>N022101 001</u>	Feb 27, 2008
<u>AB</u>	+!	<u>EQ 20MG BASE/PACKET</u>	<u>N021957 001</u>	Oct 20, 2006
<u>AB</u>	+!	<u>EQ 40MG BASE/PACKET</u>	<u>N021957 002</u>	Oct 20, 2006
<u>AB</u>	+	EQ 2.5MG BASE/PACKET	N021957 003	Dec 15, 2011
<u>AB</u>	+	EQ 5MG BASE/PACKET	N021957 004	Dec 15, 2011

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE;ORAL

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 20MG BASE;375MG</u>	<u>A213699 001</u>	Oct 06, 2022
<u>AB</u>		<u>EQ 20MG BASE;500MG</u>	<u>A213699 002</u>	Oct 06, 2022
<u>AB</u>	DR REDDYS	<u>EQ 20MG BASE;375MG</u>	<u>A204206 001</u>	Feb 18, 2020
<u>AB</u>	!	<u>EQ 20MG BASE;500MG</u>	<u>A204206 002</u>	Feb 18, 2020
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 20MG BASE;375MG</u>	<u>A217738 001</u>	Oct 11, 2023
<u>AB</u>		<u>EQ 20MG BASE;500MG</u>	<u>A217738 002</u>	Oct 11, 2023

ESOMEPRAZOLE SODIUM

INJECTABLE;INTRAVENOUS

ESOMEPRAZOLE SODIUM

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 40MG BASE/VIAL</u>	<u>A205379 001</u>	Sep 25, 2015
<u>AP</u>	DEVA HOLDING AS	<u>EQ 40MG BASE/VIAL</u>	<u>A207181 001</u>	Mar 06, 2017
<u>AP</u>	EUGIA PHARMA	<u>EQ 40MG BASE/VIAL</u>	<u>A204657 002</u>	Aug 10, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A203349 002</u>	Apr 01, 2020
<u>AP</u>	HAINAN POLY	<u>EQ 40MG BASE/VIAL</u>	<u>A215732 001</u>	Feb 10, 2022
<u>AP</u>	SUN PHARM	<u>EQ 40MG BASE/VIAL</u>	<u>A200882 002</u>	Mar 18, 2013

NEXIUM IV

<u>AP</u>	+! ASTRAZENECA	<u>EQ 40MG BASE/VIAL</u>	<u>N021689 002</u>	Mar 31, 2005
	ESOMEPRAZOLE SODIUM			
	GLAND PHARMA LTD	EQ 20MG BASE/VIAL	A203349 001	Apr 01, 2020

ESTAZOLAM

TABLET;ORAL

ESTAZOLAM

<u>AB</u>	DR REDDYS LABS SA	<u>1MG</u>	<u>A074921 001</u>	Jul 10, 1997
<u>AB</u>	!	<u>2MG</u>	<u>A074921 002</u>	Jul 10, 1997
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A074826 001</u>	Jul 03, 1997
<u>AB</u>		<u>2MG</u>	<u>A074826 002</u>	Jul 03, 1997

PRESCRIPTION DRUG PRODUCT LIST

ESTAZOLAM

TABLET;ORAL

ESTAZOLAM

<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>A074818</u>	<u>001</u>	Aug 19, 1997
<u>AB</u>		<u>2MG</u>	<u>A074818</u>	<u>002</u>	Aug 19, 1997

ESTRADIOL

CREAM;VAGINAL

ESTRACE

<u>AB</u>	+!	ALLERGAN	<u>0.01%</u>	<u>A086069</u>	<u>001</u>	Jan 31, 1984
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ESTRADIOL

<u>AB</u>		ALVOGEN	<u>0.01%</u>	<u>A209767</u>	<u>001</u>	Mar 05, 2018
<u>AB</u>		MYLAN	<u>0.01%</u>	<u>A208788</u>	<u>001</u>	Dec 29, 2017
<u>AB</u>		PADAGIS ISRAEL	<u>0.01%</u>	<u>A210194</u>	<u>001</u>	Jan 22, 2018
<u>AB</u>		PRASCO	<u>0.01%</u>	<u>A212313</u>	<u>001</u>	Jul 15, 2021
<u>AB</u>		TEVA PHARMS USA	<u>0.01%</u>	<u>A210488</u>	<u>001</u>	Mar 30, 2018

FILM, EXTENDED RELEASE;TRANSDERMAL

CLIMARA

<u>AB</u>	+	BAYER HLTHCARE	<u>0.06MG/24HR</u>	<u>N020375</u>	<u>006</u>	May 27, 2003
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ESTRADIOL

<u>AB</u>		MYLAN TECHNOLOGIES	<u>0.06MG/24HR</u>	<u>A075182</u>	<u>005</u>	Jul 20, 2006
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CLIMARA

<u>AB2</u>	+	BAYER HLTHCARE	<u>0.025MG/24HR</u>	<u>N020375</u>	<u>004</u>	Mar 05, 1999
<u>AB2</u>	+		<u>0.0375MG/24HR</u>	<u>N020375</u>	<u>005</u>	May 27, 2003
<u>AB2</u>	+		<u>0.05MG/24HR</u>	<u>N020375</u>	<u>001</u>	Dec 22, 1994
<u>AB2</u>	+		<u>0.075MG/24HR</u>	<u>N020375</u>	<u>003</u>	Mar 23, 1998
<u>AB2</u>	+!		<u>0.1MG/24HR</u>	<u>N020375</u>	<u>002</u>	Dec 22, 1994

ESTRADIOL

<u>AB2</u>		MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A075182</u>	<u>003</u>	Jan 26, 2005
<u>AB2</u>			<u>0.0375MG/24HR</u>	<u>A075182</u>	<u>004</u>	Jul 20, 2006
<u>AB2</u>			<u>0.05MG/24HR</u>	<u>A075182</u>	<u>006</u>	Feb 24, 2000
<u>AB2</u>			<u>0.075MG/24HR</u>	<u>A075182</u>	<u>002</u>	Jan 26, 2005
<u>AB2</u>			<u>0.1MG/24HR</u>	<u>A075182</u>	<u>001</u>	Feb 24, 2000
<u>AB3</u>		AMNEAL	<u>0.025MG/24HR</u>	<u>A211396</u>	<u>001</u>	Sep 28, 2020
<u>AB3</u>			<u>0.0375MG/24HR</u>	<u>A211396</u>	<u>002</u>	Sep 28, 2020
<u>AB3</u>			<u>0.05MG/24HR</u>	<u>A211396</u>	<u>003</u>	Sep 28, 2020
<u>AB3</u>			<u>0.075MG/24HR</u>	<u>A211396</u>	<u>004</u>	Sep 28, 2020
<u>AB3</u>			<u>0.1MG/24HR</u>	<u>A211396</u>	<u>005</u>	Sep 28, 2020
<u>AB3</u>		MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A206685</u>	<u>001</u>	Aug 15, 2018
<u>AB3</u>			<u>0.0375MG/24HR</u>	<u>A206685</u>	<u>002</u>	Aug 15, 2018
<u>AB3</u>			<u>0.05MG/24HR</u>	<u>A206685</u>	<u>003</u>	Aug 15, 2018
<u>AB3</u>			<u>0.075MG/24HR</u>	<u>A206685</u>	<u>004</u>	Aug 15, 2018
<u>AB3</u>			<u>0.1MG/24HR</u>	<u>A206685</u>	<u>005</u>	Aug 15, 2018

MINIVELLE

<u>AB3</u>	+	NOVEN	<u>0.025MG/24HR</u>	<u>N203752</u>	<u>005</u>	Sep 23, 2014
<u>AB3</u>	+		<u>0.0375MG/24HR</u>	<u>N203752</u>	<u>001</u>	Oct 29, 2012
<u>AB3</u>	+		<u>0.05MG/24HR</u>	<u>N203752</u>	<u>003</u>	Oct 29, 2012
<u>AB3</u>	+		<u>0.075MG/24HR</u>	<u>N203752</u>	<u>002</u>	Oct 29, 2012
<u>AB3</u>	+!		<u>0.1MG/24HR</u>	<u>N203752</u>	<u>004</u>	Oct 29, 2012

GEL;TRANSDERMAL

DIVIGEL

<u>AB</u>	+!	VERTICAL PHARMS	<u>0.1%</u>	<u>N022038</u>	<u>001</u>	Jun 04, 2007
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ESTRADIOL

<u>AB</u>		CHEMO RESEARCH SL	<u>0.1%</u>	<u>A211783</u>	<u>001</u>	Aug 10, 2022
<u>AB</u>		NOVITIUM PHARMA	<u>0.1%</u>	<u>A217610</u>	<u>001</u>	Aug 24, 2023
<u>AB</u>		PADAGIS ISRAEL	<u>0.1%</u>	<u>A216524</u>	<u>001</u>	Nov 14, 2023

GEL, METERED;TRANSDERMAL

ELESTRIN

	+!	MYLAN SPECIALITY LP	0.06% (0.87GM/ACTIVATION)	N021813	001	Dec 15, 2006
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ESTROGEL

	+!	ASCEND THERAPS US	0.06% (1.25GM/ACTIVATION)	N021166	002	Feb 09, 2004
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INSERT;VAGINAL

IMVEXXY

	+	MAYNE PHARMA	0.004MG	N208564	001	May 29, 2018
	+!		0.01MG	N208564	002	May 29, 2018

INSERT, EXTENDED RELEASE;VAGINAL

ESTRING

	+!	PFIZER	0.0075MG/24HR	N020472	001	Apr 26, 1996
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SPRAY;TRANSDERMAL

EVAMIST

	+!	PADAGIS US	1.53MG/SPRAY	N022014	001	Jul 27, 2007
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PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL

SYSTEM; TRANSDERMAL

ESTRADIOL

<u>AB</u>	ZYDUS PHARMS	<u>0.014MG/24HR</u>	<u>A204379 001</u>	Apr 17, 2023
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MENOSTAR

<u>AB</u>	+! BAYER HLTHCARE	<u>0.014MG/24HR</u>	<u>N021674 001</u>	Jun 08, 2004
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ESTRADIOL

<u>AB1</u>	AMNEAL	<u>0.025MG/24HR</u>	<u>A211293 001</u>	Feb 04, 2019
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<u>AB1</u>		<u>0.0375MG/24HR</u>	<u>A211293 002</u>	Feb 04, 2019
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<u>AB1</u>		<u>0.05MG/24HR</u>	<u>A211293 003</u>	Feb 04, 2019
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<u>AB1</u>		<u>0.075MG/24HR</u>	<u>A211293 004</u>	Feb 04, 2019
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<u>AB1</u>		<u>0.1MG/24HR</u>	<u>A211293 005</u>	Feb 04, 2019
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<u>AB1</u>	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>A201675 001</u>	Dec 19, 2014
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<u>AB1</u>		<u>0.0375MG/24HR</u>	<u>A201675 002</u>	Dec 19, 2014
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<u>AB1</u>		<u>0.05MG/24HR</u>	<u>A201675 003</u>	Dec 19, 2014
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<u>AB1</u>		<u>0.075MG/24HR</u>	<u>A201675 004</u>	Dec 19, 2014
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<u>AB1</u>		<u>0.1MG/24HR</u>	<u>A201675 005</u>	Dec 19, 2014
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<u>AB1</u>	ZYDUS PHARMS	<u>0.025MG/24HR</u>	<u>A206241 001</u>	Dec 01, 2022
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<u>AB1</u>		<u>0.0375MG/24HR</u>	<u>A206241 002</u>	Dec 01, 2022
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<u>AB1</u>		<u>0.05MG/24HR</u>	<u>A206241 003</u>	Dec 01, 2022
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<u>AB1</u>		<u>0.075MG/24HR</u>	<u>A206241 004</u>	Dec 01, 2022
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<u>AB1</u>		<u>0.1MG/24HR</u>	<u>A206241 005</u>	Dec 01, 2022
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VIVELLE-DOT

<u>AB1</u>	+ SANDOZ	<u>0.025MG/24HR</u>	<u>N020538 009</u>	May 03, 2002
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<u>AB1</u>	+	<u>0.0375MG/24HR</u>	<u>N020538 005</u>	Jan 08, 1999
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<u>AB1</u>	+	<u>0.05MG/24HR</u>	<u>N020538 006</u>	Jan 08, 1999
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<u>AB1</u>	+	<u>0.075MG/24HR</u>	<u>N020538 007</u>	Jan 08, 1999
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<u>AB1</u>	+!	<u>0.1MG/24HR</u>	<u>N020538 008</u>	Jan 08, 1999
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TABLET; ORAL

ESTRADIOL

<u>AB</u>	BARR LABS INC	<u>0.5MG</u>	<u>A040197 001</u>	Oct 22, 1997
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<u>AB</u>		<u>1MG</u>	<u>A040197 002</u>	Oct 22, 1997
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<u>AB</u>	!	<u>2MG</u>	<u>A040197 003</u>	Oct 22, 1997
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<u>AB</u>	EPIC PHARMA LLC	<u>0.5MG</u>	<u>A040275 001</u>	Dec 29, 1998
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<u>AB</u>		<u>1MG</u>	<u>A040275 002</u>	Dec 29, 1998
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<u>AB</u>		<u>2MG</u>	<u>A040275 003</u>	Dec 29, 1998
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<u>AB</u>	NOVITIUM PHARMA	<u>0.5MG</u>	<u>A217334 001</u>	Sep 06, 2023
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<u>AB</u>		<u>1MG</u>	<u>A217334 002</u>	Sep 06, 2023
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<u>AB</u>		<u>2MG</u>	<u>A217334 003</u>	Sep 06, 2023
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TABLET; VAGINAL

ESTRADIOL

<u>AB</u>	AMNEAL PHARMS	<u>10MCG</u>	<u>A205256 001</u>	May 29, 2015
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<u>AB</u>	GLENMARK PHARMS LTD	<u>10MCG</u>	<u>A210264 001</u>	Sep 14, 2018
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<u>AB</u>	TEVA PHARMS USA	<u>10MCG</u>	<u>A206388 001</u>	Jul 21, 2017
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VAGIFEM

<u>AB</u>	+! NOVO NORDISK INC	<u>10MCG</u>	<u>N020908 002</u>	Nov 25, 2009
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ESTRADIOL ACETATE

INSERT, EXTENDED RELEASE; VAGINAL

FEMRING

+	MILLICENT	EQ 0.05MG BASE/24HR	N021367 001	Mar 20, 2003
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+	!	EQ 0.1MG BASE/24HR	N021367 002	Mar 20, 2003
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ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

!	PFIZER	5MG/ML	A085470 003	
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ESTRADIOL VALERATE

INJECTABLE; INJECTION

DELESTROGEN

<u>AO</u>	+!	PAR STERILE PRODUCTS	<u>10MG/ML</u>	<u>N009402 002</u>
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<u>AO</u>	+		<u>20MG/ML</u>	<u>N009402 004</u>
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<u>AO</u>	+		<u>40MG/ML</u>	<u>N009402 003</u>
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ESTRADIOL VALERATE

<u>AO</u>		AM REGENT	<u>20MG/ML</u>	<u>A090920 001</u>	Jan 19, 2010
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<u>AO</u>			<u>40MG/ML</u>	<u>A090920 002</u>	Jan 19, 2010
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<u>AO</u>		HIKMA	<u>10MG/ML</u>	<u>A203723 001</u>	Apr 21, 2020
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<u>AO</u>			<u>20MG/ML</u>	<u>A203723 002</u>	Apr 21, 2020
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<u>AO</u>			<u>40MG/ML</u>	<u>A203723 003</u>	Apr 21, 2020
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<u>AO</u>		XIROMED	<u>10MG/ML</u>	<u>A216656 001</u>	Apr 28, 2023
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<u>AO</u>			<u>20MG/ML</u>	<u>A216656 002</u>	Apr 28, 2023
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<u>AO</u>			<u>40MG/ML</u>	<u>A216656 003</u>	Apr 28, 2023
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PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL; LEVONORGESTREL

FILM, EXTENDED RELEASE;TRANSDERMAL

CLIMARA PRO

+! BAYER HLTHCARE 0.045MG/24HR;0.015MG/24HR N021258 001 Nov 21, 2003

ESTRADIOL; NORETHINDRONE ACETATE

FILM, EXTENDED RELEASE;TRANSDERMAL

COMBIPATCH

+ NOVEN PHARMS INC 0.05MG/24HR;0.14MG/24HR N020870 001 Aug 07, 1998

+! 0.05MG/24HR;0.25MG/24HR N020870 002 Aug 07, 1998

TABLET;ORAL

ACTIVELLA**AB** +! AMNEAL **1MG;0.5MG** **N020907 001** Nov 18, 1998ESTRADIOL AND NORETHINDRONE ACETATE**AB** BARR **1MG;0.5MG** **A079193 001** May 11, 2010**AB** BRECKENRIDGE PHARM **0.5MG;0.1MG** **A078324 002** Jun 09, 2011**AB** **1MG;0.5MG** **A078324 001** Apr 17, 2008**AB** MYLAN LABS LTD **0.5MG;0.1MG** **A207261 001** Feb 10, 2017**AB** **1MG;0.5MG** **A207261 002** Feb 10, 2017**AB** NOVAST LABS **0.5MG;0.1MG** **A210612 001** Apr 03, 2019**AB** **1MG;0.5MG** **A210612 002** Apr 03, 2019

AMABELZ

BX LUPIN LTD 0.5MG;0.1MG A203339 001 Jun 20, 2016

BX 1MG;0.5MG A203339 002 Jun 20, 2016

ESTRADIOL; NORETHINDRONE ACETATE; RELUGOLIX

TABLET;ORAL

MYFEMBREE

+! MYOVANT SCIENCES 1MG;0.5MG;40MG N214846 001 May 26, 2021

ESTRADIOL; NORGESTIMATE

TABLET;ORAL

ESTRADIOL AND NORGESTIMATE

! BARR 1MG,1MG;N/A,0.09MG A076812 001 Apr 29, 2005

ESTRADIOL; PROGESTERONE

CAPSULE;ORAL

BIJUVA

+ MAYNE PHARMA 0.5MG;100MG N210132 002 Dec 28, 2021

+! 1MG;100MG N210132 001 Oct 28, 2018

ESTRAMUSTINE PHOSPHATE SODIUM

CAPSULE;ORAL

EMCYT

+! PHARMACIA AND UPJOHN EQ 140MG PHOSPHATE N018045 001

ESTROGENS, CONJUGATED

CREAM;TOPICAL, VAGINAL

PREMARIN

+! WYETH PHARMS 0.625MG/GM N020216 001

INJECTABLE;INJECTION

PREMARIN

+! WYETH PHARMS 25MG/VIAL N010402 001

TABLET;ORAL

PREMARIN

+ WYETH PHARMS 0.3MG N004782 003

+ 0.45MG N004782 006 Jul 16, 2003

+! 0.625MG N004782 004

+! 0.9MG N004782 005 Jan 26, 1984

+! 1.25MG N004782 001

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET;ORAL-28

PREMPHASE 14/14

+! WYETH PHARMS 0.625MG,0.625MG;N/A,5MG N020527 002 Nov 17, 1995

PREMPRO

+! WYETH PHARMS 0.3MG;1.5MG N020527 005 Jun 04, 2003

+! 0.45MG;1.5MG N020527 004 Mar 12, 2003

+! 0.625MG;2.5MG N020527 001 Nov 17, 1995

+! 0.625MG;5MG N020527 003 Jan 09, 1998

PRESCRIPTION DRUG PRODUCT LIST

ESTROGENS, ESTERIFIED

TABLET; ORAL

MENEST

MONARCH PHARMS	0.3MG	A084951	001
	0.625MG	A084948	001
	1.25MG	A084950	001
!	2.5MG	A084949	001

ESTROPIPATE

TABLET; ORAL

OGEN 5

+ PFIZER	6MG	A083220	004
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ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

<u>AB</u>	AUROBINDO PHARMA	<u>1MG</u>	<u>A208451</u>	<u>001</u>	Sep 15, 2016
<u>AB</u>		<u>2MG</u>	<u>A208451</u>	<u>002</u>	Sep 15, 2016
<u>AB</u>		<u>3MG</u>	<u>A208451</u>	<u>003</u>	Sep 15, 2016
<u>AB</u>	DR REDDYS	<u>1MG</u>	<u>A091024</u>	<u>001</u>	Apr 15, 2014
<u>AB</u>		<u>2MG</u>	<u>A091024</u>	<u>002</u>	Apr 15, 2014
<u>AB</u>		<u>3MG</u>	<u>A091024</u>	<u>003</u>	Apr 15, 2014
<u>AB</u>	GLENMARK GENERICS	<u>1MG</u>	<u>A091166</u>	<u>001</u>	Apr 15, 2014
<u>AB</u>		<u>2MG</u>	<u>A091166</u>	<u>002</u>	Apr 15, 2014
<u>AB</u>		<u>3MG</u>	<u>A091166</u>	<u>003</u>	Apr 15, 2014
<u>AB</u>	HETERO LABS LTD V	<u>1MG</u>	<u>A205504</u>	<u>001</u>	Jan 04, 2024
<u>AB</u>		<u>2MG</u>	<u>A205504</u>	<u>002</u>	Jan 04, 2024
<u>AB</u>		<u>3MG</u>	<u>A205504</u>	<u>003</u>	Jan 04, 2024
<u>AB</u>	IPCA LABS LTD	<u>1MG</u>	<u>A206222</u>	<u>001</u>	Dec 29, 2023
<u>AB</u>		<u>2MG</u>	<u>A206222</u>	<u>002</u>	Dec 29, 2023
<u>AB</u>		<u>3MG</u>	<u>A206222</u>	<u>003</u>	Dec 29, 2023
<u>AB</u>	LUPIN LTD	<u>1MG</u>	<u>A091124</u>	<u>001</u>	Sep 13, 2011
<u>AB</u>		<u>2MG</u>	<u>A091124</u>	<u>002</u>	Sep 13, 2011
<u>AB</u>		<u>3MG</u>	<u>A091124</u>	<u>003</u>	Sep 13, 2011
<u>AB</u>	MACLEODS PHARMS LTD	<u>1MG</u>	<u>A202929</u>	<u>001</u>	Jan 30, 2015
<u>AB</u>		<u>2MG</u>	<u>A202929</u>	<u>002</u>	Jan 30, 2015
<u>AB</u>		<u>3MG</u>	<u>A202929</u>	<u>003</u>	Jan 30, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>1MG</u>	<u>A091151</u>	<u>001</u>	Mar 26, 2013
<u>AB</u>		<u>2MG</u>	<u>A091151</u>	<u>002</u>	Mar 26, 2013
<u>AB</u>		<u>3MG</u>	<u>A091151</u>	<u>003</u>	Mar 26, 2013
<u>AB</u>	ORBION PHARMS	<u>1MG</u>	<u>A091113</u>	<u>001</u>	Jun 10, 2014
<u>AB</u>		<u>2MG</u>	<u>A091113</u>	<u>002</u>	Jun 10, 2014
<u>AB</u>		<u>3MG</u>	<u>A091113</u>	<u>003</u>	Jun 10, 2014
<u>AB</u>	SUN PHARM	<u>1MG</u>	<u>A091103</u>	<u>001</u>	Apr 03, 2013
<u>AB</u>		<u>2MG</u>	<u>A091103</u>	<u>002</u>	Apr 03, 2013
<u>AB</u>		<u>3MG</u>	<u>A091103</u>	<u>003</u>	Apr 03, 2013
<u>AB</u>	TEVA	<u>1MG</u>	<u>A091169</u>	<u>001</u>	May 23, 2011
<u>AB</u>		<u>2MG</u>	<u>A091169</u>	<u>002</u>	May 23, 2011
<u>AB</u>		<u>3MG</u>	<u>A091169</u>	<u>003</u>	May 23, 2011
	<u>LUNESTA</u>				
<u>AB</u>	+ WOODWARD	<u>1MG</u>	<u>N021476</u>	<u>001</u>	Dec 15, 2004
<u>AB</u>	+	<u>2MG</u>	<u>N021476</u>	<u>002</u>	Dec 15, 2004
<u>AB</u>	+!	<u>3MG</u>	<u>N021476</u>	<u>003</u>	Dec 15, 2004

EETELCALCETIDE

SOLUTION; INTRAVENOUS

PARSABIV

+! KAI PHARMS INC	2.5MG/0.5ML (2.5MG/0.5ML)	N208325	001	Feb 07, 2017
+!	5MG/ML (5MG/ML)	N208325	002	Feb 07, 2017
+!	10MG/2ML (5MG/ML)	N208325	003	Feb 07, 2017

EETPLIRSEN

SOLUTION; INTRAVENOUS

EXONDYS 51

+! SAREPTA THERAPS INC	100MG/2ML (50MG/ML)	N206488	001	Sep 19, 2016
+!	500MG/10ML (50MG/ML)	N206488	002	Sep 19, 2016

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

EDECIN

<u>AP</u>	+! BAUSCH	<u>EQ 50MG BASE/VIAL</u>	<u>N016093</u>	<u>001</u>
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ETHACRYNATE SODIUM

<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 50MG BASE/VIAL</u>	<u>A204634</u>	<u>001</u>	Aug 23, 2016
<u>AP</u>	PAR STERILE PRODUCTS	<u>EQ 50MG BASE/VIAL</u>	<u>A205473</u>	<u>001</u>	Jul 29, 2015

PRESCRIPTION DRUG PRODUCT LIST

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

ETHACRYNATE SODIUM

AP	VPI PHARMS INC	EQ 50MG BASE/VIAL	A208663 001	Jun 09, 2020
AP	ZYDUS PHARMS	EQ 50MG BASE/VIAL	A207758 001	Nov 17, 2017

ETHACRYNIC ACID

TABLET; ORAL

EDECRIN

AB	+ ! BAUSCH	25MG	N016092 001	
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ETHACRYNIC ACID

AB	AGNITIO	25MG	A211809 001	Jul 12, 2019
AB	AMNEAL PHARMS CO	25MG	A208805 001	May 08, 2018
AB	CHARTWELL RX	25MG	A213240 001	Oct 19, 2020
AB	EDENBRIDGE PHARMS	25MG	A205609 001	Jun 30, 2016
AB	LUPIN LTD	25MG	A211719 001	Sep 06, 2019
AB	PAR PHARM INC	25MG	A208501 001	Jul 21, 2017
AB	SCIEGEN PHARMS INC	25MG	A211232 001	Aug 27, 2019
AB	UPSHER SMITH LABS	25MG	A212417 001	Feb 19, 2020

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

ETHAMBUTOL HYDROCHLORIDE

AB	EPIC PHARMA LLC	100MG	A075095 001	Nov 30, 1999
AB		400MG	A075095 002	Nov 30, 1999
AB	LUPIN	100MG	A078939 001	Jun 17, 2009
AB		400MG	A078939 002	Jun 17, 2009

MYAMBUTOL

AB	+ STI PHARMA LLC	100MG	N016320 001	
AB	+ !	400MG	N016320 003	

ETHANOLAMINE OLEATE

INJECTABLE; INJECTION

ETHAMOLIN

+ !	QOL MEDCL	50MG/ML	N019357 001	Dec 22, 1988
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ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-28

ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL

AB	MYLAN LABS LTD	0.035MG;1MG	A204703 001	Jul 28, 2016
AB		0.05MG;1MG	A204704 001	Feb 09, 2016
	KELNOR			
AB	BARR	0.035MG;1MG	A076785 001	May 23, 2005
	LO-MALMOREDE			
AB	NOVAST LABS	0.035MG;1MG	A209548 001	Feb 11, 2019
	MALMOREDE			
AB	NOVAST LABS	0.05MG;1MG	A209547 001	Jul 25, 2018
	ZOVIA 1/50E-28			
AB	! WATSON LABS	0.05MG;1MG	A072723 001	Dec 30, 1991

ETHINYL ESTRADIOL; ETONOGESTREL

RING; VAGINAL

ELURYNG

AB	AMNEAL	0.015MG/24HR;0.12MG/24HR	A210830 001	Dec 11, 2019
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ENILLORING

AB	XIROMED	0.015MG/24HR;0.12MG/24HR	A211157 001	Jun 29, 2023
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ETHINYL ESTRADIOL; ETONOGESTREL

AB	TEVA PHARMS USA INC	0.015MG/24HR;0.12MG/24HR	A204305 001	Jan 13, 2021
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HALOETTE

AB	DR REDDYS LABS SA	0.015MG/24HR;0.12MG/24HR	A211328 001	Aug 05, 2022
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NUVARING

AB	+ ! ORGANON USA ORGANON	0.015MG/24HR;0.12MG/24HR	N021187 001	Oct 03, 2001
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ETHINYL ESTRADIOL; LEVONORGESTREL

SYSTEM; TRANSDERMAL

TWIRLA

+ !	AGILE	0.03MG/24HR;0.12MG/24HR	N204017 001	Feb 14, 2020
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TABLET; ORAL

ASHLYNA

AB	GLENMARK GENERICS	0.03MG,0.01MG;0.15MG,N/A	A203163 001	Feb 23, 2015
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DAYSEE

AB	LUPIN LTD	0.03MG,0.01MG;0.15MG,N/A	A091467 001	Apr 10, 2013
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DOLISHALE

AB	NOVAST LABS	0.02MG;0.09MG	A091692 001	Oct 22, 2020
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PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

<u>ICLEVIA</u>					
AB	AUROBINDO PHARMA LTD	0.03MG;0.15MG	A206850	001	Jun 29, 2018
<u>INTROVALE</u>					
AB	XIROMED	0.03MG;0.15MG	A079064	001	Sep 27, 2010
<u>JAIMIESS</u>					
AB	XIROMED	0.03MG,0.01MG;0.15MG,N/A	A203770	001	Dec 27, 2017
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB	AMNEAL PHARMS	0.03MG;0.15MG	A203871	001	Nov 13, 2015
AB		0.03MG,0.01MG;0.15MG,N/A	A203872	001	Dec 22, 2015
AB	GLENMARK GENERICS	0.02MG;0.09MG	A202791	001	Apr 09, 2015
AB	GLENMARK PHARMS LTD	0.03MG;0.15MG	A203164	001	Jun 12, 2015
AB	LUPIN LTD	0.03MG;0.15MG	A091440	001	Oct 23, 2012
AB	MYLAN LABS LTD	0.03MG;0.15MG	A200490	001	Apr 21, 2015
AB	! WATSON LABS	0.02MG;0.09MG	A079218	001	Jun 06, 2011
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL</u>					
AB	DR REDDYS LABS SA	0.03MG,0.01MG;0.15MG,N/A	A078834	001	May 31, 2011
AB	LUPIN LTD	0.02MG,0.1MG;0.01MG,N/A	A091674	001	Oct 26, 2011
AB	MYLAN LABS LTD	0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A	A206053	001	Oct 02, 2017
AB		0.02MG,0.1MG;0.01MG,N/A	A200493	001	Jun 17, 2015
AB		0.03MG,0.01MG;0.15MG,N/A	A200492	001	May 27, 2015
AB	XIROMED	0.02MG,0.1MG;0.01MG,N/A	A205131	001	Dec 14, 2017
<u>LO SIMPESE</u>					
AB	AUROBINDO PHARMA	0.02MG,0.1MG;0.01MG,N/A	A206852	001	Apr 28, 2017
<u>LOSEASONIQUE</u>					
AB	TEVA BRANDED PHARM	0.02MG,0.1MG;0.01MG,N/A	N022262	001	Oct 24, 2008
<u>QUARTETTE</u>					
AB	+! TEVA BRANDED PHARM	0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A	N204061	001	Mar 28, 2013
<u>QUASENSE</u>					
AB	WATSON LABS	0.03MG;0.15MG	A077101	001	Sep 06, 2006
<u>SEASONALE</u>					
AB	+! TEVA BRANDED PHARM	0.03MG;0.15MG	N021544	001	Sep 05, 2003
<u>SEASONIQUE</u>					
AB	+! TEVA BRANDED PHARM	0.03MG,0.01MG;0.15MG,N/A	N021840	001	May 25, 2006
<u>SETLAKIN</u>					
AB	NOVAST LABS	0.03MG;0.15MG	A090716	001	Sep 15, 2014
<u>SIMPESE</u>					
AB	AUROBINDO PHARMA	0.03MG,0.01MG;0.15MG,N/A	A206851	001	Apr 07, 2017
<u>BALCOLTRA</u>					
AB3	+! AVION PHARMS	0.02MG;0.1MG	N208612	001	Jan 09, 2018
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>					
AB3	XIROMED	0.02MG;0.1MG	A214640	001	Aug 16, 2023
TYBLUME					
	+! EXELTIS USA INC	0.02MG;0.1MG	N209405	001	Mar 30, 2020
TABLET; ORAL-28					
<u>ALTAVERA</u>					
AB	XIROMED	0.03MG;0.15MG	A079102	001	Aug 03, 2010
<u>AYUNA</u>					
AB	AUROBINDO PHARMA	0.03MG;0.15MG	A206866	001	Sep 23, 2016
<u>ENPRESSE-28</u>					
AB	DURAMED PHARMS BARR	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A075809	002	Jul 16, 2001
<u>KURVELO</u>					
AB	LUPIN LTD	0.03MG;0.15MG	A091408	001	Oct 17, 2012
<u>LEVONEST</u>					
AB	NOVAST LABS LTD	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A090719	001	Dec 29, 2010
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
AB	LUPIN LTD	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A200248	001	Nov 19, 2015
AB	MYLAN LABS LTD	0.03MG;0.15MG	A091663	001	Dec 21, 2012
AB	NAARI PTE LTD	0.03MG;0.15MG	A207033	001	Oct 09, 2020
<u>LEVORA 0.15/30-28</u>					
AB	! DR REDDYS LABS SA	0.03MG;0.15MG	A073594	001	Dec 13, 1993
<u>MARLISSA</u>					
AB	GLENMARK GENERICS	0.03MG;0.15MG	A091452	001	Feb 29, 2012
<u>MYZILRA</u>					
AB	VINTAGE PHARMS LLC	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	A077502	001	Nov 23, 2011

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-28

PORTIA-28

AB	BARR	<u>0.03MG;0.15MG</u>	<u>A075866</u>	<u>002</u>	May 23, 2002
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TRIVORA-28

AB	!	DR REDDYS LABS SA	<u>0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG</u>	<u>A074538</u>	<u>002</u>	Dec 18, 1997
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AFIRMELLE

AB1	AUROBINDO PHARMA	<u>0.02MG;0.1MG</u>	<u>A206886</u>	<u>001</u>	Nov 14, 2016
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AVIANE-28

AB1	DURAMED PHARMS BARR	<u>0.02MG;0.1MG</u>	<u>A075796</u>	<u>001</u>	Apr 30, 2001
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FALMINA

AB1	NOVAST LABS LTD	<u>0.02MG;0.1MG</u>	<u>A090721</u>	<u>001</u>	Mar 28, 2012
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LEVONORGESTREL AND ETHINYL ESTRADIOL

AB1	!	DR REDDYS LABS SA	<u>0.02MG;0.1MG</u>	<u>A076625</u>	<u>001</u>	Nov 18, 2004
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AB1	HETERO LABS	<u>0.02MG;0.1MG</u>	<u>A212298</u>	<u>001</u>	Feb 13, 2023
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AB1	LUPIN LTD	<u>0.02MG;0.1MG</u>	<u>A091425</u>	<u>001</u>	Jan 18, 2013
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AB1	MYLAN LABS LTD	<u>0.02MG;0.1MG</u>	<u>A200245</u>	<u>001</u>	Oct 09, 2013
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AB1	NAARI PTE LTD	<u>0.02MG;0.1MG</u>	<u>A207065</u>	<u>001</u>	Aug 17, 2020
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VIENVA

AB1	XIROMED	<u>0.02MG;0.1MG</u>	<u>A201088</u>	<u>001</u>	May 21, 2015
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LESSINA-28

AB2	BARR	<u>0.02MG;0.1MG</u>	<u>A075803</u>	<u>002</u>	Mar 20, 2002
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LEVONORGESTREL AND ETHINYL ESTRADIOL

AB2	!	DR REDDYS LABS SA	<u>0.02MG;0.1MG</u>	<u>A077681</u>	<u>001</u>	May 31, 2006
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ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

ETHINYL ESTRADIOL AND NORELGESTROMIN

AB	AMNEAL	<u>0.035MG/24HR;0.15MG/24HR</u>	<u>A213950</u>	<u>001</u>	Feb 25, 2021
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AB	ZYDUS NOVELTECH INC	<u>0.035MG/24HR;0.15MG/24HR</u>	<u>A214594</u>	<u>001</u>	Sep 14, 2023
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ONSURA

AB	TEVA PHARMS USA	<u>0.035MG/24HR;0.15MG/24HR</u>	<u>A213977</u>	<u>001</u>	Aug 25, 2021
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XULANE

AB	!	MYLAN TECHNOLOGIES	<u>0.035MG/24HR;0.15MG/24HR</u>	<u>A200910</u>	<u>001</u>	Apr 16, 2014
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ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL

NEXESTA FE

AB	AUROBINDO PHARMA	<u>0.035MG;0.4MG</u>	<u>A207535</u>	<u>001</u>	Feb 02, 2017
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NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB	AMNEAL PHARMS	<u>0.035MG;0.4MG</u>	<u>A078892</u>	<u>001</u>	Sep 26, 2011
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AB	BARR	<u>0.035MG;0.4MG</u>	<u>A078965</u>	<u>001</u>	Aug 05, 2010
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AB	LUPIN LTD	<u>0.035MG;0.4MG</u>	<u>A091332</u>	<u>001</u>	Mar 23, 2016
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AB	!	MYLAN LABS LTD	<u>0.035MG;0.4MG</u>	<u>A202086</u>	<u>001</u>	Apr 01, 2015
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AB	NAARI PTE LTD	<u>0.035MG;0.4MG</u>	<u>A207066</u>	<u>001</u>	Mar 29, 2017
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TABLET; ORAL-21

NORTREL 1/35-21

AB	BARR	<u>0.035MG;1MG</u>	<u>A072693</u>	<u>001</u>	Feb 28, 1992
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NORTREL 7/7/7

	BARR	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A075478</u>	<u>001</u>	Aug 30, 2002
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TABLET; ORAL-28

ALYACEN 1/35

AB	GLENMARK GENERICS	<u>0.035MG;1MG</u>	<u>A091634</u>	<u>001</u>	Jan 19, 2012
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ALYACEN 7/7/7

AB	GLENMARK GENERICS	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A091636</u>	<u>001</u>	Jan 19, 2012
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ARANELLE

AB	BARR	<u>0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG</u>	<u>A076783</u>	<u>001</u>	Sep 29, 2004
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BALZIVA-28

AB	!	BARR	<u>0.035MG;0.4MG</u>	<u>A076238</u>	<u>001</u>	Apr 22, 2004
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BRIELLYN

AB	GLENMARK GENERICS	<u>0.035MG;0.4MG</u>	<u>A090538</u>	<u>001</u>	Mar 22, 2011
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CYONANZ

AB	AUROBINDO PHARMA	<u>0.035MG;0.5MG</u>	<u>A207055</u>	<u>001</u>	Oct 21, 2016
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DASETTA 1/35

AB	NOVAST LABS LTD	<u>0.035MG;1MG</u>	<u>A090948</u>	<u>001</u>	Dec 22, 2011
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DASETTA 7/7/7

AB	!	NOVAST LABS LTD	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A090946</u>	<u>001</u>	Dec 22, 2011
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GILDAGIA

AB	VINTAGE PHARMS	<u>0.035MG;0.4MG</u>	<u>A078376</u>	<u>001</u>	Nov 06, 2012
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PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

NORETHINDRONE AND ETHINYL ESTRADIOL

AB	NAARI PTE LTD	0.035MG;1MG	A206864	001	Apr 28, 2017
AB	WATSON LABS	0.035MG;0.4MG	A078323	001	Feb 04, 2010
AB	WATSON LABS TEVA	0.035MG;1MG	A070687	001	Jan 29, 1987
<u>NORTREL 0.5/35-28</u>					
AB	BARR	0.035MG;0.5MG	A072695	001	Feb 28, 1992
<u>NORTREL 1/35-28</u>					
AB	! BARR	0.035MG;1MG	A072696	001	Feb 28, 1992
<u>NORTREL 7/7/7</u>					
AB	BARR	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A075478	002	Aug 30, 2002
<u>NYLIA 1/35</u>					
AB	AUROBINDO PHARMA	0.035MG;1MG	A207056	001	Oct 21, 2016
<u>NYLIA 7/7/7</u>					
AB	AUROBINDO PHARMA	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A207054	001	Oct 21, 2016
<u>PHILITH</u>					
AB	NOVAST LABS LTD	0.035MG;0.4MG	A090947	001	Dec 22, 2011
<u>TRI-NORINYL 28-DAY</u>					
AB	+! DR REDDYS LABS SA	0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG	N018977	002	Apr 13, 1984
<u>VYFEMLA</u>					
AB	LUPIN LTD	0.035MG;0.4MG	A201886	001	Sep 26, 2013
<u>WERA</u>					
AB	! NOVAST LABS LTD	0.035MG;0.5MG	A091204	001	Mar 27, 2012
NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)					
	WATSON LABS TEVA	0.035MG,0.035MG;0.5MG,1MG	A071044	001	Apr 01, 1988
TABLET, CHEWABLE; ORAL					

KAITLIB FE

AB	LUPIN LTD	0.025MG;0.8MG	A203448	001	Dec 17, 2015
<u>NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>					
AB	+! APIL	0.025MG;0.8MG	N022573	001	Dec 22, 2010
AB	MYLAN LABS LTD	0.025MG;0.8MG	A203371	001	Apr 23, 2014

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

CAPSULE; ORAL

GEMMILY

AB	XIROMED	0.02MG;1MG	A213317	001	Nov 09, 2020
<u>MERZEE</u>					
AB	SLAYBACK PHARMA LLC	0.02MG;1MG	A212706	001	Dec 18, 2020
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>					
AB	AMNEAL PHARMS	0.02MG;1MG	A214292	001	Jul 20, 2021
AB	GLENMARK PHARMS LTD	0.02MG;1MG	A213418	001	Jul 27, 2022
<u>TAYTULLA</u>					
AB	+! APIL	0.02MG;1MG	N204426	001	Apr 19, 2013

TABLET; ORAL

AUROVELA 24 FE

AB	AUROBINDO PHARMA	0.02MG;1MG	A207504	001	Jun 15, 2017
<u>BLISOVI 24 FE</u>					
AB	LUPIN LTD	0.02MG;1MG	A091398	001	Oct 28, 2015
<u>FINZALA</u>					
AB	TEVA PHARMS USA INC	0.02MG;1MG	A210087	001	Apr 07, 2020
<u>FYAVOLV</u>					
AB	LUPIN LTD	0.005MG;1MG	A204213	002	Dec 10, 2015
AB		0.0025MG;0.5MG	A204213	001	Dec 10, 2015
<u>GILDESS 24 FE</u>					
AB	VINTAGE PHARMS	0.02MG;1MG	A090293	001	Dec 01, 2014
<u>HAILEY 24 FE</u>					
AB	! GLENMARK PHARMS LTD	0.02MG;1MG	A204847	001	Nov 17, 2017
<u>LARIN 24 FE</u>					
AB	NOVAST LABS	0.02MG;1MG	A202994	001	Feb 18, 2015
<u>LERIBANE</u>					
AB	NOVAST LABS	0.0025MG;0.5MG	A203435	002	Jun 03, 2016
AB		0.005MG;1MG	A203435	001	Jun 03, 2016
<u>MIBELAS 24 FE</u>					
AB	LUPIN	0.02MG;1MG	A206287	001	May 24, 2016
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>					
AB	! BARR LABS INC	0.005MG;1MG	A076221	001	Nov 06, 2009
AB	GLENMARK GENERICS	0.0025MG;0.5MG	A203038	001	Apr 02, 2015
AB		0.005MG;1MG	A203038	002	Apr 02, 2015
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>					
AB	BARR LABS INC	0.02MG;1MG	A090938	001	Dec 01, 2014

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB	!	GLENMARK PHARMS LTD	0.02MG;1MG	A210369	001	Dec 26, 2017
AB		XIROMED	0.02MG;1MG	A209609	001	Jul 16, 2018
<u>OSHIH</u>						
AB		AUROBINDO PHARMA	0.02MG;1MG	A216558	001	Dec 06, 2023
LO LOESTRIN FE						
+! APIL						
0.01MG,0.01MG;1MG,N/A						
N022501 001 Oct 21, 2010						
TABLET; ORAL-21						
<u>AUROVELA 1.5/30</u>						
AB		AUROBINDO PHARMA	0.03MG;1.5MG	A207581	001	Jun 26, 2017
<u>AUROVELA 1/20</u>						
AB		AUROBINDO PHARMA	0.02MG;1MG	A207506	001	Jun 16, 2017
<u>HAILEY 1.5/30</u>						
AB		GLENMARK PHARMS	0.03MG;1.5MG	A209297	001	Jun 05, 2018
<u>JUNEL 1.5/30</u>						
AB		BARR	0.03MG;1.5MG	A076381	001	May 30, 2003
<u>JUNEL 1/20</u>						
AB		BARR	0.02MG;1MG	A076380	001	May 30, 2003
<u>LARIN 1.5/30</u>						
AB		NOVAST LABS	0.03MG;1.5MG	A202996	001	Mar 20, 2014
<u>LARIN 1/20</u>						
AB		NOVAST LABS	0.02MG;1MG	A202995	001	Dec 04, 2013
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>						
AB		GLENMARK PHARMS LTD	0.02MG;1MG	A206969	001	Jan 20, 2016
AB		MYLAN LABS LTD	0.02MG;1MG	A202771	001	Nov 06, 2013
AB			0.03MG;1.5MG	A202770	001	Feb 19, 2015

TABLET; ORAL-28

<u>AUROVELA FE 1.5/30</u>						
AB		AUROBINDO PHARMA	0.03MG;1.5MG	A207580	001	Jun 15, 2017
<u>AUROVELA FE 1/20</u>						
AB		AUROBINDO PHARMA	0.02MG;1MG	A207505	001	Jun 16, 2017
<u>BLISOVI FE 1/20</u>						
AB		LUPIN LTD	0.02MG;1MG	A201584	001	Nov 18, 2015
<u>CHABELINA FE</u>						
AB	!	NOVAST LABS	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A202962	001	Apr 15, 2020
<u>HAILEY FE 1.5/30</u>						
AB		GLENMARK PHARMS	0.03MG;1.5MG	A209031	001	Jun 05, 2018
<u>HAILEY FE 1/20</u>						
AB		GLENMARK PHARMS LTD	0.02MG;1MG	A206597	001	Nov 21, 2017
<u>JUNEL FE 1.5/30</u>						
AB	!	BARR	0.03MG;1.5MG	A076064	001	Sep 18, 2003
<u>JUNEL FE 1/20</u>						
AB		BARR	0.02MG;1MG	A076081	001	Sep 18, 2003
<u>LARIN FE 1.5/30</u>						
AB		NOVAST LABS	0.03MG;1.5MG	A091453	001	Aug 23, 2013
<u>LARIN FE 1/20</u>						
AB		NOVAST LABS	0.02MG;1MG	A091454	001	Aug 26, 2013
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>						
AB		MYLAN	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A205069	001	Jun 22, 2018
AB		MYLAN LABS LTD	0.02MG;1MG	A202772	001	Nov 14, 2013
AB			0.03MG;1.5MG	A202741	001	Feb 20, 2015
<u>TRI-LEGEST FE</u>						
AB		BARR	0.02MG,0.03MG,0.035MG;1MG,1MG,1MG	A076105	001	Oct 26, 2007
BLISOVI FE 1.5/30						
BX		LUPIN LTD	0.03MG;1.5MG	A201585	001	Nov 18, 2015

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

<u>ESTARYLLA</u>						
AB		XIROMED	0.035MG;0.25MG	A090794	001	Jan 30, 2013
<u>MILI</u>						
AB		AUROBINDO PHARMA	0.035MG;0.25MG	A205449	001	Jul 07, 2016
LTD						
<u>MONO-LINYAH</u>						
AB		NOVAST LABS LTD	0.035MG;0.25MG	A090523	001	May 23, 2012
<u>NORGESTIMATE AND ETHINYL ESTRADIOL</u>						
AB		AMNEAL PHARMS	0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG	A203873	001	May 12, 2016
AB	!	GLENMARK GENERICS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A200494	001	Jun 17, 2011
AB			0.035MG;0.25MG	A200538	001	Apr 05, 2012

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL

AB	!	GLENMARK PHARMS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A204057 001</u>	Feb 23, 2016
AB		LUPIN LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205588 001</u>	Apr 26, 2016
AB			<u>0.035MG; 0.25MG</u>	<u>A205630 001</u>	Oct 27, 2016
AB		LUPIN PHARMS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A200541 001</u>	Jun 25, 2012
AB		NAARI PTE LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A200383 001</u>	Apr 07, 2015
AB			<u>0.035MG; 0.25MG</u>	<u>A200384 001</u>	Apr 07, 2015
		<u>PREVIFEM</u>			
AB		VINTAGE PHARMS LLC	<u>0.035MG; 0.25MG</u>	<u>A076334 001</u>	Jan 09, 2004
		<u>SPRINTEC</u>			
AB	!	BARR	<u>0.035MG; 0.25MG</u>	<u>A075804 001</u>	Sep 25, 2002
		<u>TRI LO SPRINTEC</u>			
AB		BARR LABS INC	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A076784 001</u>	Jun 29, 2009
		<u>TRI-ESTARYLLA</u>			
AB		XIROMED	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090793 001</u>	Jan 30, 2013
		<u>TRI-LINYAH</u>			
AB		NOVAST LABS LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090524 001</u>	May 30, 2012
		<u>TRI-LO-ESTARYLLA</u>			
AB		XIROMED	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A091232 001</u>	Jun 29, 2015
		<u>TRI-LO-LINYAH</u>			
AB		NOVAST LABS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090541 001</u>	Sep 02, 2022
		<u>TRI-LO-MILI</u>			
AB		AUROBINDO PHARMA	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205762 001</u>	Nov 04, 2016
		<u>TRI-MILI</u>			
AB		AUROBINDO PHARMA LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205441 001</u>	Jul 06, 2016
		<u>TRI-SPRINTEC</u>			
AB		BARR	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A075808 001</u>	Dec 29, 2003

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

CRYSELLE

AB		DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 001</u>	Nov 30, 2001
		TABLET; ORAL-28			
		<u>CRYSELLE</u>			
AB	!	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 002</u>	Nov 30, 2001
		<u>ELINEST</u>			
AB		NOVAST LABS LTD	<u>0.03MG; 0.3MG</u>	<u>A091105 001</u>	Mar 28, 2012
		<u>LOW-OGESTREL-28</u>			
AB		DR REDDYS LABS SA	<u>0.03MG; 0.3MG</u>	<u>A075288 002</u>	Jul 28, 1999
		<u>TUROOZ</u>			
AB		LUPIN LTD	<u>0.03MG; 0.3MG</u>	<u>A202980 001</u>	Jul 31, 2023

ETHINYL ESTRADIOL; SEGESTERONE ACETATE

RING; VAGINAL

ANNOVERA

+! MAYNE PHARMA 0.013MG/24HR; 0.15MG/24HR N209627 001 Aug 10, 2018

ETHIODIZED OIL

OIL; INTRALYMPHATIC, INTRAUTERINE

LIPIODOL

+! GUERBET EQ 4.8GM IODINE/10ML (EQ 480MG IODINE/ML) N009190 001

ETHIONAMIDE

TABLET; ORAL

TRECATOR

+! WYETH PHARMS 250MG N013026 002

PRESCRIPTION DRUG PRODUCT LIST

ETHOSUXIMIDE

CAPSULE; ORAL

ETHOSUXIMIDE

<u>AB</u>	AKORN	<u>250MG</u>	<u>A040686</u>	<u>001</u>	May 28, 2008
<u>AB</u>	BIONPHARMA	<u>250MG</u>	<u>A040430</u>	<u>001</u>	Oct 28, 2002
<u>AB</u>	HERITAGE PHARMS INC	<u>250MG</u>	<u>A200892</u>	<u>001</u>	Sep 25, 2012
<u>AB</u>	PURACAP PHARM LLC	<u>250MG</u>	<u>A210654</u>	<u>001</u>	Mar 16, 2020
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A211928</u>	<u>001</u>	Feb 19, 2019

ZARONTIN

<u>AB</u>	+!	PARKE DAVIS	<u>250MG</u>	<u>N012380</u>	<u>001</u>
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SYRUP; ORAL

ETHOSUXIMIDE

<u>AA</u>	MIKART	<u>250MG/5ML</u>	<u>A040506</u>	<u>001</u>	Dec 22, 2003
<u>AA</u>	PHARM ASSOC	<u>250MG/5ML</u>	<u>A040253</u>	<u>001</u>	Nov 22, 2000

ZARONTIN

<u>AA</u>	+!	PARKE-DAVIS	<u>250MG/5ML</u>	<u>A080258</u>	<u>001</u>
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ETODOLAC

CAPSULE; ORAL

ETODOLAC

<u>AB</u>	ANI PHARMS	<u>200MG</u>	<u>A075126</u>	<u>001</u>	Sep 16, 1999
<u>AB</u>		<u>300MG</u>	<u>A075126</u>	<u>002</u>	Sep 16, 1999
<u>AB</u>	APOTEX	<u>200MG</u>	<u>A075419</u>	<u>001</u>	Jul 28, 2000
<u>AB</u>		<u>300MG</u>	<u>A075419</u>	<u>002</u>	Jul 28, 2000
<u>AB</u>	TARO	<u>200MG</u>	<u>A075078</u>	<u>001</u>	Apr 30, 1998
<u>AB</u>	!	<u>300MG</u>	<u>A075078</u>	<u>002</u>	Apr 30, 1998

TABLET; ORAL

ETODOLAC

<u>AB</u>	AMNEAL PHARMS CO	<u>400MG</u>	<u>A208834</u>	<u>001</u>	Jun 07, 2018
<u>AB</u>		<u>500MG</u>	<u>A208834</u>	<u>002</u>	Jun 07, 2018
<u>AB</u>	APOTEX INC	<u>400MG</u>	<u>A076004</u>	<u>001</u>	Dec 03, 2002
<u>AB</u>		<u>500MG</u>	<u>A076004</u>	<u>002</u>	Dec 03, 2002
<u>AB</u>	BAYSHORE PHARMS LLC	<u>400MG</u>	<u>A210704</u>	<u>001</u>	Dec 16, 2020
<u>AB</u>		<u>500MG</u>	<u>A210704</u>	<u>002</u>	Dec 16, 2020
<u>AB</u>	SANDOZ	<u>400MG</u>	<u>A074903</u>	<u>001</u>	Apr 11, 1997
<u>AB</u>		<u>500MG</u>	<u>A074903</u>	<u>002</u>	Apr 19, 1999
<u>AB</u>	TARO PHARM INDS	<u>400MG</u>	<u>A075074</u>	<u>001</u>	Mar 11, 1998
<u>AB</u>	!	<u>500MG</u>	<u>A075074</u>	<u>002</u>	Apr 25, 2000

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

<u>AB</u>	BAYSHORE PHARMS LLC	<u>400MG</u>	<u>A212263</u>	<u>001</u>	Nov 24, 2020
<u>AB</u>		<u>500MG</u>	<u>A212263</u>	<u>002</u>	Nov 24, 2020
<u>AB</u>		<u>600MG</u>	<u>A212263</u>	<u>003</u>	Nov 24, 2020
<u>AB</u>	TARO	<u>400MG</u>	<u>A076174</u>	<u>001</u>	Mar 13, 2003
<u>AB</u>		<u>500MG</u>	<u>A076174</u>	<u>002</u>	Mar 13, 2003
<u>AB</u>		<u>600MG</u>	<u>A076174</u>	<u>003</u>	Mar 13, 2003
<u>AB</u>	TEVA	<u>400MG</u>	<u>A075665</u>	<u>003</u>	Feb 05, 2001
<u>AB</u>		<u>500MG</u>	<u>A075665</u>	<u>002</u>	Jul 31, 2000
<u>AB</u>	!	<u>600MG</u>	<u>A075665</u>	<u>001</u>	Jul 31, 2000
<u>AB</u>	ZYDUS PHARMS	<u>400MG</u>	<u>A091134</u>	<u>001</u>	Jan 23, 2014
<u>AB</u>		<u>500MG</u>	<u>A091134</u>	<u>002</u>	Jan 23, 2014
<u>AB</u>		<u>600MG</u>	<u>A091134</u>	<u>003</u>	Jan 23, 2014

ETOMIDATE

INJECTABLE; INJECTION

AMIDATE

<u>AP</u>	+!	HOSPIRA	<u>2MG/ML</u>	<u>N018227</u>	<u>001</u>	Sep 07, 1982
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ETOMIDATE

<u>AP</u>	CAPLIN	<u>2MG/ML</u>	<u>A215028</u>	<u>001</u>	Dec 18, 2020
<u>AP</u>	EUGIA PHARMA	<u>2MG/ML</u>	<u>A206126</u>	<u>001</u>	Feb 24, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>2MG/ML</u>	<u>A209058</u>	<u>001</u>	Apr 18, 2017
<u>AP</u>	HIKMA	<u>2MG/ML</u>	<u>A074593</u>	<u>001</u>	Nov 04, 1996
<u>AP</u>		<u>2MG/ML</u>	<u>A202354</u>	<u>001</u>	Feb 25, 2016
<u>AP</u>	MYLAN LABS LTD	<u>2MG/ML</u>	<u>A201044</u>	<u>001</u>	Feb 07, 2017
<u>AP</u>	ZYDUS PHARMS	<u>2MG/ML</u>	<u>A202360</u>	<u>001</u>	Jul 18, 2014

ETONOGESTREL

IMPLANT; IMPLANTATION

NEXPLANON

	+!	ORGANON	<u>68MG/IMPLANT</u>	<u>N021529</u>	<u>002</u>	May 13, 2011
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PRESCRIPTION DRUG PRODUCT LIST

ETOPOSID

CAPSULE; ORAL

ETOPOSID

! MYLAN

50MG

A075635 001 Sep 19, 2001

INJECTABLE; INJECTION

ETOPOSID

<u>AP</u>	ACCORD HLTHCARE	<u>20MG/ML</u>	<u>A074513</u>	<u>001</u>	Mar 14, 1996
<u>AP</u>	! FRESINIUS KABI USA	<u>20MG/ML</u>	<u>A074983</u>	<u>001</u>	Sep 30, 1998
<u>AP</u>	HIKMA	<u>20MG/ML</u>	<u>A074290</u>	<u>001</u>	Jul 17, 1995
<u>AP</u>	MEITHEAL	<u>20MG/ML</u>	<u>A074529</u>	<u>001</u>	Jul 24, 1996

ETOPOSID PHOSPHATE

INJECTABLE; INJECTION

ETOPHOS PRESERVATIVE FREE

+! CHEPLAPHARM

EQ 100MG BASE/VIAL

N020457 001 May 17, 1996

ETRASIMOD ARGININE

TABLET; ORAL

VELSIPITY

+! PFIZER

EQ 2MG BASE

N216956 001 Oct 12, 2023

ETRAVIRINE

TABLET; ORAL

ETRAVIRINE

<u>AB</u>	AMNEAL	<u>100MG</u>	<u>A214196</u>	<u>002</u>	Jun 14, 2021
<u>AB</u>		<u>200MG</u>	<u>A214196</u>	<u>003</u>	Jun 14, 2021
<u>AB</u>	CARNEGIE	<u>100MG</u>	<u>A215402</u>	<u>001</u>	Apr 13, 2022
<u>AB</u>		<u>200MG</u>	<u>A215402</u>	<u>002</u>	Apr 13, 2022
	<u>INTELENCE</u>				
<u>AB</u>	+ JANSSEN R AND D	<u>100MG</u>	<u>N022187</u>	<u>001</u>	Jan 18, 2008
<u>AB</u>	+!	<u>200MG</u>	<u>N022187</u>	<u>002</u>	Dec 22, 2010
	+	25MG	N022187	003	Mar 26, 2012

EVEROLIMUS

TABLET; ORAL

AFINITOR

<u>AB</u>	+ NOVARTIS	<u>2.5MG</u>	<u>N022334</u>	<u>003</u>	Jul 09, 2010
<u>AB</u>	+!	<u>5MG</u>	<u>N022334</u>	<u>001</u>	Mar 30, 2009
<u>AB</u>	+	<u>7.5MG</u>	<u>N022334</u>	<u>004</u>	Mar 30, 2012
<u>AB</u>	+	<u>10MG</u>	<u>N022334</u>	<u>002</u>	Mar 30, 2009

EVEROLIMUS

<u>AB</u>	ALKEM LABS LTD	<u>0.25MG</u>	<u>A214138</u>	<u>001</u>	Nov 26, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A214138</u>	<u>002</u>	Nov 26, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A214138</u>	<u>003</u>	Nov 26, 2021
<u>AB</u>		<u>1MG</u>	<u>A214138</u>	<u>004</u>	Nov 26, 2021
<u>AB</u>	BIOCON PHARMA	<u>2.5MG</u>	<u>A214182</u>	<u>001</u>	Feb 11, 2021
<u>AB</u>		<u>5MG</u>	<u>A214182</u>	<u>002</u>	Feb 11, 2021
<u>AB</u>		<u>7.5MG</u>	<u>A214182</u>	<u>003</u>	Feb 11, 2021
<u>AB</u>		<u>10MG</u>	<u>A214182</u>	<u>004</u>	Feb 11, 2021
<u>AB</u>	BRECKENRIDGE	<u>0.25MG</u>	<u>A205432</u>	<u>001</u>	May 20, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A205432</u>	<u>002</u>	May 20, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A205432</u>	<u>003</u>	May 20, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A205426</u>	<u>001</u>	Mar 05, 2021
<u>AB</u>		<u>5MG</u>	<u>A205426</u>	<u>002</u>	Mar 05, 2021
<u>AB</u>		<u>7.5MG</u>	<u>A205426</u>	<u>003</u>	Mar 05, 2021
<u>AB</u>		<u>10MG</u>	<u>A205426</u>	<u>004</u>	Mar 05, 2021
<u>AB</u>	HIKMA	<u>0.25MG</u>	<u>A206133</u>	<u>001</u>	Apr 12, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A206133</u>	<u>002</u>	Apr 12, 2018
<u>AB</u>		<u>0.75MG</u>	<u>A206133</u>	<u>003</u>	Apr 12, 2018
<u>AB</u>		<u>1MG</u>	<u>A206133</u>	<u>004</u>	Nov 18, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A207486</u>	<u>001</u>	Jun 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A207486</u>	<u>002</u>	Jun 08, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A207486</u>	<u>003</u>	Jun 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A207486</u>	<u>004</u>	Nov 23, 2021
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A212936</u>	<u>001</u>	Jun 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A212936</u>	<u>002</u>	Jun 08, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A212936</u>	<u>003</u>	Jun 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A212936</u>	<u>004</u>	Jun 08, 2020
<u>AB</u>	PAR PHARM	<u>0.25MG</u>	<u>A205775</u>	<u>001</u>	Oct 18, 2021
<u>AB</u>		<u>0.5MG</u>	<u>A205775</u>	<u>002</u>	Oct 18, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A205775</u>	<u>003</u>	Oct 18, 2021
<u>AB</u>		<u>1MG</u>	<u>A205775</u>	<u>004</u>	Oct 18, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A207934</u>	<u>001</u>	Dec 09, 2019
<u>AB</u>		<u>5MG</u>	<u>A207934</u>	<u>002</u>	Dec 09, 2019

PRESCRIPTION DRUG PRODUCT LIST

EVEROLIMUS

TABLET; ORAL

EVEROLIMUS

<u>AB</u>		<u>7.5MG</u>	<u>A207934 003</u>	Dec 09, 2019
<u>AB</u>		<u>10MG</u>	<u>A207934 004</u>	Dec 09, 2020
<u>AB</u>	TEVA PHARMS USA	<u>2.5MG</u>	<u>A210050 001</u>	Dec 09, 2019
<u>AB</u>		<u>5MG</u>	<u>A210050 002</u>	Dec 09, 2019
<u>AB</u>		<u>7.5MG</u>	<u>A210050 003</u>	Dec 09, 2019
<u>AB</u>		<u>10MG</u>	<u>A210050 004</u>	Dec 09, 2019

ZORTRESS

<u>AB</u>	+	NOVARTIS	<u>0.25MG</u>	<u>N021560 001</u>	Apr 20, 2010
<u>AB</u>	+		<u>0.5MG</u>	<u>N021560 002</u>	Apr 20, 2010
<u>AB</u>	+		<u>0.75MG</u>	<u>N021560 003</u>	Apr 20, 2010
<u>AB</u>	+	!	<u>1MG</u>	<u>N021560 004</u>	Aug 10, 2018

TABLET, FOR SUSPENSION; ORAL

AFINITOR DISPERZ

<u>AB</u>	+	NOVARTIS PHARM	<u>2MG</u>	<u>N203985 001</u>	Aug 29, 2012
<u>AB</u>	+		<u>3MG</u>	<u>N203985 002</u>	Aug 29, 2012
<u>AB</u>	+	!	<u>5MG</u>	<u>N203985 003</u>	Aug 29, 2012

EVEROLIMUS

<u>AB</u>		MYLAN	<u>2MG</u>	<u>A210130 001</u>	Apr 19, 2019
<u>AB</u>			<u>3MG</u>	<u>A210130 002</u>	Apr 19, 2019
<u>AB</u>			<u>5MG</u>	<u>A210130 003</u>	Apr 19, 2019

EXEMESTANE

TABLET; ORAL

AROMASIN

<u>AB</u>	+	!	PFIZER	<u>25MG</u>	<u>N020753 001</u>	Oct 21, 1999
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EXEMESTANE

<u>AB</u>		BRECKENRIDGE	<u>25MG</u>	<u>A211031 001</u>	Feb 21, 2019
<u>AB</u>		CIPLA	<u>25MG</u>	<u>A210323 001</u>	Apr 27, 2018
<u>AB</u>		EUGIA PHARMA	<u>25MG</u>	<u>A216454 001</u>	May 20, 2022
<u>AB</u>		HIKMA	<u>25MG</u>	<u>A077431 001</u>	Apr 01, 2011
<u>AB</u>		QILU	<u>25MG</u>	<u>A213547 001</u>	Apr 13, 2020
<u>AB</u>		RISING	<u>25MG</u>	<u>A203315 001</u>	Mar 10, 2017
<u>AB</u>		UPSHER SMITH LABS	<u>25MG</u>	<u>A209208 001</u>	Jul 26, 2017
<u>AB</u>		ZYDUS PHARMS	<u>25MG</u>	<u>A202602 001</u>	Oct 03, 2018

EXENATIDE SYNTHETIC

INJECTABLE; SUBCUTANEOUS

BYETTA

	+	!	ASTRAZENECA AB	300MCG/1.2ML (250MCG/ML)	<u>N021773 001</u>	Apr 28, 2005
	+	!		600MCG/2.4ML (250MCG/ML)	<u>N021773 002</u>	Apr 28, 2005

SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

BYDUREON BCISE

	+	!	ASTRAZENECA AB	2MG/0.85ML (2MG/0.85ML)	<u>N209210 001</u>	Oct 20, 2017
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EZETIMIBE

TABLET; ORAL

EZETIMIBE

<u>AB</u>		ACCORD HLTHCARE	<u>10MG</u>	<u>A211550 001</u>	Oct 26, 2018
<u>AB</u>		ALKEM LABS LTD	<u>10MG</u>	<u>A209234 001</u>	Dec 21, 2017
<u>AB</u>		AMNEAL PHARMS CO	<u>10MG</u>	<u>A208803 001</u>	Jun 12, 2017
<u>AB</u>		AUROBINDO PHARMA	<u>10MG</u>	<u>A209838 001</u>	Aug 25, 2017
<u>AB</u>		GLENMARK PHARMS LTD	<u>10MG</u>	<u>A078560 001</u>	Jun 26, 2015
<u>AB</u>		HETERO LABS LTD III	<u>10MG</u>	<u>A210859 001</u>	Jul 26, 2022
<u>AB</u>		OHM LABS INC	<u>10MG</u>	<u>A207311 001</u>	Jun 12, 2017
<u>AB</u>		ORIENT PHARMA	<u>10MG</u>	<u>A215693 001</u>	Sep 13, 2022
<u>AB</u>		SANDOZ	<u>10MG</u>	<u>A203931 001</u>	Jun 12, 2017
<u>AB</u>		SCIEGEN PHARMS INC	<u>10MG</u>	<u>A210673 001</u>	Oct 23, 2020
<u>AB</u>		WATSON LABS INC	<u>10MG</u>	<u>A200831 001</u>	Jun 12, 2017
<u>AB</u>		ZYDUS PHARMS	<u>10MG</u>	<u>A204331 001</u>	Jun 12, 2017

ZETIA

<u>AB</u>	+	!	ORGANON	<u>10MG</u>	<u>N021445 001</u>	Oct 25, 2002
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EZETIMIBE; SIMVASTATIN

TABLET; ORAL

EZETIMIBE AND SIMVASTATIN

<u>AB</u>		ALKEM LABS LTD	<u>10MG; 10MG</u>	<u>A209222 001</u>	Dec 22, 2017
<u>AB</u>			<u>10MG; 20MG</u>	<u>A209222 002</u>	Dec 22, 2017
<u>AB</u>			<u>10MG; 40MG</u>	<u>A209222 003</u>	Dec 22, 2017
<u>AB</u>			<u>10MG; 80MG</u>	<u>A209222 004</u>	Dec 22, 2017
<u>AB</u>		AMNEAL PHARMS CO	<u>10MG; 10MG</u>	<u>A208831 001</u>	Nov 21, 2017
<u>AB</u>			<u>10MG; 20MG</u>	<u>A208831 002</u>	Nov 21, 2017

PRESCRIPTION DRUG PRODUCT LIST

EZETIMIBE; SIMVASTATIN

TABLET; ORAL

EZETIMIBE AND SIMVASTATIN

<u>AB</u>		<u>10MG; 40MG</u>	<u>A208831 003</u>	Nov 21, 2017
<u>AB</u>		<u>10MG; 80MG</u>	<u>A208831 004</u>	Nov 21, 2017
<u>AB</u>	DR REDDYS LABS SA	<u>10MG; 10MG</u>	<u>A200909 001</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 20MG</u>	<u>A200909 002</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 40MG</u>	<u>A200909 003</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 80MG</u>	<u>A200909 004</u>	Apr 26, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG; 10MG</u>	<u>A208699 001</u>	Jun 27, 2019
<u>AB</u>		<u>10MG; 20MG</u>	<u>A208699 002</u>	Jun 27, 2019
<u>AB</u>		<u>10MG; 40MG</u>	<u>A208699 003</u>	Jun 27, 2019
<u>AB</u>		<u>10MG; 80MG</u>	<u>A208699 004</u>	Jun 27, 2019
<u>AB</u>	WATSON LABS INC	<u>10MG; 10MG</u>	<u>A202968 001</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 20MG</u>	<u>A202968 002</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 40MG</u>	<u>A202968 003</u>	Apr 26, 2017
<u>AB</u>		<u>10MG; 80MG</u>	<u>A202968 004</u>	Apr 26, 2017

VYTORIN

<u>AB</u>	+	ORGANON	<u>10MG; 10MG</u>	<u>N021687 001</u>	Jul 23, 2004
<u>AB</u>	+		<u>10MG; 20MG</u>	<u>N021687 002</u>	Jul 23, 2004
<u>AB</u>	+		<u>10MG; 40MG</u>	<u>N021687 003</u>	Jul 23, 2004
<u>AB</u>	+	!	<u>10MG; 80MG</u>	<u>N021687 004</u>	Jul 23, 2004

FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

<u>AB</u>		APOTEX	<u>125MG</u>	<u>A091480 001</u>	Jul 22, 2011
<u>AB</u>			<u>250MG</u>	<u>A091480 002</u>	Jul 22, 2011
<u>AB</u>			<u>500MG</u>	<u>A091480 003</u>	Jul 22, 2011
<u>AB</u>		AUROBINDO PHARMA LTD	<u>125MG</u>	<u>A091114 001</u>	Mar 21, 2011
<u>AB</u>			<u>250MG</u>	<u>A091114 002</u>	Mar 21, 2011
<u>AB</u>			<u>500MG</u>	<u>A091114 003</u>	Mar 21, 2011
<u>AB</u>		HETERO LABS LTD V	<u>125MG</u>	<u>A202438 001</u>	Sep 10, 2014
<u>AB</u>			<u>250MG</u>	<u>A202438 002</u>	Sep 10, 2014
<u>AB</u>			<u>500MG</u>	<u>A202438 003</u>	Sep 10, 2014
<u>AB</u>		MACLEODS PHARMS LTD	<u>125MG</u>	<u>A201022 001</u>	Jan 12, 2012
<u>AB</u>			<u>250MG</u>	<u>A201022 002</u>	Jan 12, 2012
<u>AB</u>			<u>500MG</u>	<u>A201022 003</u>	Jan 12, 2012
<u>AB</u>		TEVA PHARMS	<u>125MG</u>	<u>A077487 001</u>	Aug 24, 2007
<u>AB</u>			<u>250MG</u>	<u>A077487 002</u>	Aug 24, 2007
<u>AB</u>		!	<u>500MG</u>	<u>A077487 003</u>	Aug 24, 2007

FAMOTIDINE

FOR SUSPENSION; ORAL

FAMOTIDINE

<u>AB</u>		AJANTA PHARMA LTD	<u>40MG/5ML</u>	<u>A217529 001</u>	Sep 18, 2023
<u>AB</u>		AKORN	<u>40MG/5ML</u>	<u>A201995 001</u>	May 30, 2014
<u>AB</u>		ALKEM LABS LTD	<u>40MG/5ML</u>	<u>A216400 001</u>	May 15, 2023
<u>AB</u>		AMNEALS PHARMS	<u>40MG/5ML</u>	<u>A216427 001</u>	Aug 04, 2022
<u>AB</u>		ANNORA PHARMA	<u>40MG/5ML</u>	<u>A217330 001</u>	Aug 17, 2023
<u>AB</u>		CARNEGIE	<u>40MG/5ML</u>	<u>A217137 001</u>	Jul 07, 2023
<u>AB</u>		LUPIN LTD	<u>40MG/5ML</u>	<u>A090440 001</u>	Jun 29, 2010
<u>AB</u>		MICRO LABS	<u>40MG/5ML</u>	<u>A217842 001</u>	Sep 14, 2023
<u>AB</u>		NAVINTA LLC	<u>40MG/5ML</u>	<u>A091020 001</u>	May 27, 2010
<u>AB</u>		NOVEL LABS INC	<u>40MG/5ML</u>	<u>A201695 001</u>	Dec 17, 2012
<u>AB</u>	!	NOVITIUM PHARMA	<u>40MG/5ML</u>	<u>A215043 001</u>	Apr 20, 2021
<u>AB</u>		UPSHER SMITH LABS	<u>40MG/5ML</u>	<u>A217655 001</u>	Jun 16, 2023

INJECTABLE; INJECTION

FAMOTIDINE

<u>AP</u>		FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A075709 001</u>	Apr 16, 2001
<u>AP</u>	!	HIKMA	<u>10MG/ML</u>	<u>A075488 001</u>	Apr 16, 2001
<u>AP</u>		MYLAN LABS LTD	<u>10MG/ML</u>	<u>A078641 001</u>	Jun 25, 2008
<u>AP</u>		SAGENT	<u>10MG/ML</u>	<u>A075651 001</u>	Apr 16, 2001
<u>AP</u>			<u>10MG/ML</u>	<u>A075684 001</u>	Apr 16, 2001

FAMOTIDINE PRESERVATIVE FREE

<u>AP</u>		FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A075813 001</u>	Apr 16, 2001
<u>AP</u>	!	HIKMA	<u>10MG/ML</u>	<u>A075486 001</u>	Apr 16, 2001
<u>AP</u>		MYLAN LABS LTD	<u>10MG/ML</u>	<u>A078642 001</u>	Jun 25, 2008
<u>AP</u>		SAGENT	<u>10MG/ML</u>	<u>A075622 001</u>	Apr 16, 2001
<u>AP</u>			<u>10MG/ML</u>	<u>A075825 001</u>	Apr 17, 2001

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

!	BAXTER HLTHCARE	0.4MG/ML	A075591 001	May 10, 2001
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PRESCRIPTION DRUG PRODUCT LIST

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>20MG</u>	<u>A078916 001</u>	May 22, 2009
<u>AB</u>		<u>40MG</u>	<u>A078916 002</u>	May 22, 2009
<u>AB</u>	ALKEM LABS LTD	<u>20MG</u>	<u>A215630 001</u>	Jan 07, 2022
<u>AB</u>		<u>20MG</u>	<u>A217375 001</u>	Apr 24, 2023
<u>AB</u>		<u>40MG</u>	<u>A215630 002</u>	Jan 07, 2022
<u>AB</u>		<u>40MG</u>	<u>A217375 002</u>	Apr 24, 2023
<u>AB</u>	ANNORA PHARMA	<u>20MG</u>	<u>A215767 001</u>	Nov 04, 2021
<u>AB</u>		<u>40MG</u>	<u>A215767 002</u>	Nov 04, 2021
<u>AB</u>	APOTEX	<u>20MG</u>	<u>A075611 001</u>	Jul 23, 2001
<u>AB</u>		<u>40MG</u>	<u>A075611 002</u>	Jul 23, 2001
<u>AB</u>	ASCENT PHARMS INC	<u>20MG</u>	<u>A215689 001</u>	Oct 15, 2021
<u>AB</u>		<u>40MG</u>	<u>A215689 002</u>	Oct 15, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A206530 001</u>	Dec 22, 2015
<u>AB</u>	!	<u>40MG</u>	<u>A206530 002</u>	Dec 22, 2015
<u>AB</u>	CARLSBAD	<u>20MG</u>	<u>A075805 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075805 002</u>	Apr 16, 2001
<u>AB</u>	CHARTWELL RX	<u>20MG</u>	<u>A075786 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075786 002</u>	Apr 16, 2001
<u>AB</u>	CONTRACT PHARMACAL	<u>20MG</u>	<u>A217669 001</u>	Dec 20, 2023
<u>AB</u>		<u>40MG</u>	<u>A217669 002</u>	Dec 20, 2023
<u>AB</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A075718 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075718 002</u>	Apr 16, 2001
<u>AB</u>	GRAVITI PHARMS	<u>20MG</u>	<u>A218181 001</u>	Dec 22, 2023
<u>AB</u>		<u>40MG</u>	<u>A218181 002</u>	Dec 22, 2023
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>20MG</u>	<u>A075511 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075511 002</u>	Apr 16, 2001
<u>AB</u>	MANKIND PHARMA	<u>20MG</u>	<u>A075302 001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075302 002</u>	Apr 16, 2001
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A216441 001</u>	Jun 03, 2022
<u>AB</u>		<u>40MG</u>	<u>A216441 002</u>	Jun 03, 2022

FAMOTIDINE; IBUPROFEN

TABLET; ORAL

DUEXIS

<u>AB</u>	+! HORIZON	<u>26.6MG; 800MG</u>	<u>N022519 001</u>	Apr 23, 2011
	<u>IBUPROFEN AND FAMOTIDINE</u>			
<u>AB</u>	ALKEM LABS LTD	<u>26.6MG; 800MG</u>	<u>A211890 001</u>	Aug 03, 2021
<u>AB</u>	ASCENT PHARMS INC	<u>26.6MG; 800MG</u>	<u>A216814 001</u>	Mar 15, 2023
<u>AB</u>	TEVA PHARMS USA	<u>26.6MG; 800MG</u>	<u>A211278 001</u>	Oct 29, 2021

FEBUXOSTAT

TABLET; ORAL

FEBUXOSTAT

<u>AB</u>	ALEMBIC	<u>40MG</u>	<u>A205421 001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205421 002</u>	Jul 01, 2019
<u>AB</u>	ALKEM LABS LTD	<u>40MG</u>	<u>A212924 001</u>	Dec 07, 2021
<u>AB</u>		<u>80MG</u>	<u>A212924 002</u>	Dec 07, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>40MG</u>	<u>A210741 001</u>	Oct 25, 2023
<u>AB</u>		<u>80MG</u>	<u>A210741 002</u>	Oct 25, 2023
<u>AB</u>	DR REDDYS	<u>40MG</u>	<u>A205374 001</u>	Oct 22, 2020
<u>AB</u>		<u>80MG</u>	<u>A205374 002</u>	Oct 22, 2020
<u>AB</u>	HIKMA	<u>40MG</u>	<u>A205414 001</u>	Oct 15, 2019
<u>AB</u>		<u>80MG</u>	<u>A205414 002</u>	Oct 15, 2019
<u>AB</u>	INDOCO	<u>40MG</u>	<u>A210292 001</u>	Dec 30, 2019
<u>AB</u>		<u>80MG</u>	<u>A210292 002</u>	Dec 30, 2019
<u>AB</u>	MACLEODS PHARMS LTD	<u>40MG</u>	<u>A207293 001</u>	Sep 28, 2023
<u>AB</u>		<u>80MG</u>	<u>A207293 002</u>	Sep 28, 2023
<u>AB</u>	MSN	<u>40MG</u>	<u>A210461 001</u>	Dec 30, 2019
<u>AB</u>		<u>80MG</u>	<u>A210461 002</u>	Dec 30, 2019
<u>AB</u>	PRINSTON INC	<u>40MG</u>	<u>A206266 001</u>	Mar 28, 2022
<u>AB</u>		<u>80MG</u>	<u>A206266 002</u>	Mar 28, 2022
<u>AB</u>	SUN PHARM	<u>40MG</u>	<u>A205467 001</u>	Jul 01, 2019
<u>AB</u>		<u>80MG</u>	<u>A205467 002</u>	Jul 01, 2019
<u>AB</u>	SUNSHINE	<u>40MG</u>	<u>A213069 001</u>	Jun 02, 2020
<u>AB</u>		<u>80MG</u>	<u>A213069 002</u>	Jun 02, 2020
<u>AB</u>	ZYDUS LIFESCIENCES	<u>40MG</u>	<u>A205443 001</u>	Jan 09, 2023
<u>AB</u>		<u>80MG</u>	<u>A205443 002</u>	Jan 09, 2023

PRESCRIPTION DRUG PRODUCT LIST

FEBUXOSTAT

TABLET; ORAL

ULORIC

<u>AB</u>	+	TAKEDA PHARMS USA	<u>40MG</u>	<u>N021856</u>	<u>001</u>	Feb 13, 2009
<u>AB</u>	+	!	<u>80MG</u>	<u>N021856</u>	<u>002</u>	Feb 13, 2009

FEDRATINIB HYDROCHLORIDE

CAPSULE; ORAL

INREBIC

+	!	IMPACT	EQ 100MG BASE	N212327	001	Aug 16, 2019
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FELBAMATE

SUSPENSION; ORAL

FELBAMATE

<u>AB</u>		AMNEAL PHARMS	<u>600MG/5ML</u>	<u>A202385</u>	<u>001</u>	Dec 16, 2011
<u>AB</u>		NOVITIUM PHARMA	<u>600MG/5ML</u>	<u>A211333</u>	<u>001</u>	May 31, 2019
<u>AB</u>		TARO	<u>600MG/5ML</u>	<u>A206314</u>	<u>001</u>	Jun 16, 2017

FELBATOL

<u>AB</u>	+	!	MYLAN SPECIALITY LP	<u>600MG/5ML</u>	<u>N020189</u>	<u>003</u>	Jul 29, 1993
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TABLET; ORAL

FELBAMATE

<u>AB</u>		ALVOGEN	<u>400MG</u>	<u>A204595</u>	<u>001</u>	Jan 11, 2016
<u>AB</u>			<u>600MG</u>	<u>A204595</u>	<u>002</u>	Jan 11, 2016
<u>AB</u>		AMNEAL PHARMS	<u>400MG</u>	<u>A201680</u>	<u>001</u>	Sep 13, 2011
<u>AB</u>			<u>600MG</u>	<u>A201680</u>	<u>002</u>	Sep 13, 2011
<u>AB</u>		ANI PHARMS	<u>400MG</u>	<u>A202284</u>	<u>001</u>	Nov 04, 2015
<u>AB</u>			<u>600MG</u>	<u>A202284</u>	<u>002</u>	Nov 04, 2015
<u>AB</u>		CADILA	<u>400MG</u>	<u>A208970</u>	<u>001</u>	May 30, 2017
<u>AB</u>			<u>600MG</u>	<u>A208970</u>	<u>002</u>	May 30, 2017
<u>AB</u>		TARO	<u>400MG</u>	<u>A207093</u>	<u>001</u>	Apr 20, 2017
<u>AB</u>			<u>600MG</u>	<u>A207093</u>	<u>002</u>	Apr 20, 2017

FELBATOL

<u>AB</u>	+	!	MYLAN SPECIALITY LP	<u>400MG</u>	<u>N020189</u>	<u>001</u>	Jul 29, 1993
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<u>AB</u>	+	!		<u>600MG</u>	<u>N020189</u>	<u>002</u>	Jul 29, 1993
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FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A203417</u>	<u>001</u>	Jan 17, 2013
<u>AB</u>			<u>5MG</u>	<u>A203417</u>	<u>002</u>	Jan 17, 2013
<u>AB</u>			<u>10MG</u>	<u>A203417</u>	<u>003</u>	Jan 17, 2013
<u>AB</u>		GLENMARK GENERICS	<u>2.5MG</u>	<u>A090365</u>	<u>001</u>	Dec 17, 2010
<u>AB</u>			<u>5MG</u>	<u>A090365</u>	<u>002</u>	Dec 17, 2010
<u>AB</u>			<u>10MG</u>	<u>A090365</u>	<u>003</u>	Dec 17, 2010
<u>AB</u>		HERITAGE PHARMS	<u>2.5MG</u>	<u>A201964</u>	<u>001</u>	Nov 08, 2013
<u>AB</u>			<u>5MG</u>	<u>A201964</u>	<u>002</u>	Nov 08, 2013
<u>AB</u>			<u>10MG</u>	<u>A201964</u>	<u>003</u>	Nov 08, 2013
<u>AB</u>		ORBION PHARMS	<u>2.5MG</u>	<u>A203032</u>	<u>001</u>	May 21, 2015
<u>AB</u>			<u>5MG</u>	<u>A203032</u>	<u>002</u>	May 21, 2015
<u>AB</u>			<u>10MG</u>	<u>A203032</u>	<u>003</u>	May 21, 2015
<u>AB</u>		SUN PHARM INDS LTD	<u>2.5MG</u>	<u>A091200</u>	<u>001</u>	Dec 13, 2013
<u>AB</u>			<u>5MG</u>	<u>A091200</u>	<u>002</u>	Dec 13, 2013
<u>AB</u>			<u>10MG</u>	<u>A091200</u>	<u>003</u>	Dec 13, 2013
<u>AB</u>		TORRENT PHARMS LTD	<u>2.5MG</u>	<u>A202170</u>	<u>001</u>	Nov 28, 2011
<u>AB</u>			<u>5MG</u>	<u>A202170</u>	<u>002</u>	Nov 28, 2011
<u>AB</u>		!	<u>10MG</u>	<u>A202170</u>	<u>003</u>	Nov 28, 2011
<u>AB</u>		VINTAGE PHARMS LLC	<u>2.5MG</u>	<u>A200815</u>	<u>001</u>	Oct 28, 2011
<u>AB</u>			<u>5MG</u>	<u>A200815</u>	<u>002</u>	Oct 28, 2011
<u>AB</u>			<u>10MG</u>	<u>A200815</u>	<u>003</u>	Oct 28, 2011
<u>AB</u>		YILING	<u>2.5MG</u>	<u>A210847</u>	<u>001</u>	Oct 26, 2018
<u>AB</u>			<u>5MG</u>	<u>A210847</u>	<u>002</u>	Oct 26, 2018
<u>AB</u>			<u>10MG</u>	<u>A210847</u>	<u>003</u>	Oct 26, 2018
<u>AB</u>		YUNG SHIN PHARM	<u>2.5MG</u>	<u>A204800</u>	<u>001</u>	Apr 29, 2019
<u>AB</u>			<u>5MG</u>	<u>A204800</u>	<u>002</u>	Apr 29, 2019
<u>AB</u>			<u>10MG</u>	<u>A204800</u>	<u>003</u>	Apr 29, 2019

FENFLURAMINE HYDROCHLORIDE

SOLUTION; ORAL

FINTEPLA

+	!	UCB INC	EQ 2.2MG BASE/ML	N212102	001	Jun 25, 2020
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PRESCRIPTION DRUG PRODUCT LIST

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

<u>AB</u>	+	LUPIN	<u>43MG</u>	<u>N021695</u>	<u>001</u>	Nov 30, 2004
<u>AB</u>	+	!	<u>130MG</u>	<u>N021695</u>	<u>003</u>	Nov 30, 2004

FENOFIBRATE

<u>AB</u>		SUN PHARM INDS LTD	<u>43MG</u>	<u>A201748</u>	<u>001</u>	Oct 31, 2014
<u>AB</u>			<u>130MG</u>	<u>A201748</u>	<u>002</u>	Oct 31, 2014

FENOFIBRATE (MICRONIZED)

<u>AB</u>		AJANTA PHARMA LTD	<u>67MG</u>	<u>A210705</u>	<u>001</u>	Sep 10, 2018
<u>AB</u>			<u>134MG</u>	<u>A210705</u>	<u>002</u>	Sep 10, 2018
<u>AB</u>	!		<u>200MG</u>	<u>A210705</u>	<u>003</u>	Sep 10, 2018
<u>AB</u>		ALEMBIC	<u>67MG</u>	<u>A213842</u>	<u>001</u>	Oct 19, 2020
<u>AB</u>			<u>134MG</u>	<u>A213842</u>	<u>002</u>	Oct 19, 2020
<u>AB</u>			<u>200MG</u>	<u>A213842</u>	<u>003</u>	Oct 19, 2020
<u>AB</u>		ANI PHARMS	<u>67MG</u>	<u>A209504</u>	<u>001</u>	Apr 30, 2018
<u>AB</u>			<u>134MG</u>	<u>A209504</u>	<u>002</u>	Apr 30, 2018
<u>AB</u>			<u>200MG</u>	<u>A209504</u>	<u>003</u>	Apr 30, 2018
<u>AB</u>		APOTEX	<u>43MG</u>	<u>A202252</u>	<u>001</u>	Jul 26, 2013
<u>AB</u>			<u>130MG</u>	<u>A202252</u>	<u>002</u>	Jul 26, 2013
<u>AB</u>		AUROBINDO PHARMA LTD	<u>67MG</u>	<u>A212232</u>	<u>001</u>	Sep 20, 2021
<u>AB</u>			<u>134MG</u>	<u>A212232</u>	<u>002</u>	Sep 20, 2021
<u>AB</u>			<u>200MG</u>	<u>A212232</u>	<u>003</u>	Sep 20, 2021
<u>AB</u>		DR REDDYS LABS SA	<u>43MG</u>	<u>A090859</u>	<u>001</u>	Mar 01, 2012
<u>AB</u>			<u>130MG</u>	<u>A090859</u>	<u>002</u>	Mar 01, 2012
<u>AB</u>		GLENMARK PHARMS LTD	<u>67MG</u>	<u>A205566</u>	<u>001</u>	Apr 07, 2017
<u>AB</u>			<u>134MG</u>	<u>A205566</u>	<u>002</u>	Apr 07, 2017
<u>AB</u>			<u>200MG</u>	<u>A205566</u>	<u>003</u>	Apr 07, 2017
<u>AB</u>		INVAGEN PHARMS	<u>67MG</u>	<u>A207378</u>	<u>001</u>	Mar 28, 2017
<u>AB</u>			<u>134MG</u>	<u>A207378</u>	<u>002</u>	Mar 28, 2017
<u>AB</u>			<u>200MG</u>	<u>A207378</u>	<u>003</u>	Mar 28, 2017
<u>AB</u>		REYOUNG	<u>67MG</u>	<u>A207805</u>	<u>001</u>	Nov 16, 2017
<u>AB</u>			<u>134MG</u>	<u>A207805</u>	<u>002</u>	Nov 16, 2017
<u>AB</u>			<u>200MG</u>	<u>A207805</u>	<u>003</u>	Nov 16, 2017
<u>AB</u>		RHODES PHARMS	<u>67MG</u>	<u>A075753</u>	<u>001</u>	Sep 03, 2002
<u>AB</u>			<u>134MG</u>	<u>A075753</u>	<u>002</u>	Apr 09, 2002
<u>AB</u>			<u>200MG</u>	<u>A075753</u>	<u>003</u>	Apr 09, 2002
<u>AB</u>		TORRENT	<u>67MG</u>	<u>A210782</u>	<u>001</u>	Jun 26, 2018
<u>AB</u>			<u>134MG</u>	<u>A210782</u>	<u>002</u>	Jun 26, 2018
<u>AB</u>			<u>200MG</u>	<u>A210782</u>	<u>003</u>	Jun 26, 2018
		LIPOFEN				
	+	CIPHER PHARMS INC	50MG	N021612	001	Jan 11, 2006
	+	!	150MG	N021612	003	Jan 11, 2006

TABLET; ORAL

FENOFIBRATE

<u>AB</u>		AJANTA PHARMA LTD	<u>54MG</u>	<u>A210138</u>	<u>001</u>	Jul 23, 2018
<u>AB</u>			<u>160MG</u>	<u>A210138</u>	<u>002</u>	Jul 23, 2018
<u>AB</u>		ALEMBIC	<u>48MG</u>	<u>A210476</u>	<u>001</u>	Aug 09, 2019
<u>AB</u>			<u>145MG</u>	<u>A210476</u>	<u>002</u>	Aug 09, 2019
<u>AB</u>		AMNEAL	<u>48MG</u>	<u>A209951</u>	<u>001</u>	Feb 09, 2018
<u>AB</u>			<u>54MG</u>	<u>A209950</u>	<u>001</u>	Mar 19, 2018
<u>AB</u>			<u>145MG</u>	<u>A209951</u>	<u>002</u>	Feb 09, 2018
<u>AB</u>			<u>160MG</u>	<u>A209950</u>	<u>002</u>	Mar 19, 2018
<u>AB</u>		AUROBINDO PHARMA	<u>48MG</u>	<u>A205118</u>	<u>001</u>	May 05, 2016
<u>AB</u>			<u>54MG</u>	<u>A216798</u>	<u>001</u>	Sep 27, 2022
<u>AB</u>			<u>145MG</u>	<u>A205118</u>	<u>002</u>	May 05, 2016
<u>AB</u>			<u>160MG</u>	<u>A216798</u>	<u>002</u>	Sep 27, 2022
<u>AB</u>		AUSTARPHARMA	<u>48MG</u>	<u>A208476</u>	<u>001</u>	Feb 10, 2021
<u>AB</u>			<u>54MG</u>	<u>A207803</u>	<u>001</u>	Dec 19, 2017
<u>AB</u>			<u>145MG</u>	<u>A208476</u>	<u>002</u>	Feb 10, 2021
<u>AB</u>			<u>160MG</u>	<u>A207803</u>	<u>002</u>	Dec 19, 2017
<u>AB</u>		CHARTWELL RX	<u>54MG</u>	<u>A209660</u>	<u>001</u>	Feb 11, 2019
<u>AB</u>			<u>160MG</u>	<u>A209660</u>	<u>002</u>	Feb 11, 2019
<u>AB</u>		CIPLA	<u>48MG</u>	<u>A208709</u>	<u>001</u>	Dec 15, 2016
<u>AB</u>			<u>145MG</u>	<u>A208709</u>	<u>002</u>	Dec 15, 2016
<u>AB</u>		CREEKWOOD PHARMS	<u>40MG</u>	<u>A217732</u>	<u>001</u>	Sep 07, 2023
<u>AB</u>			<u>120MG</u>	<u>A217732</u>	<u>002</u>	Sep 07, 2023
<u>AB</u>		DR REDDYS	<u>54MG</u>	<u>A210670</u>	<u>001</u>	Sep 06, 2019
<u>AB</u>			<u>160MG</u>	<u>A210670</u>	<u>002</u>	Sep 06, 2019
<u>AB</u>		GRAVITI PHARMS	<u>48MG</u>	<u>A211122</u>	<u>001</u>	Mar 18, 2020
<u>AB</u>			<u>54MG</u>	<u>A210606</u>	<u>001</u>	Aug 17, 2018
<u>AB</u>			<u>145MG</u>	<u>A211122</u>	<u>002</u>	Mar 18, 2020

PRESCRIPTION DRUG PRODUCT LIST

FENOFIBRATE

TABLET; ORAL

FENOFIBRATE

<u>AB</u>		<u>160MG</u>	<u>A210606 002</u>	Aug 17, 2018
<u>AB</u>	HETERO LABS LTD III	<u>48MG</u>	<u>A204598 001</u>	Jul 12, 2016
<u>AB</u>		<u>145MG</u>	<u>A204598 002</u>	Jul 12, 2016
<u>AB</u>	IMPAX LABS	<u>54MG</u>	<u>A076509 001</u>	Mar 26, 2008
<u>AB</u>	!	<u>160MG</u>	<u>A076509 002</u>	Mar 26, 2008
<u>AB</u>	LUPIN LTD	<u>48MG</u>	<u>A090856 001</u>	Dec 23, 2011
<u>AB</u>		<u>54MG</u>	<u>A204019 001</u>	Aug 17, 2015
<u>AB</u>		<u>145MG</u>	<u>A090856 002</u>	Dec 23, 2011
<u>AB</u>		<u>160MG</u>	<u>A204019 002</u>	Aug 17, 2015
<u>AB</u>	MANKIND PHARMA	<u>54MG</u>	<u>A213864 001</u>	Jun 12, 2020
<u>AB</u>		<u>160MG</u>	<u>A213864 002</u>	Jun 12, 2020
<u>AB</u>	MYLAN	<u>54MG</u>	<u>A076520 001</u>	Oct 25, 2007
<u>AB</u>		<u>160MG</u>	<u>A076520 003</u>	Oct 25, 2007
<u>AB</u>	MYLAN PHARMS INC	<u>40MG</u>	<u>A204475 001</u>	Jun 23, 2016
<u>AB</u>		<u>48MG</u>	<u>A202856 001</u>	Dec 07, 2012
<u>AB</u>		<u>120MG</u>	<u>A204475 002</u>	Jun 23, 2016
<u>AB</u>		<u>145MG</u>	<u>A202856 002</u>	Dec 07, 2012
<u>AB</u>	PRINSTON INC	<u>48MG</u>	<u>A211080 001</u>	Aug 28, 2018
<u>AB</u>		<u>145MG</u>	<u>A211080 002</u>	Aug 28, 2018
<u>AB</u>	RHODES PHARMS	<u>54MG</u>	<u>A076433 001</u>	May 13, 2005
<u>AB</u>		<u>160MG</u>	<u>A076433 002</u>	May 13, 2005
<u>AB</u>	SUN PHARM	<u>48MG</u>	<u>A200884 001</u>	Sep 07, 2017
<u>AB</u>		<u>145MG</u>	<u>A200884 002</u>	Sep 07, 2017
<u>AB</u>	SUN PHARM INDS LTD	<u>54MG</u>	<u>A076635 001</u>	Oct 31, 2005
<u>AB</u>		<u>160MG</u>	<u>A076635 003</u>	Oct 31, 2005
<u>AB</u>	VALEANT PHARMS NORTH	<u>48MG</u>	<u>A090715 001</u>	Apr 05, 2012
<u>AB</u>		<u>145MG</u>	<u>A090715 002</u>	Apr 05, 2012

FENOGLIDE

<u>AB</u>	+ SALIX	<u>40MG</u>	<u>N022118 001</u>	Aug 10, 2007
<u>AB</u>	+!	<u>120MG</u>	<u>N022118 002</u>	Aug 10, 2007

TRICOR

<u>AB</u>	+ ABBVIE	<u>48MG</u>	<u>N021656 001</u>	Nov 05, 2004
<u>AB</u>	+!	<u>145MG</u>	<u>N021656 002</u>	Nov 05, 2004
BX	TRIGLIDE + SKYEPHARMA AG	160MG	N021350 002	May 07, 2005
	FENOFIBRATE SUN PHARM INDS LTD	107MG	A076635 002	Oct 31, 2005

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

CORLOPAM

<u>AP</u>	+! HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>N019922 001</u>	Sep 23, 1997
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FENOLDOPAM MESYLATE

<u>AP</u>	HIKMA	<u>EQ 10MG BASE/ML</u>	<u>A076582 001</u>	Oct 12, 2004
<u>AP</u>	SANDOZ	<u>EQ 10MG BASE/ML</u>	<u>A077155 001</u>	Feb 15, 2005

FENOPROFEN CALCIUM

CAPSULE; ORAL

FENOPROFEN CALCIUM

<u>AB</u>	MISEMER	<u>EQ 200MG BASE</u>	<u>A215548 001</u>	May 16, 2023
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A215548 003</u>	May 16, 2023
<u>AB</u>	RISING	<u>EQ 400MG BASE</u>	<u>A214475 001</u>	Jul 18, 2022

NALFON

<u>AB</u>	+ XSPIRE PHARMA	<u>EQ 200MG BASE</u>	<u>N017604 003</u>	
<u>AB</u>	+!	<u>EQ 400MG BASE</u>	<u>N017604 004</u>	Jul 21, 2009

FENOPROFEN CALCIUM

MISEMER

		EQ 300MG BASE	A215548 002	May 16, 2023
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TABLET; ORAL

FENOPROFEN CALCIUM

	!	XSPIRE PHARMA	EQ 600MG BASE	A072267 001	Aug 17, 1988
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FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL-100

<u>AB</u>	AVEVA	<u>100MCG/HR</u>	<u>A077449 004</u>	Oct 20, 2008
<u>AB</u>	KINDEVA	<u>100MCG/HR</u>	<u>A202097 005</u>	Nov 04, 2016
<u>AB</u>	MYLAN TECHNOLOGIES	<u>100MCG/HR</u>	<u>A076258 004</u>	Jan 28, 2005
<u>AB</u>	SPECIX LLC	<u>100MCG/HR</u>	<u>A077154 004</u>	Feb 09, 2011

FENTANYL-12

<u>AB</u>	AVEVA	<u>12.5MCG/HR</u>	<u>A077449 005</u>	Sep 11, 2015
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PRESCRIPTION DRUG PRODUCT LIST

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL-12

AB	KINDEVA	12.5MCG/HR	A202097 001	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	12.5MCG/HR	A076258 005	Jan 23, 2007
AB	SPECGX LLC	12.5MCG/HR	A077154 005	Jun 11, 2015

FENTANYL-25

AB	AVEVA	25MCG/HR	A077449 001	Oct 20, 2008
AB	KINDEVA	25MCG/HR	A202097 002	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	25MCG/HR	A076258 001	Jan 28, 2005
AB	! SPECGX LLC	25MCG/HR	A077154 001	Feb 09, 2011

FENTANYL-37

AB	AVEVA	37.5MCG/HR	A077449 006	Dec 06, 2017
AB	MYLAN TECHNOLOGIES	37.5MCG/HR	A076258 006	Dec 29, 2014
AB	SPECGX LLC	37.5MCG/HR	A077154 006	Jan 14, 2020

FENTANYL-50

AB	AVEVA	50MCG/HR	A077449 002	Oct 20, 2008
AB	KINDEVA	50MCG/HR	A202097 003	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	50MCG/HR	A076258 002	Jan 28, 2005
AB	SPECGX LLC	50MCG/HR	A077154 002	Feb 09, 2011

FENTANYL-62

AB	AVEVA	62.5MCG/HR	A077449 007	Dec 06, 2017
AB	MYLAN TECHNOLOGIES	62.5MCG/HR	A076258 007	Dec 29, 2014
AB	SPECGX LLC	62.5MCG/HR	A077154 007	Jan 14, 2020

FENTANYL-75

AB	AVEVA	75MCG/HR	A077449 003	Oct 20, 2008
AB	KINDEVA	75MCG/HR	A202097 004	Nov 04, 2016
AB	MYLAN TECHNOLOGIES	75MCG/HR	A076258 003	Jan 28, 2005
AB	SPECGX LLC	75MCG/HR	A077154 003	Feb 09, 2011

FENTANYL-87

AB	AVEVA	87.5MCG/HR	A077449 008	Dec 06, 2017
AB	MYLAN TECHNOLOGIES	87.5MCG/HR	A076258 008	Dec 29, 2014

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE

AP	FRESENIUS KABI USA	EQ 0.05MG BASE/ML	A212086 001	Sep 01, 2020
AP	+! HIKMA	EQ 0.05MG BASE/ML	N019101 001	Jul 11, 1984
AP	HOSPIRA	EQ 0.05MG BASE/ML	N019115 001	Jan 12, 1985

FENTANYL CITRATE PRESERVATIVE FREE

AP	FRESENIUS KABI USA	EQ 0.05MG BASE/ML	A210762 001	May 03, 2019
AP	HOSPIRA	EQ 0.05MG BASE/ML	A072786 001	Sep 24, 1991

SUBLIMAZE PRESERVATIVE FREE

AP	+! RISING	EQ 0.05MG BASE/ML	N016619 001	
	FENTANYL CITRATE			
	+! HIKMA	EQ 0.025MG BASE/0.5ML	N019101 002	Jan 20, 2023

TABLET; BUCCAL, SUBLINGUAL

FENTORA

+	CEPHALON	EQ 0.1MG BASE	N021947 001	Sep 25, 2006
+		EQ 0.2MG BASE	N021947 002	Sep 25, 2006
+!		EQ 0.4MG BASE	N021947 003	Sep 25, 2006
+		EQ 0.6MG BASE	N021947 004	Sep 25, 2006
+		EQ 0.8MG BASE	N021947 005	Sep 25, 2006

TROCHE/LOZENGE; TRANSMUCOSAL

ACTIQ

+	CEPHALON	EQ 0.2MG BASE	N020747 001	Nov 04, 1998
+!		EQ 0.4MG BASE	N020747 002	Nov 04, 1998
+		EQ 0.6MG BASE	N020747 003	Nov 04, 1998
+		EQ 0.8MG BASE	N020747 004	Nov 04, 1998
+		EQ 1.2MG BASE	N020747 005	Nov 04, 1998
+		EQ 1.6MG BASE	N020747 006	Nov 04, 1998

FERRIC CARBOXYMALTOS

SOLUTION; INTRAVENOUS

INJECTAFER

+!	AM REGENT	750MG IRON/15ML (50MG IRON/ML)	N203565 001	Jul 25, 2013
+!		100MG IRON/2ML (50MG IRON/ML)	N203565 004	Feb 04, 2022
+!		500MG IRON/10ML (50MG IRON/ML)	N203565 002	Oct 08, 2020
+!		1GM IRON/20ML (50MG IRON/ML)	N203565 003	Apr 28, 2021

PRESCRIPTION DRUG PRODUCT LIST

FERRIC CITRATE

TABLET; ORAL

AURYXIA

+! KERYX BIOPHARMS EQ 210MG IRON N205874 001 Sep 05, 2014

FERRIC DERISOMALTOSE

SOLUTION; INTRAVENOUS

MONOFERRIC

+! PHARMACOSMOS AS 1GM/10ML (100MG/ML) N208171 003 Jan 16, 2020

FERRIC HEXACYANOFERRATE (II)

CAPSULE; ORAL

RADIOGARDASE (PRUSSIAN BLUE)

+! HEYL CHEMISCH 500MG N021626 001 Oct 02, 2003

FERRIC MALTOI

CAPSULE; ORAL

ACCRUFER

+! SHIELD TX 30MG IRON N212320 001 Jul 25, 2019

FERRIC OXYHYDROXIDE

INJECTABLE; INJECTION

FERLECIT**AB** +! SANOFI AVENTIS US EQ 62.5MG IRON/5ML (EQ 12.5MG IRON/ML) **N020955 001** Feb 18, 1999SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE**AB** WEST-WARD PHARMS EQ 62.5MG IRON/5ML (EQ 12.5MG IRON/ML) **A078215 001** Mar 31, 2011

INT

INFED

BP +! ALLERGAN EQ 100MG IRON/2ML (EQ 50MG IRON/ML) N017441 001

INJECTABLE; INTRAVENOUS

VENOFER

+ AM REGENT EQ 50MG IRON/2.5ML (EQ 20MG IRON/ML) N021135 002 Mar 20, 2005

+! EQ 100MG IRON/5ML (EQ 20MG IRON/ML) N021135 001 Nov 06, 2000

+ EQ 200MG IRON/10ML (EQ 20MG IRON/ML) N021135 004 Feb 09, 2007

TABLET, CHEWABLE; ORAL

VELPHORO

+! VIFOR FRESENIUS EQ 500MG IRON N205109 001 Nov 27, 2013

FERUMOXYTOL

SOLUTION; INTRAVENOUS

FERAHEME**AB** +! COVIS EQ 510MG IRON/17ML (EQ 30MG IRON/ML) **N022180 001** Jun 30, 2009FERUMOXYTOL**AB** SANDOZ EQ 510MG IRON/17ML (EQ 30MG IRON/ML) **A206604 001** Jan 15, 2021FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

FESOTERODINE FUMARATE**AB** ALEMBIC 4MG **A204973 001** Jan 04, 2023**AB** 8MG **A204973 002** Jan 04, 2023**AB** ALKEM LABS LTD 4MG **A204827 001** Dec 10, 2015**AB** 8MG **A204827 002** Dec 10, 2015**AB** AMNEAL PHARMS NY 4MG **A205002 001** Jan 04, 2023**AB** 8MG **A205002 002** Jan 04, 2023**AB** ANI PHARMS 4MG **A204504 001** Jan 04, 2023**AB** 8MG **A204504 002** Jan 04, 2023**AB** AUROBINDO PHARMA 4MG **A205007 001** Feb 17, 2017**AB** 8MG **A205007 002** Feb 17, 2017**AB** DR REDDYS 4MG **A204975 001** Aug 13, 2019**AB** 8MG **A204975 002** Aug 13, 2019**AB** HETERO LABS LTD V 4MG **A204792 001** Jan 09, 2024**AB** 8MG **A204792 002** Jan 09, 2024**AB** ZYDUS PHARMS 4MG **A204946 001** Oct 03, 2017**AB** 8MG **A204946 002** Oct 03, 2017TOVIAZ**AB** + PFIZER 4MG **N022030 001** Oct 31, 2008**AB** +! 8MG **N022030 002** Oct 31, 2008FEXINIDAZOLE

TABLET; ORAL

FEXINIDAZOLE

+! SANOFI 600MG N214429 001 Jul 16, 2021

PRESCRIPTION DRUG PRODUCT LIST

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

FEXOFENADINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>30MG</u>	<u>A076502 001</u>	Apr 11, 2006
<u>AB</u>		<u>60MG</u>	<u>A076502 002</u>	Apr 11, 2006
<u>AB</u>		<u>180MG</u>	<u>A076502 003</u>	Apr 11, 2006
<u>AB</u>	RISING	<u>60MG</u>	<u>A077081 003</u>	Apr 11, 2008
<u>AB</u>		<u>180MG</u>	<u>A077081 001</u>	Apr 16, 2007
<u>AB</u>	TEVA	<u>30MG</u>	<u>A076447 001</u>	Sep 01, 2005
<u>AB</u>		<u>60MG</u>	<u>A076447 002</u>	Sep 01, 2005
<u>AB</u>		<u>180MG</u>	<u>A076447 003</u>	Sep 01, 2005

FEZOLINETANT

TABLET; ORAL

VEOZAH

+	!	ASTELLAS	45MG	N216578 001	May 12, 2023
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FIDAXOMICIN

FOR SUSPENSION; ORAL

DIFICID

+	!	CUBIST PHARMS LLC	40MG/ML	N213138 001	Jan 24, 2020
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TABLET; ORAL

DIFICID

+	!	CUBIST PHARMS LLC	200MG	N201699 001	May 27, 2011
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FINASTERIDE

TABLET; ORAL

FINASTERIDE

<u>AB</u>	ACCORD HLTHCARE	<u>1MG</u>	<u>A091643 001</u>	Nov 05, 2013
<u>AB</u>		<u>5MG</u>	<u>A090121 001</u>	Feb 23, 2010
<u>AB</u>	ALKEM LABS LTD	<u>1MG</u>	<u>A207750 001</u>	Jan 06, 2017
<u>AB</u>		<u>5MG</u>	<u>A204304 001</u>	Jan 05, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078341 001</u>	Oct 30, 2007
<u>AB</u>	AUROBINDO PHARMA LTD	<u>1MG</u>	<u>A203687 001</u>	Nov 05, 2013
<u>AB</u>	CIPLA	<u>1MG</u>	<u>A077335 001</u>	Nov 20, 2014
<u>AB</u>	DR REDDYS LABS INC	<u>1MG</u>	<u>A076436 001</u>	Jul 28, 2006
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076437 001</u>	Feb 28, 2007
<u>AB</u>	HETERO LABS LTD III	<u>1MG</u>	<u>A090060 001</u>	Jul 01, 2013
<u>AB</u>		<u>5MG</u>	<u>A090061 001</u>	Jun 07, 2010
<u>AB</u>	SUN PHARM	<u>1MG</u>	<u>A090508 001</u>	Jul 01, 2013
<u>AB</u>		<u>5MG</u>	<u>A090507 001</u>	Aug 16, 2011
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076511 001</u>	Dec 15, 2006
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A078900 001</u>	Dec 28, 2009

PROPECIA

<u>AB</u>	+	!	ORGANON	<u>1MG</u>	<u>N020788 001</u>	Dec 19, 1997
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PROSCAR

<u>AB</u>	+	!	ORGANON	<u>5MG</u>	<u>N020180 001</u>	Jun 19, 1992
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FINASTERIDE; TADALAFIL

CAPSULE; ORAL

ENTADFI

+	!	BLUE WATER BIOTECH	5MG; 5MG	N215423 001	Dec 09, 2021
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FINERENONE

TABLET; ORAL

KERENDIA

+		BAYER HLTHCARE	10MG	N215341 001	Jul 09, 2021
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+	!		20MG	N215341 002	Jul 09, 2021
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FINGOLIMOD HYDROCHLORIDE

CAPSULE; ORAL

FINGOLIMOD HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 0.5MG BASE</u>	<u>A207991 001</u>	Oct 28, 2020
<u>AB</u>	ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A208004 001</u>	Dec 30, 2020
<u>AB</u>	APOTEX	<u>EQ 0.5MG BASE</u>	<u>A207993 001</u>	Dec 18, 2020
<u>AB</u>	BIONPHARMA	<u>EQ 0.5MG BASE</u>	<u>A210252 001</u>	May 24, 2023
<u>AB</u>	DR REDDYS	<u>EQ 0.5MG BASE</u>	<u>A208000 001</u>	Mar 05, 2021
<u>AB</u>	EZRA VENTURES	<u>EQ 0.5MG BASE</u>	<u>A207945 001</u>	Dec 06, 2023
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 0.5MG BASE</u>	<u>A207985 001</u>	Jun 18, 2020
<u>AB</u>	HEC PHARM CO LTD	<u>EQ 0.5MG BASE</u>	<u>A207939 001</u>	Nov 10, 2021
<u>AB</u>	HETERO LABS LTD V	<u>EQ 0.5MG BASE</u>	<u>A207933 001</u>	May 18, 2020
<u>AB</u>	MYLAN	<u>EQ 0.5MG BASE</u>	<u>A208005 001</u>	Jan 19, 2021
<u>AB</u>	PRINSTON INC	<u>EQ 0.5MG BASE</u>	<u>A208003 001</u>	Sep 07, 2022
<u>AB</u>	SUN PHARM	<u>EQ 0.5MG BASE</u>	<u>A208014 001</u>	Dec 04, 2019

PRESCRIPTION DRUG PRODUCT LIST

FINGOLIMOD HYDROCHLORIDE

CAPSULE; ORAL

FINGOLIMOD HYDROCHLORIDE

AB	TEVA PHARMS USA	EQ 0.5MG BASE	A208008 001	Jul 02, 2020
AB	ZYDUS PHARMS	EQ 0.5MG BASE	A207994 001	Oct 14, 2020

GILENYA

AB	+! NOVARTIS	EQ 0.5MG BASE	N022527 001	Sep 21, 2010
	+	EQ 0.25MG BASE	N022527 002	May 11, 2018

FINGOLIMOD LAURYL SULFATE

TABLET, ORALLY DISINTEGRATING; ORAL

TASCENSO ODT

+	CYCLE	EQ 0.25MG BASE	N214962 001	Dec 23, 2021
+	!	EQ 0.5MG BASE	N214962 002	Dec 09, 2022

FISH OIL TRIGLYCERIDES

EMULSION; INTRAVENOUS

OMEGAVEN

+	!	FRESENIUS KABI USA	5GM/50ML (0.1GM/ML)	N210589 001	Jul 27, 2018
+	!		10GM/100ML (0.1GM/ML)	N210589 002	Jul 27, 2018

FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL

EMULSION; INTRAVENOUS

SMOFLIPID 20%

+	!	FRESENIUS KABI USA	3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (100ML)	N207648 001	Jul 13, 2016
+	!		3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (250ML)	N207648 002	Jul 13, 2016
+	!		3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (500ML)	N207648 003	Jul 13, 2016
+	!		3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (1000ML)	N207648 004	Aug 10, 2018

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

AB	EPIC PHARMA	100MG	A076835 001	Nov 30, 2005
AB	! PADAGIS US	100MG	A076831 001	Dec 16, 2004

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

AB	AMNEAL PHARM	50MG	A075442 001	Jul 31, 2001
AB		100MG	A075442 002	Jul 31, 2001
AB		150MG	A075442 003	Jul 31, 2001
AB	ANI PHARMS	50MG	A075882 001	Oct 28, 2002
AB		100MG	A075882 002	Oct 28, 2002
AB		150MG	A075882 003	Oct 28, 2002
AB	AUROBINDO PHARMA LTD	50MG	A202821 001	Nov 03, 2017
AB		100MG	A202821 002	Nov 03, 2017
AB		150MG	A202821 003	Nov 03, 2017
AB	BEXIMCO PHARMS USA	50MG	A210683 001	Sep 16, 2020
AB		100MG	A210683 002	Sep 16, 2020
AB		150MG	A210683 003	Sep 16, 2020
AB	HIKMA	50MG	A076278 001	Jan 14, 2003
AB		100MG	A076278 002	Jan 14, 2003
AB	!	150MG	A076278 003	Jan 14, 2003
AB	SUN PHARM INDS LTD	50MG	A076421 001	Mar 28, 2003
AB		100MG	A076421 002	Mar 28, 2003
AB		150MG	A076421 003	Mar 28, 2003
AB	YICHANG HUMANWELL	50MG	A215599 001	Sep 08, 2022
AB		100MG	A215599 002	Sep 08, 2022
AB		150MG	A215599 003	Sep 08, 2022

FLIBANSERIN

TABLET; ORAL

ADDYI

+	!	SPROUT PHARMS	100MG	N022526 001	Aug 18, 2015
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FLORBETABEN F-18

SOLUTION; INTRAVENOUS

NEURACEQ

+	!	LIFE MOLECULAR	1.4-135mCi/ML	N204677 001	Mar 19, 2014
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PRESCRIPTION DRUG PRODUCT LIST

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

+!	AVID RADIOPHARMS INC	10-100ML (13.5-51mCi/ML)	N202008 004	Oct 13, 2023
+!		10-50ML (13.5-51mCi/ML)	N202008 003	Apr 06, 2012

FLORTAUCIPIR F-18

SOLUTION; INTRAVENOUS

TAUVID

+!	AVID RADIOPHARMS INC	30ML (8.1-51mCi/ML)	N212123 001	May 28, 2020
+!		50ML (8.1-51mCi/ML)	N212123 002	May 28, 2020
+!		50ML (8.1-100mCi/ML)	N212123 003	Jul 01, 2022
+!		100ML (8.1-100mCi/ML)	N212123 004	Jul 01, 2022

FLOTUFOLASTAT F-18 GALLIUM

SOLUTION; INTRAVENOUS

POSLUMA

+!	BLUE EARTH	25ML (8-158mCi/ML)	N216023 001	May 25, 2023
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FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

<u>AP</u>	FRESENIUS KABI USA	<u>500MG/VIAL</u>	<u>A075837 001</u>	Feb 22, 2001
<u>AP</u>	! HIKMA	<u>500MG/VIAL</u>	<u>A075387 001</u>	Apr 16, 2000

FLUCICLOVINE F-18

SOLUTION; INTRAVENOUS

AXUMIN

+!	BLUE EARTH	9-221mCi/ML	N208054 001	May 27, 2016
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FLUCONAZOLE

FOR SUSPENSION; ORAL

DIFLUCAN

<u>AB</u>	+ PFIZER	<u>50MG/5ML</u>	<u>N020090 001</u>	Dec 23, 1993
<u>AB</u>	+!	<u>200MG/5ML</u>	<u>N020090 002</u>	Dec 23, 1993

FLUCONAZOLE

<u>AB</u>	AUROBINDO PHARMA	<u>50MG/5ML</u>	<u>A079150 001</u>	Sep 18, 2009
<u>AB</u>		<u>200MG/5ML</u>	<u>A079150 002</u>	Sep 18, 2009
<u>AB</u>	HAINAN POLY	<u>200MG/5ML</u>	<u>A215738 001</u>	Sep 21, 2023

INJECTABLE; INJECTION

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

<u>AP</u>	FRESENIUS KABI USA	<u>200MG/100ML (2MG/ML)</u>	<u>A076145 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076145 002</u>	Jul 29, 2004
<u>AP</u>	HIKMA	<u>200MG/100ML (2MG/ML)</u>	<u>A076087 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076087 003</u>	Jul 29, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>200MG/100ML (2MG/ML)</u>	<u>A076736 001</u>	Aug 23, 2005

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML (2MG/ML)</u>	<u>A076766 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076766 002</u>	Jul 29, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>200MG/100ML (2MG/ML)</u>	<u>A078698 001</u>	Jan 30, 2012
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A078698 002</u>	Jan 30, 2012
<u>AP</u>	HOSPIRA	<u>200MG/100ML (2MG/ML)</u>	<u>A076303 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076303 002</u>	Jul 29, 2004
<u>AP</u>	! INFORLIFE	<u>200MG/100ML (2MG/ML)</u>	<u>A079104 001</u>	Jul 30, 2009
<u>AP</u>	!	<u>400MG/200ML (2MG/ML)</u>	<u>A079104 002</u>	Jul 30, 2009
<u>AP</u>	WEST-WARD PHARMS INT	<u>200MG/100ML (2MG/ML)</u>	<u>A078107 001</u>	Jul 30, 2008
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A078107 002</u>	Jul 30, 2008
<u>AP</u>	WOODWARD	<u>200MG/100ML (2MG/ML)</u>	<u>A077909 001</u>	May 26, 2010
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A077909 002</u>	May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

	HIKMA	100MG/50ML (2MG/ML)	A076087 002	Sep 26, 2008
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FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	WOODWARD	100MG/50ML (2MG/ML)	A077909 003	Apr 20, 2015
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TABLET; ORAL

DIFLUCAN

<u>AB</u>	+ PFIZER	<u>50MG</u>	<u>N019949 001</u>	Jan 29, 1990
<u>AB</u>	+	<u>100MG</u>	<u>N019949 002</u>	Jan 29, 1990
<u>AB</u>	+	<u>150MG</u>	<u>N019949 004</u>	Jun 30, 1994
<u>AB</u>	+!	<u>200MG</u>	<u>N019949 003</u>	Jan 29, 1990

FLUCONAZOLE

<u>AB</u>	ANI PHARMS	<u>50MG</u>	<u>A078423 001</u>	Mar 07, 2011
<u>AB</u>		<u>100MG</u>	<u>A078423 002</u>	Mar 07, 2011

PRESCRIPTION DRUG PRODUCT LIST

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

<u>AB</u>		<u>150MG</u>	<u>A078423 003</u>	Mar 07, 2011
<u>AB</u>		<u>200MG</u>	<u>A078423 004</u>	Mar 07, 2011
<u>AB</u>	AUROBINDO PHARMA	<u>50MG</u>	<u>A077731 001</u>	Oct 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A077731 002</u>	Oct 07, 2008
<u>AB</u>		<u>150MG</u>	<u>A077731 003</u>	Oct 07, 2008
<u>AB</u>		<u>200MG</u>	<u>A077731 004</u>	Oct 07, 2008
<u>AB</u>	CHARTWELL	<u>50MG</u>	<u>A076665 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076665 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076665 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076665 004</u>	Jul 29, 2004
<u>AB</u>	DR REDDYS LABS INC	<u>50MG</u>	<u>A076658 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076658 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076658 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076658 004</u>	Jul 29, 2004
<u>AB</u>	GLENMARK GENERICS	<u>50MG</u>	<u>A077253 001</u>	Jan 25, 2006
<u>AB</u>		<u>100MG</u>	<u>A077253 002</u>	Jan 25, 2006
<u>AB</u>		<u>150MG</u>	<u>A077253 003</u>	Jan 25, 2006
<u>AB</u>		<u>200MG</u>	<u>A077253 004</u>	Jan 25, 2006
<u>AB</u>	TARO	<u>50MG</u>	<u>A076507 001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076507 002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076507 003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076507 004</u>	Jul 29, 2004
<u>AB</u>	THINQ PHARM-CRO PVT	<u>50MG</u>	<u>A076957 001</u>	Sep 28, 2005
<u>AB</u>		<u>100MG</u>	<u>A076957 002</u>	Sep 28, 2005
<u>AB</u>		<u>150MG</u>	<u>A076957 004</u>	Feb 27, 2017
<u>AB</u>		<u>200MG</u>	<u>A076957 003</u>	Sep 28, 2005
<u>AB</u>	ZYDUS PHARMS	<u>50MG</u>	<u>A208963 001</u>	Feb 16, 2017
<u>AB</u>		<u>100MG</u>	<u>A208963 002</u>	Feb 16, 2017
<u>AB</u>		<u>150MG</u>	<u>A208963 003</u>	Feb 16, 2017
<u>AB</u>		<u>200MG</u>	<u>A208963 004</u>	Feb 16, 2017

FLUCYTOSINE

CAPSULE; ORAL

ANCOBON

<u>AB</u>	+	BAUSCH	<u>250MG</u>	<u>N017001 001</u>
<u>AB</u>	+	!	<u>500MG</u>	<u>N017001 002</u>

FLUCYTOSINE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A213665 001</u>	May 01, 2020
<u>AB</u>		<u>500MG</u>	<u>A213665 002</u>	May 01, 2020
<u>AB</u>	NOVEL LABS INC	<u>250MG</u>	<u>A204652 001</u>	Jul 07, 2017
<u>AB</u>		<u>500MG</u>	<u>A204652 002</u>	Jul 07, 2017
<u>AB</u>	SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A201566 001</u>	Jun 28, 2011
<u>AB</u>		<u>500MG</u>	<u>A201566 002</u>	Jun 28, 2011
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A212632 001</u>	Apr 17, 2020
<u>AB</u>		<u>500MG</u>	<u>A212632 002</u>	Apr 17, 2020

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARABINE PHOSPHATE

<u>AP</u>	!	ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078610 001</u>	Feb 11, 2009
<u>AP</u>		FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>A078544 001</u>	Oct 15, 2007
<u>AP</u>		HIKMA	<u>50MG/VIAL</u>	<u>A076349 001</u>	Aug 28, 2003
<u>AP1</u>		AREVA PHARMS	<u>50MG/2ML (25MG/ML)</u>	<u>A090724 001</u>	Sep 27, 2010
<u>AP1</u>		FRESENIUS KABI USA	<u>50MG/2ML (25MG/ML)</u>	<u>A078393 001</u>	Oct 15, 2007
<u>AP1</u>		SAGENT PHARMS INC	<u>50MG/2ML (25MG/ML)</u>	<u>A076661 001</u>	Apr 28, 2004

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

<u>AP</u>		3D IMAGING DRUG	<u>20-300mCi/ML</u>	<u>A203778 001</u>	Oct 30, 2015
<u>AP</u>		BIOMEDCL RES FDN	<u>20-300mCi/ML</u>	<u>A203710 001</u>	May 01, 2015
<u>AP</u>			<u>20-300mCi/ML</u>	<u>A203837 001</u>	May 01, 2015
<u>AP</u>		BRIGHAM WOMENS	<u>20-300mCi/ML</u>	<u>A203816 001</u>	Oct 30, 2014
<u>AP</u>		CARDINAL HEALTH 414	<u>20-300mCi/ML</u>	<u>A203603 001</u>	Nov 13, 2015
<u>AP</u>			<u>20-500mCi/ML</u>	<u>A203603 002</u>	Sep 27, 2018
<u>AP</u>		CHILDRENS HOSP MI	<u>20-300mCi/ML</u>	<u>A204385 001</u>	Oct 29, 2014
<u>AP</u>		DECATUR	<u>20-300mCi/ML</u>	<u>A204463 001</u>	Oct 21, 2014
<u>AP</u>	+	FEINSTEIN	<u>20-400mCi/ML</u>	<u>N021870 002</u>	Nov 21, 2008
<u>AP</u>		ISOLOGIC INNOVATIVE	<u>20-300mCi/ML</u>	<u>A204525 001</u>	Oct 29, 2014
<u>AP</u>		JUBILANT DRAXIMAGE	<u>20-300mCi/ML</u>	<u>A203920 001</u>	Jun 23, 2015

PRESCRIPTION DRUG PRODUCT LIST

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

<u>AP</u>	KETTERING MEDCTR	<u>4-40mCi/ML</u>	<u>A204759 001</u>	Oct 27, 2015
<u>AP</u>	KREITCHMAN PET CTR	<u>10-100mCi/ML</u>	<u>A203942 001</u>	Apr 11, 2016
<u>AP</u>	MA GENERAL HOSP	<u>20-300mCi/ML</u>	<u>A204333 001</u>	Sep 25, 2014
<u>AP</u>	MCPRF	<u>20-240mCi/ML</u>	<u>A203612 001</u>	Aug 05, 2013
<u>AP</u>	MEM SLOAN-KETTERING	<u>20-300mCi/ML</u>	<u>A208679 001</u>	Dec 08, 2016
<u>AP</u>	METHODIST HOSP RES	<u>20-300mCi/ML</u>	<u>A203904 001</u>	Apr 23, 2015
<u>AP</u>	MIPS CRF	<u>20-300mCi/ML</u>	<u>A204472 001</u>	Sep 11, 2015
<u>AP</u>	NCM USA BRONX LLC	<u>20-300mCi/ML</u>	<u>A204512 001</u>	Jan 07, 2015
<u>AP</u>	! PETNET	<u>20-200mCi/ML</u>	<u>A079086 001</u>	Feb 25, 2011
<u>AP</u>	PHARMALOGIC	<u>20-300mCi/ML</u>	<u>A204264 001</u>	Dec 18, 2014
<u>AP</u>	PHARMALOGIC HLDGS	<u>20-200mCi/ML</u>	<u>A203664 001</u>	Feb 04, 2014
<u>AP</u>	PRECISION NUCLEAR	<u>20-500mCi/ML</u>	<u>A204546 001</u>	Apr 07, 2015
<u>AP</u>	! QUEEN HAMAMATSU PET	<u>10-100mCi/ML</u>	<u>A203771 001</u>	Aug 31, 2015
<u>AP</u>	SOFIE	<u>20-300mCi/ML</u>	<u>A203591 001</u>	Aug 31, 2015
<u>AP</u>	! TRUSTEES UNIV PA	<u>20-500mCi/ML</u>	<u>A203665 001</u>	Feb 14, 2013
<u>AP</u>	! UCLA BIOMEDICAL	<u>20-200mCi/ML</u>	<u>A203801 001</u>	Oct 29, 2014
<u>AP</u>	! UCSF RODIOPHARM	<u>4-40mCi/ML</u>	<u>A203811 001</u>	Jun 27, 2013
<u>AP</u>	UCSF RODIOPHARM	<u>20-300mCi/ML</u>	<u>A203902 001</u>	May 09, 2014
<u>AP</u>	UIHC PET IMAGING	<u>20-300mCi/ML</u>	<u>A203990 001</u>	Aug 06, 2014
<u>AP</u>	UNIV MICHIGAN	<u>20-300mCi/ML</u>	<u>A204531 001</u>	Jul 17, 2015
<u>AP</u>	UNIV SOUTHERN CA	<u>20-300mCi/ML</u>	<u>A209341 001</u>	Dec 16, 2020
<u>AP</u>	UNIV TX MD ANDERSON	<u>20-300mCi/ML</u>	<u>A203246 002</u>	Jan 13, 2014
<u>AP</u>	UNIV UTAH CYCLOTRON	<u>20-300mCi/ML</u>	<u>A204498 001</u>	Jun 23, 2015
<u>AP</u>	WISCONSIN	<u>20-500mCi/ML</u>	<u>A203709 001</u>	Oct 23, 2013
<u>AP</u>	WUSM CYCLOTRON	<u>20-300mCi/ML</u>	<u>A203935 001</u>	Feb 05, 2014
	NORTHLAND	4-500mCi/ML	A203994	001 Feb 04, 2015
	NUKEMED	4-500mCi/ML	A203911	001 Apr 22, 2015
	UNIV TX MD ANDERSON	20-150mCi/ML	A203246	001 Jan 13, 2014

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLUDROCORTISONE ACETATE

<u>AB</u>	BARR	<u>0.1MG</u>	<u>A040425 001</u>	Jan 21, 2003
<u>AB</u>	! IMPAX LABS	<u>0.1MG</u>	<u>A040431 001</u>	Mar 18, 2002
<u>AB</u>	NOVITIUM PHARMA	<u>0.1MG</u>	<u>A215279 001</u>	May 31, 2022

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<u>AP</u>	BAXTER HLTHCARE CORP	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076755 002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076755 001</u>	Oct 12, 2004
<u>AP</u>	FRESENIUS KABI USA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076955 002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076955 001</u>	Oct 12, 2004
<u>AP</u>	HIKMA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076256 002</u>	Oct 12, 2004
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076787 002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076256 001</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076787 001</u>	Oct 12, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078527 001</u>	Mar 23, 2009
<u>AP</u>	! RISING	<u>1MG/10ML (0.1MG/ML)</u>	<u>A078527 002</u>	Mar 23, 2009
<u>AP</u>	SAGENT PHARMS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078595 001</u>	May 13, 2008
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A090584 001</u>	Aug 28, 2012
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A090584 002</u>	Aug 28, 2012
<u>AP</u>	SANDOZ	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A077071 001</u>	May 03, 2005
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A077071 002</u>	May 03, 2005

FLUNISOLIDE

SPRAY, METERED; NASAL

FLUNISOLIDE

<u>AB</u>	! BAUSCH	<u>0.025MG/SPRAY</u>	<u>A074805 001</u>	Feb 20, 2002
<u>AB</u>	RICONPHARMA LLC	<u>0.025MG/SPRAY</u>	<u>A207802 001</u>	Jun 16, 2022
<u>AB</u>	RISING	<u>0.025MG/SPRAY</u>	<u>A077704 001</u>	Aug 03, 2006

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	COSETTE	<u>0.01%</u>	<u>A089526 001</u>	Jul 26, 1988
<u>AT</u>		<u>0.025%</u>	<u>A210747 001</u>	Nov 05, 2018
<u>AT</u>	FOUGERA PHARMS INC	<u>0.01%</u>	<u>A088170 001</u>	Dec 16, 1982
<u>AT</u>		<u>0.025%</u>	<u>A088169 001</u>	Dec 16, 1982
<u>AT</u>	TARO	<u>0.025%</u>	<u>A087104 001</u>	Apr 27, 1982

PRESCRIPTION DRUG PRODUCT LIST

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

SYNALAR

AT	+	MEDIMETRIKS PHARMS	0.01%	N012787	004	
AT	+		0.025%	N012787	002	
AT	+		0.025%	N012787	005	

IMPLANT; INTRAVITREAL

ILUVIEN

+	ALIMERA SCIENCES INC	0.19MG	N201923	001	Sep 26, 2014
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RETISERT

+	BAUSCH AND LOMB	0.59MG	N021737	001	Apr 08, 2005
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YUTIQ

+	ALIMERA SCIENCES INC	0.18MG	N210331	001	Oct 12, 2018
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OIL; TOPICAL

DERMA-SMOOTH/FS

AT	+	HILL DERMAC	0.01%	N019452	001	Feb 03, 1988
AT	+		0.01%	N019452	002	Nov 09, 2005

FLUCINOLONE ACETONIDE

AT		GLENMARK PHARMS LTD	0.01%	A210556	001	Oct 25, 2018
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FLUOCINOLONE ACETONIDE

AT		LYNE	0.01%	A090982	001	Apr 25, 2016
AT			0.01%	A203377	001	Apr 25, 2016
AT		NOVEL LABS INC	0.01%	A207345	001	Jul 31, 2023
AT			0.01%	A207347	001	Aug 07, 2023
AT		PADAGIS ISRAEL	0.01%	A202847	001	Aug 09, 2013
AT			0.01%	A202848	001	Aug 09, 2013
AT		QUAGEN	0.01%	A212760	001	Apr 02, 2021
AT			0.01%	A212761	001	Apr 02, 2021
AT		TARO	0.01%	A202368	001	May 19, 2016
AT			0.01%	A209336	001	May 19, 2016

FLUOCINONIDE ACETONIDE

AT		GLENMARK PHARMS LTD	0.01%	A210539	001	Oct 26, 2018
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OIL/DROPS; OTIC

DERMOTIC

AT	+	HILL DERMAC	0.01%	N019452	003	Nov 09, 2005
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FLAC

AT		PATRIN	0.01%	A210736	001	Apr 11, 2018
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FLUOCINOLONE ACETONIDE

AT		LYNE	0.01%	A203378	001	Apr 25, 2016
AT		PADAGIS ISRAEL	0.01%	A202849	001	Jul 17, 2017
AT		QUAGEN	0.01%	A212762	001	Apr 02, 2021
AT		TASMAN PHARMA	0.01%	A213264	001	Feb 05, 2021

FLUOCINONIDE ACETONIDE

AT		GLENMARK PHARMS LTD	0.01%	A211815	001	Dec 14, 2018
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OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

AT		COSETTE	0.025%	A089524	001	Jul 26, 1988
AT		FOUGERA PHARMS INC	0.025%	A088168	001	Dec 16, 1982
AT		TARO	0.025%	A040041	001	Sep 15, 1994

SYNALAR

AT	+	MEDIMETRIKS PHARMS	0.025%	N013960	001	
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SHAMPOO; TOPICAL

CAPEX

+	GALDERMA LABS LP	0.01%	N020001	001	Aug 27, 1990
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SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

AT		CHARTWELL RX	0.01%	A209596	001	Dec 26, 2017
AT		ENCUBE ETHICALS	0.01%	A209913	001	Feb 13, 2019
AT		FOUGERA PHARMS INC	0.01%	A088167	001	Dec 16, 1982
AT		LUPIN	0.01%	A206422	001	Sep 02, 2015
AT		TARO	0.01%	A089124	001	Sep 11, 1985

SYNALAR

AT	+	MEDIMETRIKS PHARMS	0.01%	N015296	001	
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FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN

CREAM; TOPICAL

TRI-LUMA

+	GALDERMA LABS LP	0.01%; 4%; 0.05%	N021112	001	Jan 18, 2002
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PRESCRIPTION DRUG PRODUCT LIST

FLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-SYNALAR

! MEDIMETRIKS PHARMS 0.025%;EQ 3.5MG BASE/GM A060700 001

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

AB	AMNEAL	<u>0.1%</u>	<u>A211111</u>	<u>001</u>	Jun 04, 2018
AB	CADILA	<u>0.1%</u>	<u>A208989</u>	<u>001</u>	Feb 10, 2020
AB	FOUGERA PHARMS INC	<u>0.1%</u>	<u>A200735</u>	<u>001</u>	Jul 14, 2014
AB	GLENMARK GENERICS	<u>0.1%</u>	<u>A091282</u>	<u>001</u>	Jul 14, 2014
AB	PADAGIS ISRAEL	<u>0.1%</u>	<u>A090256</u>	<u>001</u>	Jan 14, 2014
AB	TARO	<u>0.1%</u>	<u>A200734</u>	<u>001</u>	Jul 14, 2014

VANOS

AB	+! BAUSCH	<u>0.1%</u>	<u>N021758</u>	<u>001</u>	Feb 11, 2005
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FLUOCINONIDE

AB1	! AMNEAL	<u>0.05%</u>	<u>A210554</u>	<u>001</u>	Aug 21, 2018
AB1	COSETTE	<u>0.05%</u>	<u>A073085</u>	<u>001</u>	Feb 14, 1992
AB1	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A073030</u>	<u>001</u>	Oct 17, 1994
AB1	TARO	<u>0.05%</u>	<u>A071500</u>	<u>001</u>	Jun 10, 1987
AB1	TEVA	<u>0.05%</u>	<u>A072488</u>	<u>001</u>	Feb 06, 1989

FLUOCINONIDE EMULSIFIED BASE

AB2	COSETTE	<u>0.05%</u>	<u>A074204</u>	<u>001</u>	Jun 13, 1995
AB2	FOUGERA PHARMS	<u>0.05%</u>	<u>A076586</u>	<u>001</u>	Jun 23, 2004
AB2	! TARO	<u>0.05%</u>	<u>A072494</u>	<u>001</u>	Jan 19, 1989
AB2	TEVA	<u>0.05%</u>	<u>A072490</u>	<u>001</u>	Feb 07, 1989

GEL; TOPICAL

FLUOCINONIDE

AB	+ ALVOGEN	<u>0.05%</u>	<u>N017373</u>	<u>001</u>	
AB	COSETTE	<u>0.05%</u>	<u>A072537</u>	<u>001</u>	Feb 07, 1989
AB	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072933</u>	<u>001</u>	Dec 30, 1994
AB	! TARO	<u>0.05%</u>	<u>A074935</u>	<u>001</u>	Jul 29, 1997

OINTMENT; TOPICAL

FLUOCINONIDE

AB	CHARTWELL RX	<u>0.05%</u>	<u>A207538</u>	<u>001</u>	Jul 31, 2017
AB	FOUGERA PHARMS	<u>0.05%</u>	<u>A074905</u>	<u>001</u>	Aug 26, 1997
AB	! TARO	<u>0.05%</u>	<u>A075008</u>	<u>001</u>	Jun 30, 1999
AB	TEVA	<u>0.05%</u>	<u>A073481</u>	<u>001</u>	Dec 27, 1991
AB	XIROMED	<u>0.05%</u>	<u>A212976</u>	<u>001</u>	Nov 26, 2019

LIDEX

AB	+ ALVOGEN	<u>0.05%</u>	<u>N016909</u>	<u>002</u>	
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SOLUTION; TOPICAL

FLUOCINONIDE

AT	CHARTWELL RX	<u>0.05%</u>	<u>A209118</u>	<u>001</u>	Apr 23, 2018
AT	COSETTE	<u>0.05%</u>	<u>A071535</u>	<u>001</u>	Dec 02, 1988
AT	ENCUBE ETHICALS	<u>0.05%</u>	<u>A209699</u>	<u>001</u>	Nov 29, 2018
AT	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072934</u>	<u>001</u>	Feb 27, 1995
AT	MACLEODS PHARMS LTD	<u>0.05%</u>	<u>A209283</u>	<u>001</u>	Apr 23, 2018
AT	NOVEL LABS INC	<u>0.05%</u>	<u>A206003</u>	<u>001</u>	Jul 21, 2017
AT	! TARO	<u>0.05%</u>	<u>A074799</u>	<u>001</u>	Dec 31, 1996
AT	ZYDUS PHARMS	<u>0.05%</u>	<u>A208948</u>	<u>001</u>	Jul 17, 2018

FLUORESCHEIN SODIUM

INJECTABLE; INTRAVENOUS

AK-FLUOR 10%

AP	+! LONG GROVE PHARMS	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N022186</u>	<u>001</u>	Aug 08, 2008
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AK-FLUOR 25%

AP	+! LONG GROVE PHARMS	<u>EQ 500MG BASE/2ML (EQ 250MG BASE/ML)</u>	<u>N022186</u>	<u>002</u>	Aug 08, 2008
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FLUORESCHEIN SODIUM

AP	NEXUS PHARMS	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>A215709</u>	<u>001</u>	Sep 25, 2023
AP		<u>EQ 500MG BASE/2ML (EQ 250MG BASE/ML)</u>	<u>A215709</u>	<u>002</u>	Sep 25, 2023

FLUORESCITE

AP	+! ALCON LABS INC	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N021980</u>	<u>001</u>	Mar 28, 2006
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FLUORODOPA F-18

SOLUTION; INTRAVENOUS

FLUORODOPA F18

+! FEINSTEIN 0.42-8.33mCi/ML N200655 001 Oct 10, 2019

PRESCRIPTION DRUG PRODUCT LIST

FLUOROESTRADIOL F-18

SOLUTION; INTRAVENOUS

CERIANNA

+! GE HEALTHCARE 50ML (4-100mCi/ML) N212155 001 May 20, 2020

FLUOROMETHOLONE

SUSPENSION/DROPS; OPHTHALMIC

FLUOROMETHOLONE**AB** AMNEAL 0.1% **A216348 001** Jan 09, 2024FML**AB** +! ABBVIE 0.1% **N016851 002** Jul 28, 1982

FML FORTE

+! ABBVIE 0.25% N019216 001 Apr 23, 1986

FLUOROMETHOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

FLAREX

+! HARROW EYE 0.1% N019079 001 Feb 11, 1986

FLUOROURACIL

CREAM; TOPICAL

EFUDEX**AB** +! BAUSCH 5% **N016831 003**FLUOROURACIL**AB** ACCORD HLTHCARE 5% **A214845 001** Oct 07, 2021**AB** DR REDDYS LABS SA 5% **A077524 001** Apr 11, 2008**AB** TARO 5% **A090368 001** Mar 05, 2010

CARAC

+! VALEANT PHARMS NORTH 0.5% N020985 001 Oct 27, 2000

TOLAK

+! HILL DERMACEUTICALS 4% N022259 001 Sep 18, 2015

INJECTABLE; INJECTION

FLUOROURACIL**AP** ! ACCORD HLTHCARE 500MG/10ML (50MG/ML) **A040743 002** Apr 26, 2007**AP** ! 1GM/20ML (50MG/ML) **A040743 001** Apr 26, 2007**AP** ! 2.5GM/50ML (50MG/ML) **A040798 002** Apr 26, 2007**AP** ! 5GM/100ML (50MG/ML) **A040798 001** Apr 26, 2007**AP** ALEMBIC 2.5GM/50ML (50MG/ML) **A217295 001** Mar 03, 2023**AP** 5GM/100ML (50MG/ML) **A217676 001** Oct 18, 2023**AP** FRESENIUS KABI USA 500MG/10ML (50MG/ML) **A040279 002** Sep 30, 1998**AP** 1GM/20ML (50MG/ML) **A040279 001** Sep 30, 1998**AP** 2.5GM/50ML (50MG/ML) **A040278 001** Sep 30, 1998**AP** 5GM/100ML (50MG/ML) **A040278 002** Sep 30, 1998**AP** GLAND PHARMA LTD 500MG/10ML (50MG/ML) **A210123 001** Oct 27, 2017**AP** 1GM/20ML (50MG/ML) **A210123 002** Oct 27, 2017**AP** 2.5GM/50ML (50MG/ML) **A210124 001** Dec 26, 2017**AP** 5GM/100ML (50MG/ML) **A210124 002** Dec 26, 2017**AP** SAGENT PHARMS INC 500MG/10ML (50MG/ML) **A203608 001** May 11, 2017**AP** 1GM/20ML (50MG/ML) **A203608 002** May 11, 2017**AP** 2.5GM/50ML (50MG/ML) **A203609 001** Feb 17, 2016**AP** 5GM/100ML (50MG/ML) **A203609 002** Feb 17, 2016

SOLUTION; TOPICAL

EFUDEX**AT** +! BAUSCH 2% **N016831 001**FLUOROURACIL**AT** ENCUBE 5% **A215612 001** Nov 02, 2023**AT** TARO 2% **A076526 001** Nov 05, 2003**AT** ! 5% **A076526 002** Nov 05, 2003FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE HYDROCHLORIDE**AB** ALEMBIC PHARMS LTD EQ 40MG BASE **A090223 003** Mar 19, 2009**AB** APNAR PHARMA LP EQ 40MG BASE **A075049 003** Jan 29, 2002**AB** AUROBINDO PHARMA EQ 40MG BASE **A078619 003** Jan 31, 2008**AB** CADILA PHARMS LTD EQ 40MG BASE **A206993 003** May 23, 2019**AB** HERITAGE PHARMS EQ 40MG BASE **A201336 003** Oct 01, 2012**AB** IVAX SUB TEVA EQ 40MG BASE **A075245 003** Sep 28, 2004

PHARMS

AB MARKSANS PHARMA EQ 40MG BASE **A075465 003** Aug 02, 2001**AB** MICRO LABS EQ 40MG BASE **A216232 003** Mar 29, 2022**AB** SCIEGEN PHARMS INC EQ 40MG BASE **A204597 003** Mar 16, 2015**AB** SUN PHARM INDS LTD EQ 40MG BASE **A076990 001** Dec 13, 2004**AB** TEVA EQ 40MG BASE **A075452 003** Jan 29, 2002

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

PROZAC

<u>AB</u>	<u>+!</u>	ELI LILLY AND CO	<u>EQ 40MG BASE</u>	<u>N018936 003</u>	Jun 15, 1999
<u>FLUOXETINE HYDROCHLORIDE</u>					
<u>AB1</u>		ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A090223 001</u>	Mar 19, 2009
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A090223 002</u>	Mar 19, 2009
<u>AB1</u>		APNAR PHARMA LP	<u>EQ 10MG BASE</u>	<u>A075049 001</u>	Aug 02, 2001
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075049 002</u>	Jan 29, 2002
<u>AB1</u>		AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A078619 001</u>	Jan 31, 2008
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A078619 002</u>	Jan 31, 2008
<u>AB1</u>		CADILA PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A206993 001</u>	May 23, 2019
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A206993 002</u>	May 23, 2019
<u>AB1</u>		HERITAGE PHARMS	<u>EQ 10MG BASE</u>	<u>A201336 001</u>	Oct 01, 2012
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A201336 002</u>	Oct 01, 2012
<u>AB1</u>		IVAX SUB TEVA PHARMS	<u>EQ 10MG BASE</u>	<u>A075245 002</u>	Jan 31, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075245 001</u>	Jan 31, 2002
<u>AB1</u>		LANDELA PHARM	<u>EQ 10MG BASE</u>	<u>A075464 001</u>	Jan 30, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075464 002</u>	Jan 30, 2002
<u>AB1</u>		MARKSANS PHARMA	<u>EQ 10MG BASE</u>	<u>A075465 001</u>	Jan 29, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075465 002</u>	Jan 29, 2002
<u>AB1</u>		MICRO LABS	<u>EQ 10MG BASE</u>	<u>A216232 001</u>	Mar 29, 2022
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A216232 002</u>	Mar 29, 2022
<u>AB1</u>		SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A204597 001</u>	Mar 16, 2015
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A204597 002</u>	Mar 16, 2015
<u>AB1</u>		TEVA	<u>EQ 10MG BASE</u>	<u>A075452 001</u>	Jan 29, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A075452 002</u>	Jan 29, 2002
<u>AB1</u>		TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A076001 001</u>	Jan 29, 2002
<u>AB1</u>			<u>EQ 20MG BASE</u>	<u>A076001 002</u>	Jan 29, 2002

PROZAC

<u>AB1</u>	<u>+</u>	ELI LILLY AND CO	<u>EQ 10MG BASE</u>	<u>N018936 006</u>	Dec 23, 1992
<u>AB1</u>	<u>+</u>		<u>EQ 20MG BASE</u>	<u>N018936 001</u>	Dec 29, 1987

CAPSULE, DELAYED REL PELLETS;ORAL

FLUOXETINE HYDROCHLORIDE

! DR REDDYS LABS LTD EQ 90MG BASE

A078572 001 Mar 22, 2010

SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

<u>AA</u>		APTAPHARMA INC	<u>EQ 20MG BASE/5ML</u>	<u>A216953 001</u>	Nov 15, 2022
<u>AA</u>		AUROBINDO PHARMA	<u>EQ 20MG BASE/5ML</u>	<u>A079209 001</u>	Mar 20, 2009
<u>AA</u>		LANNETT CO INC	<u>EQ 20MG BASE/5ML</u>	<u>A077849 001</u>	Feb 09, 2007
<u>AA</u>		NOSTRUM LABS INC	<u>EQ 20MG BASE/5ML</u>	<u>A075292 001</u>	Feb 07, 2002
<u>AA</u>		NOVITIUM PHARMA	<u>EQ 20MG BASE/5ML</u>	<u>A216448 001</u>	Nov 09, 2022
<u>AA</u>	<u>!</u>	PHARM ASSOC	<u>EQ 20MG BASE/5ML</u>	<u>A076015 001</u>	Jan 30, 2002
<u>AA</u>		TEVA	<u>EQ 20MG BASE/5ML</u>	<u>A075506 001</u>	Aug 02, 2001

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC	<u>EQ 10MG BASE</u>	<u>A208698 001</u>	Apr 05, 2017
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A208698 002</u>	Apr 05, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A213286 001</u>	Apr 08, 2020
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A213286 002</u>	Apr 08, 2020
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A213265 001</u>	Jun 10, 2020
<u>AB</u>		DR REDDYS	<u>EQ 10MG BASE</u>	<u>A076006 001</u>	Jan 30, 2002
<u>AB</u>	<u>!</u>		<u>EQ 20MG BASE</u>	<u>A076006 002</u>	Apr 23, 2018
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A211721 001</u>	Jan 25, 2019
<u>AB</u>		INVENTIA HLTHCARE	<u>EQ 60MG BASE</u>	<u>A209695 001</u>	Nov 20, 2017
<u>AB</u>		LUPIN LTD	<u>EQ 10MG BASE</u>	<u>A211653 001</u>	Apr 15, 2019
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A211653 002</u>	Apr 15, 2019
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A211632 001</u>	Feb 08, 2019
<u>AB</u>		PAR FORM	<u>EQ 10MG BASE</u>	<u>A203836 001</u>	Aug 19, 2016
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203836 002</u>	Aug 19, 2016
<u>AB</u>		PAR PHARM INC	<u>EQ 60MG BASE</u>	<u>A209419 001</u>	Nov 16, 2017
<u>AB</u>		SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A210935 001</u>	Mar 20, 2019
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A210935 002</u>	Mar 20, 2019
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A211282 001</u>	Jan 10, 2019
<u>AB</u>		TARO	<u>EQ 60MG BASE</u>	<u>A211477 001</u>	Nov 21, 2018
<u>AB</u>		TEVA	<u>EQ 10MG BASE</u>	<u>A075872 001</u>	Jan 29, 2002
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A075872 002</u>	Jan 04, 2019
<u>AB</u>		TEVA PHARMS USA	<u>EQ 60MG BASE</u>	<u>A211051 001</u>	Dec 03, 2018
<u>AB</u>	<u>+!</u>	TWI PHARMS	<u>EQ 60MG BASE</u>	<u>N202133 001</u>	Oct 06, 2011
<u>AB</u>		UPSHER SMITH LABS	<u>EQ 10MG BASE</u>	<u>A211696 001</u>	Jan 30, 2019
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A211696 002</u>	Jan 30, 2019

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE

TABLET; ORAL

FLUOXETINE HYDROCHLORIDE

TORRENT

EQ 10MG BASE

A206937 001 Oct 21, 2016

!

EQ 20MG BASE

A206937 002 Oct 21, 2016

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA LLC	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A078901 005</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A078901 001</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A078901 003</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A078901 002</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A078901 004</u>	Nov 16, 2012
<u>AB</u>	PAR PHARM	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A077742 001</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077742 002</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077742 003</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077742 004</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077742 005</u>	Nov 02, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A202074 001</u>	Mar 25, 2013
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077528 001</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077528 002</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077528 003</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077528 004</u>	Jun 19, 2012
<u>SYMBYAX</u>				
<u>AB</u>	+ LILLY	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>N021520 001</u>	Apr 09, 2007
<u>AB</u>	+	<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>N021520 002</u>	Dec 24, 2003
<u>AB</u>	+	<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>N021520 004</u>	Dec 24, 2003
<u>AB</u>	+!	<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>N021520 003</u>	Dec 24, 2003
<u>AB</u>	+	<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>N021520 005</u>	Dec 24, 2003

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

<u>AO</u>	EUGIA PHARMA	<u>25MG/ML</u>	<u>A207739 001</u>	Oct 17, 2017
<u>AO</u>	! FRESENIUS KABI USA	<u>25MG/ML</u>	<u>A071413 001</u>	Jul 14, 1987
<u>AO</u>	GLAND PHARMA LTD	<u>25MG/ML</u>	<u>A215509 001</u>	Mar 30, 2023
<u>AO</u>	HIKMA	<u>25MG/ML</u>	<u>A074531 001</u>	Aug 30, 1996
<u>AO</u>	MSN	<u>25MG/ML</u>	<u>A215365 001</u>	Nov 02, 2023
<u>AO</u>	MYLAN LABS LTD	<u>25MG/ML</u>	<u>A075918 001</u>	Aug 17, 2001
<u>AO</u>	PAR STERILE PRODUCTS	<u>25MG/ML</u>	<u>A203732 001</u>	Jul 03, 2014

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

5MG/ML

A074725 001 Sep 16, 1996

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

2.5MG/5ML

A040146 001 Aug 21, 1996

INJECTABLE; INJECTION

FLUPHENAZINE HYDROCHLORIDE

! FRESENIUS KABI USA

2.5MG/ML

A089556 001 Apr 16, 1987

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>1MG</u>	<u>A217410 001</u>	Jan 05, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A217410 002</u>	Jan 05, 2023
<u>AB</u>		<u>5MG</u>	<u>A217410 003</u>	Jan 05, 2023
<u>AB</u>		<u>10MG</u>	<u>A217410 004</u>	Jan 05, 2023
<u>AB</u>	AMNEAL	<u>1MG</u>	<u>A213647 001</u>	Jul 09, 2020
<u>AB</u>		<u>2.5MG</u>	<u>A213647 002</u>	Jul 09, 2020
<u>AB</u>		<u>5MG</u>	<u>A213647 003</u>	Jul 09, 2020
<u>AB</u>		<u>10MG</u>	<u>A213647 004</u>	Jul 09, 2020
<u>AB</u>	APOTEX	<u>1MG</u>	<u>A216649 001</u>	Jul 15, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A216649 002</u>	Jul 15, 2022
<u>AB</u>		<u>5MG</u>	<u>A216649 003</u>	Jul 15, 2022
<u>AB</u>		<u>10MG</u>	<u>A216649 004</u>	Jul 15, 2022
<u>AB</u>	AUROBINDO PHARMA LTD	<u>1MG</u>	<u>A218055 001</u>	Aug 18, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A218055 002</u>	Aug 18, 2023
<u>AB</u>		<u>5MG</u>	<u>A218055 003</u>	Aug 18, 2023
<u>AB</u>		<u>10MG</u>	<u>A218055 004</u>	Aug 18, 2023
<u>AB</u>	CHARTWELL RX	<u>1MG</u>	<u>A215141 001</u>	Oct 20, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A215141 002</u>	Oct 20, 2021

PRESCRIPTION DRUG PRODUCT LIST

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AB</u>		<u>5MG</u>	<u>A215141 003</u>	Oct 20, 2021
<u>AB</u>		<u>10MG</u>	<u>A215141 004</u>	Oct 20, 2021
<u>AB</u>	DR REDDYS	<u>1MG</u>	<u>A214534 001</u>	Jan 07, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214534 002</u>	Jan 07, 2021
<u>AB</u>		<u>5MG</u>	<u>A214534 003</u>	Jan 07, 2021
<u>AB</u>		<u>10MG</u>	<u>A214534 004</u>	Jan 07, 2021
<u>AB</u>	GLENMARK PHARMS LTD	<u>1MG</u>	<u>A216350 001</u>	Nov 06, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A216350 002</u>	Nov 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A216350 003</u>	Nov 06, 2023
<u>AB</u>		<u>10MG</u>	<u>A216350 004</u>	Nov 06, 2023
<u>AB</u>	LANNETT CO INC	<u>1MG</u>	<u>A089743 002</u>	Aug 25, 1988
<u>AB</u>		<u>2.5MG</u>	<u>A089743 003</u>	Aug 25, 1988
<u>AB</u>	!	<u>5MG</u>	<u>A089743 004</u>	Aug 25, 1988
<u>AB</u>		<u>10MG</u>	<u>A089743 001</u>	Aug 25, 1988
<u>AB</u>	MSN	<u>1MG</u>	<u>A217189 001</u>	Sep 11, 2023
<u>AB</u>		<u>2.5MG</u>	<u>A217189 002</u>	Sep 11, 2023
<u>AB</u>		<u>5MG</u>	<u>A217189 003</u>	Sep 11, 2023
<u>AB</u>		<u>10MG</u>	<u>A217189 004</u>	Sep 11, 2023
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A214674 001</u>	Mar 01, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214674 002</u>	Mar 01, 2021
<u>AB</u>		<u>5MG</u>	<u>A214674 003</u>	Mar 01, 2021
<u>AB</u>		<u>10MG</u>	<u>A214674 004</u>	Mar 01, 2021
<u>AB</u>	SANDOZ	<u>1MG</u>	<u>A216891 001</u>	Nov 15, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A216891 002</u>	Nov 15, 2022
<u>AB</u>		<u>5MG</u>	<u>A216891 003</u>	Nov 15, 2022
<u>AB</u>		<u>10MG</u>	<u>A216891 004</u>	Nov 15, 2022
<u>AB</u>	TARO	<u>1MG</u>	<u>A215674 001</u>	Apr 14, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A215674 002</u>	Apr 14, 2022
<u>AB</u>		<u>5MG</u>	<u>A215674 003</u>	Apr 14, 2022
<u>AB</u>		<u>10MG</u>	<u>A215674 004</u>	Apr 14, 2022
<u>AB</u>	TWI PHARMS	<u>1MG</u>	<u>A215848 004</u>	Dec 14, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A215848 001</u>	Apr 01, 2022
<u>AB</u>		<u>5MG</u>	<u>A215848 002</u>	Apr 01, 2022
<u>AB</u>		<u>10MG</u>	<u>A215848 003</u>	Apr 01, 2022
<u>AB</u>	UPSHER SMITH LABS	<u>1MG</u>	<u>A213784 001</u>	Oct 24, 2022
<u>AB</u>		<u>2.5MG</u>	<u>A213784 002</u>	Oct 24, 2022
<u>AB</u>		<u>5MG</u>	<u>A213784 003</u>	Oct 24, 2022
<u>AB</u>		<u>10MG</u>	<u>A213784 004</u>	Oct 24, 2022
<u>AB</u>	ZYDUS	<u>1MG</u>	<u>A214552 001</u>	May 27, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A214552 002</u>	May 27, 2021
<u>AB</u>		<u>5MG</u>	<u>A214552 003</u>	May 27, 2021
<u>AB</u>		<u>10MG</u>	<u>A214552 004</u>	May 27, 2021

FLURANDRENOLIDE

LOTION; TOPICAL

FLURANDRENOLIDE

! PADAGIS ISRAEL

0.05%

A207133 001 Aug 30, 2016

OINTMENT; TOPICAL

FLURANDRENOLIDE

! TELIGENT

0.05%

A207851 001 Dec 30, 2016

TAPE; TOPICAL

CORDRAN

+! ALMIRALL

0.004MG/SQ CM

N016455 001

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

<u>AB</u>	CHARTWELL RX	<u>15MG</u>	<u>A072368 001</u>	Mar 30, 1989
<u>AB</u>		<u>30MG</u>	<u>A072369 001</u>	Mar 30, 1989
<u>AB</u>	RISING	<u>15MG</u>	<u>A070345 002</u>	Nov 27, 1985
<u>AB</u>	!	<u>30MG</u>	<u>A070345 001</u>	Nov 27, 1985

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

! TEVA

100MG

A074431 001 May 31, 1995

PRESCRIPTION DRUG PRODUCT LIST

FLURBIPROFEN SODIUM

SOLUTION/DROPS;OPHTHALMIC

FLURBIPROFEN SODIUM

! BAUSCH AND LOMB

0.03%

A074447 001 Jan 04, 1995

FLUTAMIDE

CAPSULE;ORAL

FLUTAMIDE

! WAYLIS THERAP

125MG

A075298 001 Sep 18, 2001

FLUTEMETAMOL F-18

INJECTABLE;INTRAVENOUS

VIZAMYL

+! GE HEALTHCARE

121.5mCi/30ML (4.05mCi/ML)

N203137 002 Oct 25, 2013

FLUTICASON E FUROATE

POWDER;INHALATION

ARNUITY ELLIPTA

+! GLAXOSMITHKLINE

0.05MG/INH

N205625 003 May 17, 2018

+!

0.1MG/INH

N205625 001 Aug 20, 2014

+!

0.2MG/INH

N205625 002 Aug 20, 2014

SPRAY, METERED;NASAL

VERAMYST

+ GLAXOSMITHKLINE

0.0275MG/INH

N022051 001 Apr 27, 2007

FLUTICASON E FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER;INHALATION

TRELEGEY ELLIPTA

+! GLAXOSMITHKLINE

0.1MG/INH;EQ 0.0625MG BASE/INH;EQ
0.025MG BASE/INH

N209482 001 Sep 18, 2017

+!

0.2MG/INH;EQ 0.0625MG BASE/INH;EQ
0.025MG BASE/INH

N209482 002 Sep 09, 2020

FLUTICASON E FUROATE; VILANTEROL TRIFENATATE

POWDER;INHALATION

BREQ ELLIPTA

+ GLAXO GRP LTD

0.05MG/INH;EQ 0.025MG BASE/INH

N204275 003 May 12, 2023

+!

0.1MG/INH;EQ 0.025MG BASE/INH

N204275 001 May 10, 2013

+!

0.2MG/INH;EQ 0.025MG BASE/INH

N204275 002 Apr 30, 2015

FLUTICASON E PROPIONATE

AEROSOL, METERED;INHALATION

FLOVENT HFA

+! GLAXO GRP LTD

0.044MG/INH

N021433 003 May 14, 2004

+!

0.11MG/INH

N021433 002 May 14, 2004

+!

0.22MG/INH

N021433 001 May 14, 2004

CREAM;TOPICAL

FLUTICASON E PROPIONATE**AB** COSETTE**0.05%****A077055 001** Jun 30, 2006**AB** FOUGERA PHARMS**0.05%****A076451 001** May 14, 2004**AB** ! PADAGIS ISRAEL**0.05%****A076793 001** May 14, 2004

LOTION;TOPICAL

FLUTICASON E PROPIONATE**AB** ! GLENMARK GENERICS**0.05%****A090759 001** May 02, 2011**AB** PADAGIS ISRAEL**0.05%****A091553 001** Jul 30, 2013

OINTMENT;TOPICAL

FLUTICASON E PROPIONATE**AB** COSETTE**0.005%****A077168 001** Mar 03, 2006**AB** ! PADAGIS ISRAEL**0.005%****A076668 001** May 14, 2004

POWDER;INHALATION

ARMONAIR DIGIHALER

+ TEVA PHARM

0.055MG/INH

N208798 004 Feb 20, 2020

+

0.113MG/INH

N208798 005 Feb 20, 2020

+!

0.232MG/INH

N208798 006 Feb 20, 2020

FLOVENT DISKUS 100

+! GLAXO GRP LTD

0.1MG/INH

N020833 002 Sep 29, 2000

FLOVENT DISKUS 250

+! GLAXO GRP LTD

0.25MG/INH

N020833 003 Sep 29, 2000

FLOVENT DISKUS 50

+! GLAXO GRP LTD

0.05MG/INH

N020833 001 Sep 29, 2000

SPRAY, METERED;NASAL

FLUTICASON E PROPIONATE**AB** APOTEX INC**0.05MG/SPRAY****A077538 001** Sep 12, 2007**AB** CHARTWELL RX**0.05MG/SPRAY****A078492 001** Jan 09, 2012**AB** ! HIKMA**0.05MG/SPRAY****A076504 001** Feb 22, 2006**AB****0.05MG/SPRAY****A077570 001** Jan 16, 2008

PRESCRIPTION DRUG PRODUCT LIST

FLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

XHANCE

+! OPTINOSE US INC 0.093MG

N209022 001 Sep 18, 2017

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

ADVAIR HFA

+! GLAXO GRP LTD 0.045MG/INH;EQ 0.021MG BASE/INH

N021254 001 Jun 08, 2006

+! 0.115MG/INH;EQ 0.021MG BASE/INH

N021254 002 Jun 08, 2006

+! 0.23MG/INH;EQ 0.021MG BASE/INH

N021254 003 Jun 08, 2006

POWDER; INHALATION

ADVAIR DISKUS 100/50**AB** +! GLAXO GRP LTD **0.1MG/INH;EQ 0.05MG BASE/INH****N021077 001** Aug 24, 2000ADVAIR DISKUS 250/50**AB** +! GLAXO GRP LTD **0.25MG/INH;EQ 0.05MG BASE/INH****N021077 002** Aug 24, 2000ADVAIR DISKUS 500/50**AB** +! GLAXO GRP LTD **0.5MG/INH;EQ 0.05MG BASE/INH****N021077 003** Aug 24, 2000FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE**AB** HIKMA **0.1MG/INH;EQ 0.05MG BASE/INH****A203433 001** Dec 17, 2020**AB** **0.25MG/INH;EQ 0.05MG BASE/INH****A203433 002** Dec 17, 2020**AB** **0.5MG/INH;EQ 0.05MG BASE/INH****A203433 003** Dec 19, 2023**AB** TEVA PHARMS USA **0.1MG/INH;EQ 0.05MG BASE/INH****A213948 001** Dec 13, 2021**AB** **0.25MG/INH;EQ 0.05MG BASE/INH****A213948 002** Dec 13, 2021**AB** **0.5MG/INH;EQ 0.05MG BASE/INH****A213948 003** Dec 13, 2021WIXELA INHUB**AB** MYLAN **0.1MG/INH;EQ 0.05MG BASE/INH****A208891 001** Jan 30, 2019**AB** **0.25MG/INH;EQ 0.05MG BASE/INH****A208891 002** Jan 30, 2019**AB** **0.5MG/INH;EQ 0.05MG BASE/INH****A208891 003** Jan 30, 2019

AIRDUO DIGIHALER

+ TEVA PHARM 0.055MG/INH;EQ 0.014MG BASE/INH

N208799 004 Jul 12, 2019

+ 0.113MG/INH;EQ 0.014MG BASE/INH

N208799 005 Jul 12, 2019

+ 0.232MG/INH;EQ 0.014MG BASE/INH

N208799 006 Jul 12, 2019

AIRDUO RESPICLICK

+ TEVA PHARM 0.055MG/INH;EQ 0.014MG BASE/INH

N208799 001 Jan 27, 2017

+ 0.113MG/INH;EQ 0.014MG BASE/INH

N208799 002 Jan 27, 2017

+! 0.232MG/INH;EQ 0.014MG BASE/INH

N208799 003 Jan 27, 2017

FLUVASTATIN SODIUM

CAPSULE; ORAL

FLUVASTATIN SODIUM**AB** MYLAN PHARMS INC **EQ 20MG BASE****A090595 001** Apr 11, 2012**AB** ! **EQ 40MG BASE****A090595 002** Apr 11, 2012**AB** TEVA PHARMS **EQ 20MG BASE****A078407 001** Jun 12, 2012**AB** **EQ 40MG BASE****A078407 002** Jun 12, 2012

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM**AB** BEIJING **EQ 80MG BASE****A209397 001** Apr 26, 2021**AB** TEVA PHARMS USA **EQ 80MG BASE****A079011 001** Jan 27, 2016LESCOL XL**AB** +! SANDOZ **EQ 80MG BASE****N021192 001** Oct 06, 2000FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE**AB** ACTAVIS ELIZABETH **100MG****A091482 001** Apr 23, 2013**AB** ! **150MG****A091482 002** Nov 18, 2013**AB** ANCHEN PHARMS **100MG****A091476 001** Mar 13, 2013**AB** **150MG****A091476 002** Mar 13, 2013**AB** BIONPHARMA **100MG****A212182 002** Sep 16, 2020**AB** **150MG****A212182 001** May 11, 2020

TABLET; ORAL

FLUVOXAMINE MALEATE**AB** APOTEX **25MG****A075902 001** May 07, 2001**AB** **50MG****A075902 002** May 07, 2001**AB** **100MG****A075902 003** May 07, 2001**AB** UPSHER SMITH LABS **25MG****A075888 001** Nov 29, 2000**AB** **50MG****A075888 002** Nov 29, 2000**AB** ! **100MG****A075888 003** Nov 29, 2000LUVOX**AB** + ANI PHARMS **25MG****N021519 001** Dec 20, 2007**AB** + **50MG****N021519 002** Dec 20, 2007**AB** + **100MG****N021519 003** Dec 20, 2007

PRESCRIPTION DRUG PRODUCT LIST

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

<u>AP</u>	!	FRESENIUS KABI USA	<u>5MG/ML</u>	<u>A089202</u>	<u>001</u>	Feb 18, 1986
<u>AP</u>		XGEN PHARMS	<u>5MG/ML</u>	<u>A202522</u>	<u>001</u>	Nov 06, 2019

TABLET; ORAL

FOLIC ACID

<u>AA</u>	!	AMNEAL PHARM	<u>1MG</u>	<u>A040625</u>	<u>001</u>	Jul 21, 2005
<u>AA</u>		ATHEM	<u>1MG</u>	<u>A211064</u>	<u>001</u>	Mar 08, 2019
<u>AA</u>		CADILA PHARMS LTD	<u>1MG</u>	<u>A202437</u>	<u>001</u>	Jan 27, 2014
<u>AA</u>		CHARTWELL MOLECULAR	<u>1MG</u>	<u>A090035</u>	<u>001</u>	Jun 09, 2009
<u>AA</u>		LEADING	<u>1MG</u>	<u>A040796</u>	<u>001</u>	Jan 12, 2009
<u>AA</u>		NUVO PHARMS INC	<u>1MG</u>	<u>A204418</u>	<u>001</u>	Jul 28, 2015
<u>AA</u>		QINGDAO BAHEAL PHARM	<u>1MG</u>	<u>A091145</u>	<u>001</u>	Jul 12, 2013
<u>AA</u>	+	WATSON LABS	<u>1MG</u>	<u>A080680</u>	<u>001</u>	

FOMEPIZOLE

INJECTABLE; INJECTION

FOMEPIZOLE

<u>AP</u>		AM REGENT	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078368</u>	<u>001</u>	Dec 14, 2007
<u>AP</u>		GLAND PHARMA LTD	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A216791</u>	<u>001</u>	Jul 06, 2023
<u>AP</u>	!	MYLAN INSTITUTIONAL	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078639</u>	<u>001</u>	Mar 03, 2008
<u>AP</u>		NAVINTA LLC	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078537</u>	<u>001</u>	Mar 06, 2008

FONDAPARINUX SODIUM

INJECTABLE; SUBCUTANEOUS

ARIXTRA

<u>AP</u>	+	MYLAN IRELAND LTD	<u>2.5MG/0.5ML</u>	<u>N021345</u>	<u>001</u>	Dec 07, 2001
<u>AP</u>	+		<u>5MG/0.4ML</u>	<u>N021345</u>	<u>002</u>	May 28, 2004
<u>AP</u>	+		<u>7.5MG/0.6ML</u>	<u>N021345</u>	<u>003</u>	May 28, 2004
<u>AP</u>	+		<u>10MG/0.8ML</u>	<u>N021345</u>	<u>004</u>	May 28, 2004

FONDAPARINUX SODIUM

<u>AP</u>		DR REDDYS LABS LTD	<u>2.5MG/0.5ML</u>	<u>A091316</u>	<u>001</u>	Jul 11, 2011
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A091316</u>	<u>002</u>	Jul 11, 2011
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A091316</u>	<u>003</u>	Jul 11, 2011
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A091316</u>	<u>004</u>	Jul 11, 2011
<u>AP</u>		EUGIA PHARMA	<u>2.5MG/0.5ML</u>	<u>A206918</u>	<u>001</u>	Dec 26, 2017
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A206918</u>	<u>002</u>	Dec 26, 2017
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A206918</u>	<u>003</u>	Dec 26, 2017
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A206918</u>	<u>004</u>	Dec 26, 2017
<u>AP</u>		HENGRUI PHARMA	<u>2.5MG/0.5ML</u>	<u>A206812</u>	<u>001</u>	May 15, 2018
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A206812</u>	<u>002</u>	May 15, 2018
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A206812</u>	<u>003</u>	May 15, 2018
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A206812</u>	<u>004</u>	May 15, 2018
<u>AP</u>		SCINOPHARM TAIWAN	<u>2.5MG/0.5ML</u>	<u>A208615</u>	<u>001</u>	Nov 14, 2018
<u>AP</u>			<u>5MG/0.4ML</u>	<u>A208615</u>	<u>002</u>	Nov 14, 2018
<u>AP</u>			<u>7.5MG/0.6ML</u>	<u>A208615</u>	<u>003</u>	Nov 14, 2018
<u>AP</u>			<u>10MG/0.8ML</u>	<u>A208615</u>	<u>004</u>	Nov 14, 2018

FORMOTEROL FUMARATE

SOLUTION; INHALATION

FORMOTEROL FUMARATE

<u>AN</u>		ALEMBIC	<u>0.02MG/2ML</u>	<u>A215078</u>	<u>001</u>	Nov 22, 2021
<u>AN</u>		AUCTA	<u>0.02MG/2ML</u>	<u>A216486</u>	<u>001</u>	Nov 25, 2022
<u>AN</u>		LUPIN	<u>0.02MG/2ML</u>	<u>A215053</u>	<u>001</u>	Aug 22, 2022
<u>AN</u>		MANKIND PHARMA	<u>0.02MG/2ML</u>	<u>A215883</u>	<u>001</u>	Mar 22, 2023
<u>AN</u>		SLAYBACK PHARMA LLC	<u>0.02MG/2ML</u>	<u>A215621</u>	<u>001</u>	Dec 13, 2022
<u>AN</u>		TEVA PHARMS USA INC	<u>0.02MG/2ML</u>	<u>A091141</u>	<u>001</u>	Jun 22, 2021

PERFOROMIST

<u>AN</u>	+	MYLAN SPECLT	<u>0.02MG/2ML</u>	<u>N022007</u>	<u>001</u>	May 11, 2007
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FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION

BEVESPI AEROSPHERE

+	!	ASTRAZENECA	0.0048MG/INH;0.0090MG/INH	N208294	001	Apr 25, 2016
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FORMOTEROL FUMARATE; MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

DULERA

+		ORGANON LLC	0.005MG/INH;0.05MG/INH	N022518	003	Aug 12, 2019
+	!		0.005MG/INH;0.1MG/INH	N022518	001	Jun 22, 2010
+	!		0.005MG/INH;0.2MG/INH	N022518	002	Jun 22, 2010

PRESCRIPTION DRUG PRODUCT LIST

FOSAMPRENAVIR CALCIUM

TABLET; ORAL

FOSAMPRENAVIR CALCIUM

AB	MYLAN	EQ 700MG BASE	A204060 001	Apr 15, 2016
AB	SUN PHARM	EQ 700MG BASE	A204024 001	Nov 20, 2019

LEXIVA

AB	+! VIIV HLTHCARE	EQ 700MG BASE	N021548 001	Oct 20, 2003
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FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND

AP	+! MERCK AND CO INC	EQ 150MG BASE/VIAL	N022023 002	Nov 12, 2010
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FOSAPREPITANT DIMEGLUMINE

AP	ACCORD HLTHCARE	EQ 150MG BASE/VIAL	A204025 001	Aug 26, 2020
AP	ASPIRO	EQ 150MG BASE/VIAL	A214616 001	Jul 29, 2021
AP	BAXTER HLTHCARE CORP	EQ 150MG BASE/VIAL	A211860 001	Sep 05, 2019
AP	BE PHARMS	EQ 150MG BASE/VIAL	A212309 001	Sep 05, 2019
AP	CHIA TAI TIANQING	EQ 150MG BASE/VIAL	A212143 001	Mar 03, 2021
AP	DR REDDYS	EQ 150MG BASE/VIAL	A211160 001	Dec 09, 2020
AP	EUGIA PHARMA	EQ 150MG BASE/VIAL	A210625 001	Jan 12, 2021
AP	FRESENIUS KABI USA	EQ 150MG BASE/VIAL	A206197 001	Jun 09, 2016
AP	LUPIN LTD	EQ 150MG BASE/VIAL	A210689 001	Sep 05, 2019
AP	MSN	EQ 150MG BASE/VIAL	A209965 001	Sep 05, 2019
AP	MYLAN LABS LTD	EQ 150MG BASE/VIAL	A204015 002	Sep 05, 2019
AP	NAVINTA LLC	EQ 150MG BASE/VIAL	A212957 002	Aug 20, 2020
AP	PIRAMAL CRITICAL	EQ 150MG BASE/VIAL	A214683 001	May 16, 2023
AP	QILU PHARM HAINAN	EQ 150MG BASE/VIAL	A213106 001	Sep 08, 2020
	TEVA PHARMS USA	EQ 150MG BASE/VIAL	N210064 001	Sep 05, 2019

SOLUTION; INTRAVENOUS

FOCINVEZ

	+! STERISCIENCE	EQ 150MG BASE/50ML (EQ 3MG BASE/ML)	N216686 001	Aug 22, 2023
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FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

AP	AVET LIFESCIENCES	2.4GM/100ML	A213807 001	Jun 05, 2023
AP	BEIJING	2.4GM/100ML	A213987 001	Nov 29, 2023
AP	FRESENIUS KABI USA	2.4GM/100ML	A212483 001	Jan 29, 2021
AP	GLAND PHARMA LTD	2.4GM/100ML	A213001 001	Apr 21, 2021

FOSCAVIR

AP	+! CLINIGEN HLTHCARE	2.4GM/100ML	N020068 001	Sep 27, 1991
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FOSDENOPTERIN HYDROBROMIDE

POWDER; INTRAVENOUS

NULIBRY

	+! SENTYNL THERAPS INC	EQ 9.5MG BASE/VIAL	N214018 001	Feb 26, 2021
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FOSFOMYCIN TROMETHAMINE

FOR SOLUTION; ORAL

FOSFOMYCIN TROMETHAMINE

AA	ALKEM LABS LTD	EQ 3GM BASE/PACKET	A214554 001	Oct 21, 2021
AA	CIPLA	EQ 3GM BASE/PACKET	A211881 001	Jan 26, 2022
AA	XIROMED	EQ 3GM BASE/PACKET	A212548 001	Oct 06, 2020

MONUROL

AA	+! ZAMBON SPA	EQ 3GM BASE/PACKET	N050717 001	Dec 19, 1996
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FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

AB	APOTEX INC	10MG	A076906 001	May 17, 2005
AB		20MG	A076906 002	May 17, 2005
AB		40MG	A076906 003	May 17, 2005
AB	AUROBINDO PHARMA LTD	10MG	A091163 001	Mar 30, 2011
AB		20MG	A091163 002	Mar 30, 2011
AB		40MG	A091163 003	Mar 30, 2011
AB	CHARTWELL RX	10MG	A076483 001	Apr 23, 2004
AB		20MG	A076483 002	Apr 23, 2004
AB		40MG	A076483 003	Apr 23, 2004
AB	INVAGEN PHARMS	10MG	A077222 001	Apr 20, 2005
AB		20MG	A077222 002	Apr 20, 2005
AB		40MG	A077222 003	Apr 20, 2005
AB	PRINSTON INC	10MG	A205670 001	Aug 29, 2016
AB		20MG	A205670 002	Aug 29, 2016

PRESCRIPTION DRUG PRODUCT LIST

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

AB		40MG	A205670 003	Aug 29, 2016
AB	TEVA	10MG	A076139 001	Nov 25, 2003
AB		20MG	A076139 002	Nov 25, 2003
AB	!	40MG	A076139 003	Nov 25, 2003

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

AB	AUROBINDO PHARMA	10MG;12.5MG	A079245 001	Jul 09, 2009
AB	!	20MG;12.5MG	A079245 002	Jul 09, 2009
AB	INVAGEN PHARMS	10MG;12.5MG	A090228 001	Jul 09, 2009
AB		20MG;12.5MG	A090228 002	Jul 09, 2009
AB	SANDOZ	10MG;12.5MG	A076961 001	Sep 28, 2005
AB		20MG;12.5MG	A076961 002	Sep 28, 2005

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

SOLUTION; INTRAVENOUS

AKYNZEO

+	!	HELSINN HLTHCARE	EQ 235MG BASE/20ML (EQ 11.75MG BASE/ML);EQ 0.25MG BASE/20ML (EQ 0.0125MG BASE/ML)	N210493 002	May 27, 2020
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FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

CEREBYX

AP	+	!	PARKE DAVIS	EQ 50MG PHENYTOIN NA/ML	N020450 001	Aug 05, 1996
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FOSPHENYTOIN SODIUM

AP			FRESENIUS KABI USA	EQ 50MG PHENYTOIN NA/ML	A078052 001	Aug 06, 2007
AP			GLAND PHARMA LTD	EQ 50MG PHENYTOIN NA/ML	A214926 001	Oct 13, 2023
AP			HIKMA	EQ 50MG PHENYTOIN NA/ML	A077481 001	Aug 06, 2007
AP				EQ 50MG PHENYTOIN NA/ML	A077989 001	Aug 06, 2007
AP			HIKMA FARMACEUTICA	EQ 50MG PHENYTOIN NA/ML	A078765 001	Dec 02, 2009
AP			SUN PHARM	EQ 50MG PHENYTOIN NA/ML	A078417 001	Mar 18, 2008
AP			WOCKHARDT	EQ 50MG PHENYTOIN NA/ML	A078137 001	Aug 06, 2007

FOSTAMATINIB DISODIUM

TABLET; ORAL

TAVALISSE

+		RIGEL PHARMS INC	EQ 100MG BASE	N209299 001	Apr 17, 2018
+	!		EQ 150MG BASE	N209299 002	Apr 17, 2018

FOSTEMSAVIR TROMETHAMINE

TABLET, EXTENDED RELEASE; ORAL

RUKOBIA

+	!	VIIV HLTHCARE	EQ 600MG BASE	N212950 001	Jul 02, 2020
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FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVA

AB	+	!	ENDO PHARMS	EQ 2.5MG BASE	N021006 001	Nov 08, 2001
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FROVATRIPTAN SUCCINATE

AB			AMNEAL PHARMS CO	EQ 2.5MG BASE	A211292 001	Nov 06, 2018
AB			GLENMARK PHARMS LTD	EQ 2.5MG BASE	A204730 001	Mar 11, 2016
AB			RENATA	EQ 2.5MG BASE	A213891 001	Apr 06, 2022

FRUQUINTINIB

CAPSULE; ORAL

FRUZAQLA

+		TAKEDA PHARMS USA	1MG	N217564 001	Nov 08, 2023
+	!		5MG	N217564 002	Nov 08, 2023

FULVESTRANT

SOLUTION; INTRAMUSCULAR

FASLODEX

AO	+	!	ASTRAZENECA	250MG/5ML (50MG/ML)	N021344 001	Apr 25, 2002
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FULVESTRANT

AO			ACCORD HLTHCARE	250MG/5ML (50MG/ML)	A211689 001	Nov 17, 2020
AO			ALEMBIC	250MG/5ML (50MG/ML)	A215077 001	Dec 22, 2022
AO			AMNEAL	250MG/5ML (50MG/ML)	A210044 001	Mar 04, 2019
AO			CHIA TAI TIANQING	250MG/5ML (50MG/ML)	A211422 001	Feb 07, 2020
AO			DR REDDYS	250MG/5ML (50MG/ML)	A209246 001	Aug 07, 2020
AO			EUGIA PHARMA	250MG/5ML (50MG/ML)	A208811 001	Jul 23, 2019
AO				250MG/5ML (50MG/ML)	A215169 001	Jun 30, 2023
AO			GLENMARK PHARMS INC	250MG/5ML (50MG/ML)	A207754 001	Aug 22, 2019

PRESCRIPTION DRUG PRODUCT LIST

FULVESTRANT

SOLUTION; INTRAMUSCULAR

FULVESTRANT

<u>AO</u>	HBT LABS INC	<u>250MG/5ML (50MG/ML)</u>	<u>A209714 001</u>	Nov 21, 2019
<u>AO</u>	JIANGSU HANSOH PHARM	<u>250MG/5ML (50MG/ML)</u>	<u>A214682 001</u>	Feb 10, 2022
<u>AO</u>	SAGENT PHARMS INC	<u>250MG/5ML (50MG/ML)</u>	<u>A205871 001</u>	Aug 22, 2019
<u>AO</u>	SANDOZ	<u>250MG/5ML (50MG/ML)</u>	<u>A205935 001</u>	May 14, 2019
<u>AO</u>	XIROMED	<u>250MG/5ML (50MG/ML)</u>	<u>A213553 001</u>	Aug 13, 2021
<u>AO</u>	ZYDUS PHARMS	<u>250MG/5ML (50MG/ML)</u>	<u>A215234 001</u>	Jul 29, 2021
	FRESENIUS KABI USA	250MG/5ML (50MG/ML)	N210326 001	May 20, 2019
	TEVA PHARMS USA INC	250MG/5ML (50MG/ML)	N210063 001	Aug 19, 2019

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

<u>AP</u>	ACCORD HLTHCARE	<u>10MG/ML</u>	<u>A070017 001</u>	Dec 15, 1986
<u>AP</u>	AMNEAL PHARMS CO	<u>10MG/ML</u>	<u>A207552 001</u>	Jul 20, 2016
<u>AP</u>	AREVA PHARMS	<u>10MG/ML</u>	<u>A208435 001</u>	Dec 18, 2020
<u>AP</u>	AVET LIFESCIENCES	<u>10MG/ML</u>	<u>A203428 001</u>	Aug 26, 2014
<u>AP</u>	! BAXTER HLTHCARE CORP	<u>10MG/ML</u>	<u>A202747 001</u>	Jan 27, 2014
<u>AP</u>	EUGIA PHARMA	<u>10MG/ML</u>	<u>A212174 001</u>	May 03, 2019
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>N018902 001</u>	May 22, 1984
<u>AP</u>	GLAND PHARMA LTD	<u>10MG/ML</u>	<u>A213902 001</u>	Jul 01, 2020
<u>AP</u>	HOSPIRA	<u>10MG/ML</u>	<u>A075241 001</u>	May 28, 1999
<u>AP</u>		<u>10MG/ML</u>	<u>N018667 001</u>	May 28, 1982
<u>AP</u>	MANKIND PHARMA	<u>10MG/ML</u>	<u>A216860 001</u>	Dec 16, 2022
<u>AP</u>	MEITHEAL	<u>10MG/ML</u>	<u>A212803 001</u>	Jan 27, 2022
<u>AP</u>	SAGENT	<u>10MG/ML</u>	<u>A214766 001</u>	Jan 27, 2021
<u>AP</u>	WOCKHARDT	<u>10MG/ML</u>	<u>A077941 001</u>	Mar 22, 2007

SOLUTION; ORAL

FUROSEMIDE

! HIKMA

10MG/ML
40MG/5MLA070434 001 Apr 22, 1987
A070433 001 Apr 22, 1987

SOLUTION; SUBCUTANEOUS

FUROSCIX

+! SCPHARMACEUTICALS

80MG/10ML (8MG/ML)

N209988 001 Oct 07, 2022

TABLET; ORAL

FUROSEMIDE

<u>AB</u>	EPIC PHARMA LLC	<u>20MG</u>	<u>N018569 002</u>	
<u>AB</u>		<u>40MG</u>	<u>N018569 001</u>	
<u>AB</u>		<u>80MG</u>	<u>N018569 005</u>	Aug 14, 1984
<u>AB</u>	GRAVITI PHARMS	<u>20MG</u>	<u>A216629 001</u>	Oct 17, 2022
<u>AB</u>		<u>40MG</u>	<u>A216629 002</u>	Oct 17, 2022
<u>AB</u>		<u>80MG</u>	<u>A216629 003</u>	Oct 17, 2022
<u>AB</u>	HIKMA	<u>20MG</u>	<u>N018823 001</u>	Nov 10, 1983
<u>AB</u>		<u>40MG</u>	<u>N018823 002</u>	Nov 10, 1983
<u>AB</u>		<u>80MG</u>	<u>A070086 001</u>	Jan 24, 1986
<u>AB</u>	IPCA LABS LTD	<u>20MG</u>	<u>A078010 001</u>	Sep 18, 2006
<u>AB</u>		<u>40MG</u>	<u>A078010 002</u>	Sep 18, 2006
<u>AB</u>		<u>80MG</u>	<u>A078010 003</u>	Sep 18, 2006
<u>AB</u>	LEADING	<u>20MG</u>	<u>A077293 001</u>	Nov 09, 2005
<u>AB</u>		<u>40MG</u>	<u>A077293 002</u>	Nov 09, 2005
<u>AB</u>		<u>80MG</u>	<u>A077293 003</u>	Nov 09, 2005
<u>AB</u>	MYLAN	<u>20MG</u>	<u>N018487 001</u>	
<u>AB</u>		<u>40MG</u>	<u>N018487 002</u>	
<u>AB</u>		<u>80MG</u>	<u>A070082 001</u>	Oct 29, 1986
<u>AB</u>	PRINSTON INC	<u>20MG</u>	<u>A076796 001</u>	Mar 26, 2004
<u>AB</u>		<u>40MG</u>	<u>A076796 002</u>	Mar 26, 2004
<u>AB</u>		<u>80MG</u>	<u>A076796 003</u>	Mar 26, 2004
	<u>LASIX</u>			
<u>AB</u>	+ VALIDUS PHARMS	<u>20MG</u>	<u>N016273 002</u>	
<u>AB</u>	+	<u>40MG</u>	<u>N016273 001</u>	
<u>AB</u>	+!	<u>80MG</u>	<u>N016273 003</u>	

FUTIBATINIB

TABLET; ORAL

LYTGOBI

+! TAIHO ONCOLOGY

4MG

N214801 001 Sep 30, 2022

PRESCRIPTION DRUG PRODUCT LIST

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>	ACI	<u>100MG</u>	<u>A206943 001</u>	May 14, 2018
<u>AB</u>		<u>300MG</u>	<u>A206943 002</u>	May 14, 2018
<u>AB</u>		<u>400MG</u>	<u>A206943 003</u>	May 14, 2018
<u>AB</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A075350 001</u>	Sep 12, 2003
<u>AB</u>		<u>300MG</u>	<u>A075350 002</u>	Sep 12, 2003
<u>AB</u>		<u>400MG</u>	<u>A075350 003</u>	Sep 12, 2003
<u>AB</u>	ALKEM	<u>100MG</u>	<u>A090858 001</u>	Dec 17, 2010
<u>AB</u>		<u>300MG</u>	<u>A090858 002</u>	Dec 17, 2010
<u>AB</u>		<u>400MG</u>	<u>A090858 003</u>	Dec 17, 2010
<u>AB</u>	AMNEAL PHARMS NY	<u>100MG</u>	<u>A078428 001</u>	Jul 25, 2007
<u>AB</u>		<u>300MG</u>	<u>A078428 002</u>	Jul 25, 2007
<u>AB</u>		<u>400MG</u>	<u>A078428 003</u>	Jul 25, 2007
<u>AB</u>	ASCENT PHARMS INC	<u>100MG</u>	<u>A214956 001</u>	May 10, 2021
<u>AB</u>		<u>300MG</u>	<u>A214956 002</u>	May 10, 2021
<u>AB</u>		<u>400MG</u>	<u>A214956 003</u>	May 10, 2021
<u>AB</u>	AUROBINDO PHARMA	<u>100MG</u>	<u>A078787 001</u>	Jan 31, 2008
<u>AB</u>		<u>300MG</u>	<u>A078787 002</u>	Jan 31, 2008
<u>AB</u>		<u>400MG</u>	<u>A078787 003</u>	Jan 31, 2008
<u>AB</u>	CSPC OUYI	<u>100MG</u>	<u>A075477 001</u>	Mar 23, 2005
<u>AB</u>		<u>300MG</u>	<u>A075477 002</u>	Mar 23, 2005
<u>AB</u>		<u>400MG</u>	<u>A075477 003</u>	Mar 23, 2005
<u>AB</u>	GRANULES	<u>100MG</u>	<u>A075360 001</u>	Apr 06, 2005
<u>AB</u>		<u>300MG</u>	<u>A075360 002</u>	Apr 06, 2005
<u>AB</u>		<u>400MG</u>	<u>A075360 003</u>	Apr 06, 2005
<u>AB</u>	GRAVITI PHARMS	<u>100MG</u>	<u>A207099 001</u>	Mar 24, 2017
<u>AB</u>		<u>300MG</u>	<u>A207099 002</u>	Mar 24, 2017
<u>AB</u>		<u>400MG</u>	<u>A207099 003</u>	Mar 24, 2017
<u>AB</u>	INVAGEN PHARMS	<u>100MG</u>	<u>A090705 001</u>	Dec 30, 2009
<u>AB</u>		<u>300MG</u>	<u>A090705 002</u>	Dec 30, 2009
<u>AB</u>		<u>400MG</u>	<u>A090705 003</u>	Dec 30, 2009
<u>AB</u>	IPCA LABS LTD	<u>100MG</u>	<u>A208928 001</u>	Nov 20, 2023
<u>AB</u>		<u>300MG</u>	<u>A208928 002</u>	Nov 20, 2023
<u>AB</u>		<u>400MG</u>	<u>A208928 003</u>	Nov 20, 2023
<u>AB</u>	LAURUS	<u>100MG</u>	<u>A217546 001</u>	May 12, 2023
<u>AB</u>		<u>300MG</u>	<u>A217546 002</u>	May 12, 2023
<u>AB</u>		<u>400MG</u>	<u>A217546 003</u>	May 12, 2023
<u>AB</u>	MARKSANS PHARMA	<u>100MG</u>	<u>A090007 001</u>	Jul 21, 2011
<u>AB</u>		<u>300MG</u>	<u>A090007 002</u>	Jul 21, 2011
<u>AB</u>		<u>400MG</u>	<u>A090007 003</u>	Jul 21, 2011
<u>AB</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A204989 001</u>	Feb 18, 2016
<u>AB</u>		<u>300MG</u>	<u>A204989 002</u>	Feb 18, 2016
<u>AB</u>		<u>400MG</u>	<u>A204989 003</u>	Feb 18, 2016
<u>AB</u>	STRIDES PHARMA	<u>100MG</u>	<u>A211314 001</u>	Oct 16, 2018
<u>AB</u>		<u>300MG</u>	<u>A211314 002</u>	Oct 16, 2018
<u>AB</u>		<u>400MG</u>	<u>A211314 003</u>	Oct 16, 2018
<u>AB</u>	SUN PHARM INDS LTD	<u>100MG</u>	<u>A077242 001</u>	Aug 24, 2006
<u>AB</u>		<u>300MG</u>	<u>A077242 002</u>	Aug 24, 2006
<u>AB</u>		<u>400MG</u>	<u>A077242 003</u>	Aug 24, 2006
<u>AB</u>	TARO	<u>100MG</u>	<u>A077261 001</u>	Aug 02, 2013
<u>AB</u>		<u>300MG</u>	<u>A077261 002</u>	Aug 02, 2013
<u>AB</u>		<u>400MG</u>	<u>A077261 003</u>	Aug 02, 2013
<u>AB</u>	ZHEJIANG YONGTAI	<u>100MG</u>	<u>A213603 001</u>	Aug 17, 2020
<u>AB</u>		<u>300MG</u>	<u>A213603 002</u>	Aug 17, 2020
<u>AB</u>		<u>400MG</u>	<u>A213603 003</u>	Aug 17, 2020

NEURONTIN

<u>AB</u>	+ VIATRIS	<u>100MG</u>	<u>N020235 001</u>	Dec 30, 1993
<u>AB</u>	+	<u>300MG</u>	<u>N020235 002</u>	Dec 30, 1993
<u>AB</u>	+	<u>400MG</u>	<u>N020235 003</u>	Dec 30, 1993

SOLUTION; ORAL

GABAPENTIN

<u>AA</u>	ACELLA PHARMS LLC	<u>250MG/5ML</u>	<u>A076403 001</u>	May 01, 2012
<u>AA</u>	AMNEAL PHARMS	<u>250MG/5ML</u>	<u>A202024 001</u>	Mar 23, 2012
<u>AA</u>	BELCHER	<u>250MG/5ML</u>	<u>A091286 001</u>	Mar 14, 2016
<u>AA</u>	MISSION PHARMACAL	<u>250MG/5ML</u>	<u>A078974 001</u>	Feb 18, 2011
<u>AA</u>	RUBICON	<u>250MG/5ML</u>	<u>A216492 001</u>	Jan 18, 2023
<u>AA</u>	TARO	<u>250MG/5ML</u>	<u>A076672 001</u>	Jul 03, 2013

NEURONTIN

<u>AA</u>	+! VIATRIS	<u>250MG/5ML</u>	<u>N021129 001</u>	Mar 02, 2000
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PRESCRIPTION DRUG PRODUCT LIST

GABAPENTIN

TABLET; ORAL

GABAPENTIN

<u>AB</u>	ACI	<u>600MG</u>	<u>A203244 002</u>	Jul 12, 2013
<u>AB</u>		<u>800MG</u>	<u>A203244 001</u>	Jul 12, 2013
<u>AB</u>	ACTAVIS ELIZABETH	<u>600MG</u>	<u>A075694 001</u>	Oct 21, 2004
<u>AB</u>		<u>800MG</u>	<u>A075694 002</u>	Oct 21, 2004
<u>AB</u>	ALKEM LABS LTD	<u>600MG</u>	<u>A206402 001</u>	Dec 23, 2015
<u>AB</u>		<u>800MG</u>	<u>A206402 002</u>	Dec 23, 2015
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077894 001</u>	Oct 10, 2006
<u>AB</u>		<u>300MG</u>	<u>A077894 002</u>	Oct 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077894 003</u>	Oct 10, 2006
<u>AB</u>	ASCENT PHARMS INC	<u>600MG</u>	<u>A214957 001</u>	Oct 01, 2021
<u>AB</u>		<u>800MG</u>	<u>A214957 002</u>	Oct 01, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A200651 001</u>	Oct 06, 2011
<u>AB</u>		<u>800MG</u>	<u>A200651 002</u>	Oct 06, 2011
<u>AB</u>	CSPC OUYI	<u>600MG</u>	<u>A207057 001</u>	Oct 26, 2017
<u>AB</u>		<u>800MG</u>	<u>A207057 002</u>	Oct 26, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>600MG</u>	<u>A077662 001</u>	Aug 18, 2006
<u>AB</u>		<u>800MG</u>	<u>A077662 002</u>	Aug 18, 2006
<u>AB</u>	GRANULES	<u>600MG</u>	<u>A217116 001</u>	Mar 28, 2023
<u>AB</u>		<u>800MG</u>	<u>A217116 002</u>	Mar 28, 2023
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A202764 001</u>	Oct 16, 2012
<u>AB</u>		<u>800MG</u>	<u>A202764 002</u>	Oct 16, 2012
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>100MG</u>	<u>A076017 001</u>	Apr 28, 2004
<u>AB</u>		<u>300MG</u>	<u>A076017 002</u>	Apr 28, 2004
<u>AB</u>		<u>400MG</u>	<u>A076017 003</u>	Apr 28, 2004
<u>AB</u>	RISING	<u>600MG</u>	<u>A217995 001</u>	Jul 19, 2023
<u>AB</u>		<u>800MG</u>	<u>A217995 002</u>	Jul 19, 2023
<u>AB</u>	RUBICON	<u>600MG</u>	<u>A077661 004</u>	Sep 13, 2006
<u>AB</u>		<u>800MG</u>	<u>A077661 005</u>	Sep 13, 2006
<u>AB</u>	SCIEGEN PHARMS INC	<u>600MG</u>	<u>A205101 001</u>	Feb 04, 2016
<u>AB</u>		<u>800MG</u>	<u>A205101 002</u>	Feb 04, 2016
<u>AB</u>	SUN PHARM INDS LTD	<u>600MG</u>	<u>A077525 001</u>	Aug 24, 2006
<u>AB</u>		<u>800MG</u>	<u>A077525 002</u>	Aug 24, 2006
<u>AB</u>	ZYDUS PHARMS USA INC	<u>600MG</u>	<u>A078926 001</u>	Feb 11, 2011
<u>AB</u>		<u>800MG</u>	<u>A078926 002</u>	Feb 11, 2011

NEURONTIN

<u>AB</u>	+ VIATRIS	<u>600MG</u>	<u>N020882 001</u>	Oct 09, 1998
<u>AB</u>	+!	<u>800MG</u>	<u>N020882 002</u>	Oct 09, 1998

GRALISE

BX	+! ALMATICA	300MG	N022544 001	Jan 28, 2011
BX	+!	600MG	N022544 002	Jan 28, 2011
	+	450MG	N022544 003	Apr 18, 2023
	+	750MG	N022544 004	Apr 18, 2023
	+	900MG	N022544 005	Apr 18, 2023

GABAPENTIN ENACARBIL

TABLET, EXTENDED RELEASE; ORAL

HORIZANT

+	AZURITY	300MG	N022399 002	Dec 13, 2011
+	!	600MG	N022399 001	Apr 06, 2011

GADOBENATE DIMEGLUMINE

INJECTABLE; INTRAVENOUS

MULTIHANCE

+	BRACCO	2.645GM/5ML (529MG/ML)	N021357 001	Nov 23, 2004
+		5.29GM/10ML (529MG/ML)	N021357 002	Nov 23, 2004
+		7.935GM/15ML (529MG/ML)	N021357 003	Nov 23, 2004
+		10.58GM/20ML (529MG/ML)	N021357 004	Nov 23, 2004

MULTIHANCE MULTIPACK

+	BRACCO	26.45GM/50ML (529MG/ML)	N021358 001	Nov 23, 2004
+		52.9GM/100ML (529MG/ML)	N021358 002	Nov 23, 2004

GADOBUTROL

SOLUTION; INTRAVENOUS

GADAVIST

<u>AP</u>	+!	BAYER HLTHCARE	<u>1.20944GM/2ML (604.72MG/ML)</u>	<u>N201277 006</u>	Dec 18, 2013
<u>AP</u>	+		<u>4.5354GM/7.5ML (604.72MG/ML)</u>	<u>N201277 001</u>	Mar 14, 2011
<u>AP</u>	+		<u>6.0472GM/10ML (604.72MG/ML)</u>	<u>N201277 002</u>	Mar 14, 2011
<u>AP</u>	+		<u>9.0708GM/15ML (604.72MG/ML)</u>	<u>N201277 003</u>	Mar 14, 2011

PRESCRIPTION DRUG PRODUCT LIST

GADOBUTROL

SOLUTION; INTRAVENOUS

GADAVIST

<u>AP</u>	<u>+</u> !	<u>18.1416GM/30ML (604.72MG/ML)</u>	<u>N201277 004</u>	Mar 14, 2011
<u>AP</u>	<u>+</u> !	<u>39.3068GM/65ML (604.72MG/ML)</u>	<u>N201277 005</u>	Mar 14, 2011

GADOBUTROL

<u>AP</u>	HAINAN POLY PHARM	<u>4.5354GM/7.5ML (604.72MG/ML)</u>	<u>A217480 001</u>	Mar 15, 2023
<u>AP</u>		<u>9.0708GM/15ML (604.72MG/ML)</u>	<u>A217480 002</u>	Mar 15, 2023
<u>AP</u>	HENGRUI PHARMA	<u>1.20944GM/2ML (604.72MG/ML)</u>	<u>A215061 001</u>	Nov 17, 2022
<u>AP</u>		<u>4.5354GM/7.5ML (604.72MG/ML)</u>	<u>A215061 002</u>	Nov 17, 2022
<u>AP</u>		<u>6.0472GM/10ML (604.72MG/ML)</u>	<u>A215061 003</u>	Nov 17, 2022
<u>AP</u>		<u>9.0708GM/15ML (604.72MG/ML)</u>	<u>A215061 004</u>	Nov 17, 2022
<u>AP</u>		<u>18.1416GM/30ML (604.72MG/ML)</u>	<u>A216081 001</u>	Sep 08, 2023
<u>AP</u>		<u>39.3068GM/65ML (604.72MG/ML)</u>	<u>A216081 002</u>	Sep 08, 2023

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

<u>+</u> !	GE HEALTHCARE	287MG/ML	N020123 001	Jan 08, 1993
<u>+</u> !		28.7GM/100ML (287MG/ML)	N022066 002	Sep 05, 2007

GADOPICLENOL

SOLUTION; INTRAVENOUS

ELUCIREM

<u>+</u> !	GUERBET	1.4553GM/3ML (485.1MG/ML)	N216986 001	Sep 21, 2022
<u>+</u> !		3.63825GM/7.5ML (485.1MG/ML)	N216986 002	Sep 21, 2022
<u>+</u> !		4.851GM/10ML (485.1MG/ML)	N216986 003	Sep 21, 2022
<u>+</u> !		7.2765GM/15ML (485.1MG/ML)	N216986 004	Sep 21, 2022
<u>+</u> !		14.553GM/30ML (485.1MG/ML)	N216986 005	Sep 21, 2022
<u>+</u> !		24.255GM/50ML (485.1MG/ML)	N216986 006	Sep 21, 2022
<u>+</u> !		48.51GM/100ML (485.1MG/ML)	N216986 007	Sep 21, 2022

GADOTERATE MEGLUMINE

SOLUTION; INTRAVENOUS

CLARISCAN

<u>AP</u>	GE HEALTHCARE	<u>7.538GM/20ML (376.9MG/ML)</u>	<u>A210016 003</u>	Nov 01, 2019
<u>AP</u>		<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>A210016 005</u>	Nov 24, 2020
<u>AP</u>		<u>3.769GM/10ML (376.9MG/ML)</u>	<u>A210016 001</u>	Nov 01, 2019
<u>AP</u>		<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>A210016 002</u>	Nov 01, 2019
<u>AP</u>		<u>37.69GM/100ML (376.9MG/ML)</u>	<u>A210016 004</u>	Aug 04, 2020

DOTAREM

<u>AP</u>	<u>+</u> !	GUERBET	<u>37.69GM/100ML (376.9MG/ML)</u>	<u>N204781 001</u>	Mar 20, 2013
<u>AP</u>	<u>+</u> !		<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>N204781 005</u>	Mar 31, 2017
<u>AP</u>	<u>+</u> !		<u>3.769GM/10ML (376.9MG/ML)</u>	<u>N204781 002</u>	Mar 20, 2013
<u>AP</u>	<u>+</u> !		<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>N204781 003</u>	Mar 20, 2013
<u>AP</u>	<u>+</u> !		<u>7.538GM/20ML (376.9MG/ML)</u>	<u>N204781 004</u>	Mar 20, 2013

GADOTERATE MEGLUMINE

<u>AP</u>	HENGRUI PHARMA	<u>1.8845GM/5ML (376.9MG/ML)</u>	<u>A215304 001</u>	Apr 11, 2022
<u>AP</u>		<u>3.769GM/10ML (376.9MG/ML)</u>	<u>A215304 002</u>	Apr 11, 2022
<u>AP</u>		<u>5.6535GM/15ML (376.9MG/ML)</u>	<u>A215304 003</u>	Apr 11, 2022
<u>AP</u>		<u>7.538GM/20ML (376.9MG/ML)</u>	<u>A215304 004</u>	Apr 11, 2022
<u>AP</u>		<u>37.69GM/100ML (376.9MG/ML)</u>	<u>A215304 005</u>	Apr 15, 2022

GADOTERIDOL

INJECTABLE; INJECTION

PROHANCE

<u>+</u> !	BRACCO	279.3MG/ML	N020131 001	Nov 16, 1992
<u>+</u> !	PROHANCE MULTIPACK			
<u>+</u> !	BRACCO	279.3MG/ML	N021489 001	Oct 09, 2003

GADOXETATE DISODIUM

SOLUTION; INTRAVENOUS

EOVIST

<u>+</u> !	BAYER HLTHCARE	1.8143GM/10ML (181.43MG/ML)	N022090 001	Jul 03, 2008
<u>+</u>		2.72145GM/15ML (181.43MG/ML)	N022090 002	Feb 04, 2013

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

<u>AB</u>	<u>!</u>	AUROBINDO PHARMA	<u>EQ 8MG BASE</u>	<u>A204895 001</u>	Aug 05, 2016
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A204895 002</u>	Aug 05, 2016
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A204895 003</u>	Aug 05, 2016
<u>AB</u>		BARR	<u>EQ 8MG BASE</u>	<u>A078189 001</u>	Sep 15, 2008
<u>AB</u>			<u>EQ 16MG BASE</u>	<u>A078189 002</u>	Sep 15, 2008
<u>AB</u>			<u>EQ 24MG BASE</u>	<u>A078189 003</u>	Sep 15, 2008

PRESCRIPTION DRUG PRODUCT LIST

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE;ORAL

GALANTAMINE HYDROBROMIDE

AB	SUN PHARM	EQ 8MG BASE	A090178 001	Feb 02, 2011
AB		EQ 16MG BASE	A090178 002	Feb 02, 2011
AB		EQ 24MG BASE	A090178 003	Feb 02, 2011
AB	WATSON LABS	EQ 8MG BASE	A079028 001	Dec 15, 2008
AB		EQ 16MG BASE	A079028 002	Dec 15, 2008
AB		EQ 24MG BASE	A079028 003	Dec 15, 2008

SOLUTION;ORAL

GALANTAMINE HYDROBROMIDE

! HIKMA

4MG/ML

A078185 001 Jan 30, 2009

TABLET;ORAL

GALANTAMINE HYDROBROMIDE

AB	AUROBINDO PHARMA LTD	EQ 4MG BASE	A090957 001	Mar 29, 2011
AB		EQ 8MG BASE	A090957 002	Mar 29, 2011
AB		EQ 12MG BASE	A090957 003	Mar 29, 2011
AB	BARR	EQ 4MG BASE	A077605 001	Aug 28, 2008
AB		EQ 8MG BASE	A077605 002	Aug 28, 2008
AB		EQ 12MG BASE	A077605 003	Aug 28, 2008
AB	DR REDDYS LABS LTD	EQ 4MG BASE	A077593 001	Sep 11, 2008
AB		EQ 8MG BASE	A077593 002	Sep 11, 2008
AB		EQ 12MG BASE	A077593 003	Sep 11, 2008
AB	SANDOZ	EQ 4MG BASE	A077589 001	Jun 22, 2009
AB		EQ 8MG BASE	A077589 002	Jun 22, 2009
AB		EQ 12MG BASE	A077589 003	Jun 22, 2009
AB	! YABAO PHARM	EQ 4MG BASE	A077604 001	Feb 06, 2009
AB		EQ 8MG BASE	A077604 002	Feb 06, 2009
AB		EQ 12MG BASE	A077604 003	Feb 06, 2009
AB	ZYDUS PHARMS USA INC	EQ 4MG BASE	A078898 001	Feb 17, 2011
AB		EQ 8MG BASE	A078898 002	Feb 17, 2011
AB		EQ 12MG BASE	A078898 003	Feb 17, 2011

GALLIUM CITRATE GA-67

INJECTABLE;INJECTION

GALLIUM CITRATE GA 67

BS CURIUM 2mCi/ML N018058 001

GALLIUM DOTATATE GA-68

POWDER;INTRAVENOUS

NETSPOT

+! AAA USA INC

2.1-5.5mCi/ML

N208547 001 Jun 01, 2016

GALLIUM GA-68 EDOTREOTIDE

SOLUTION;INTRAVENOUS

GALLIUM GA 68 EDOTREOTIDE

+! UIHC PET IMAGING

0.5-4mCi/ML

N210828 001 Aug 21, 2019

GALLIUM GA-68 GOZETOTIDE

POWDER;INTRAVENOUS

ILLUCCIX

+! TELIX

N/A

N214032 001 Dec 17, 2021

LOCAMETZ

+! NOVARTIS

N/A

N215841 001 Mar 23, 2022

SOLUTION;INTRAVENOUS

GALLIUM GA 68 GOZETOTIDE

+! UNIV CA LOS ANGELES

0.5-5mCi/mL

N212642 001 Dec 01, 2020

+! UNIV OF CA SAN FRAN

0.5-5mCi/mL

N212643 001 Dec 01, 2020

GANAXOLONE

SUSPENSION;ORAL

ZTALMY

+! MARINUS

50MG/ML

N215904 001 Jun 01, 2022

GANCICLOVIR

GEL;OPHTHALMIC

ZIRGAN

+! BAUSCH AND LOMB

0.15%

N022211 001 Sep 15, 2009

SOLUTION;INTRAVENOUS

GANZYK-RTU

+! EXELA PHARMA

500MG/250ML (2MG/ML)

N209347 001 Feb 17, 2017

PRESCRIPTION DRUG PRODUCT LIST

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

GANCICLOVIR SODIUM

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A090658 001</u>	Jun 21, 2010
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A076222 001</u>	Jul 16, 2003
<u>AP</u>	PAR STERILE PRODUCTS	<u>EQ 500MG BASE/VIAL</u>	<u>A204950 001</u>	Dec 06, 2016
<u>AP</u>	! PHARMASCIENCE INC	<u>EQ 500MG BASE/VIAL</u>	<u>A207645 001</u>	Dec 08, 2017
<u>AP</u>	SLATE RUN PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A204204 001</u>	Nov 08, 2018

GANIRELIX ACETATE

INJECTABLE; INJECTION

FYREMADEL

<u>AP</u>	SUN PHARM	<u>250MCG/0.5ML</u>	<u>A204246 001</u>	Nov 30, 2018
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GANIRELIX ACETATE

<u>AP</u>	AMPHASTAR PHARMS INC	<u>250MCG/0.5ML</u>	<u>A212613 001</u>	Apr 07, 2022
<u>AP</u>	GLAND PHARMA LTD	<u>250MCG/0.5ML</u>	<u>A215658 001</u>	Feb 28, 2023
<u>AP</u>	LUPIN LTD	<u>250MCG/0.5ML</u>	<u>A216075 001</u>	Nov 16, 2023
<u>AP</u>	MEITHEAL	<u>250MCG/0.5ML</u>	<u>A214996 001</u>	Jun 06, 2022
<u>AP</u>	+! ORGANON USA ORGANON	<u>250MCG/0.5ML</u>	<u>N021057 001</u>	Jul 29, 1999

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

<u>AT</u>	HIKMA	<u>0.5%</u>	<u>A203189 001</u>	Sep 03, 2014
<u>AT</u>	LUPIN LTD	<u>0.5%</u>	<u>A202653 001</u>	Aug 28, 2013
<u>AT</u>	SANDOZ	<u>0.5%</u>	<u>A204227 001</u>	Jul 11, 2016

ZYMAXID

<u>AT</u>	+! ABBVIE	<u>0.5%</u>	<u>N022548 001</u>	May 18, 2010
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GEFITINIB

TABLET; ORAL

GEFITINIB

<u>AB</u>	ACTAVIS LABS FL INC	<u>250MG</u>	<u>A208913 001</u>	Apr 26, 2023
<u>AB</u>	APOTEX	<u>250MG</u>	<u>A209532 001</u>	Sep 23, 2022
<u>AB</u>	NATCO	<u>250MG</u>	<u>A212827 001</u>	May 31, 2023
<u>AB</u>	QILU PHARM HAINAN	<u>250MG</u>	<u>A211591 001</u>	Feb 13, 2023

IRESSA

<u>AB</u>	+! ASTRAZENECA	<u>250MG</u>	<u>N206995 001</u>	Jul 13, 2015
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GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 200MG BASE/VIAL</u>	<u>A091594 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A091594 002</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A091594 003</u>	Jul 25, 2011
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 200MG BASE/VIAL</u>	<u>A091365 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A091365 002</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A202997 001</u>	May 07, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A090799 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090799 002</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090242 003</u>	May 16, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090799 003</u>	May 16, 2011
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 200MG BASE/VIAL</u>	<u>A204520 001</u>	Jan 05, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A204520 002</u>	Jan 05, 2016
<u>AP</u>	HIKMA	<u>200MG/5.26ML (38MG/ML)</u>	<u>A213175 001</u>	Mar 07, 2023
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A213175 002</u>	Mar 07, 2023
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A213175 003</u>	Mar 07, 2023
<u>AP</u>	HIKMA INTL PHARMS	<u>EQ 200MG BASE/VIAL</u>	<u>A206617 001</u>	Jun 25, 2021
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A206617 002</u>	Jun 25, 2021
<u>AP</u>	HOSPIRA	<u>EQ 200MG BASE/VIAL</u>	<u>A078339 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A078339 002</u>	Jul 25, 2011
<u>AP</u>	+! HOSPIRA INC	<u>200MG/5.26ML (38MG/ML)</u>	<u>N200795 001</u>	Aug 04, 2011
<u>AP</u>	+!	<u>1GM/26.3ML (38MG/ML)</u>	<u>N200795 002</u>	Aug 04, 2011
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A079183 001</u>	Nov 15, 2010
<u>AP</u>	+!	<u>2GM/52.6ML (38MG/ML)</u>	<u>N200795 003</u>	Aug 04, 2011
<u>AP</u>	! JIANGSU HANSON PHARM	<u>EQ 200MG BASE/VIAL</u>	<u>A202485 001</u>	May 07, 2013
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A202485 002</u>	May 07, 2013
<u>AP</u>	MEITHEAL	<u>200MG/5.26ML (38MG/ML)</u>	<u>A212129 001</u>	Dec 11, 2020
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A212129 002</u>	Dec 11, 2020
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A212129 003</u>	Dec 11, 2020
<u>AP</u>	MYLAN LABS LTD	<u>200MG/5.26ML (38MG/ML)</u>	<u>A205242 001</u>	Dec 06, 2017
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A205242 002</u>	Dec 06, 2017

PRESCRIPTION DRUG PRODUCT LIST

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A205242 003</u>	Dec 06, 2017
<u>AP</u>	NOVAST LABS	<u>200MG/5.26ML (38MG/ML)</u>	<u>A210383 001</u>	Feb 14, 2019
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A210383 002</u>	Feb 14, 2019
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A210383 003</u>	Feb 14, 2019
<u>AP</u>	SAGENT PHARMS INC	<u>200MG/5.26ML (38MG/ML)</u>	<u>A209077 001</u>	Jul 20, 2018
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A209077 002</u>	Jul 20, 2018
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A209077 003</u>	Jul 20, 2018
<u>AP</u>	SHILPA	<u>EQ 200MG BASE/VIAL</u>	<u>A207575 001</u>	Feb 22, 2019
<u>AP</u>		<u>200MG/5.26ML (38MG/ML)</u>	<u>A210991 001</u>	Oct 04, 2019
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A207575 002</u>	Feb 22, 2019
<u>AP</u>		<u>1GM/26.3ML (38MG/ML)</u>	<u>A210991 002</u>	Oct 04, 2019
<u>AP</u>		<u>2GM/52.6ML (38MG/ML)</u>	<u>A210991 003</u>	Oct 04, 2019
<u>AP</u>	SUN PHARM	<u>EQ 200MG BASE/VIAL</u>	<u>A078433 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A078433 002</u>	Jul 25, 2011
<u>AP</u>	TEYRO LABS	<u>EQ 200MG BASE/VIAL</u>	<u>A078759 001</u>	Jul 25, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A078759 002</u>	Jul 25, 2011

SOLUTION; INTRAVENOUS

GEMCITABINE HYDROCHLORIDE

+	!	ACCORD HLTHCARE	1GM/10ML (100MG/ML)	N209604 002	Aug 03, 2017
+	!		1.5GM/15ML (100MG/ML)	N209604 003	Aug 03, 2017
+	!		2GM/20ML (100MG/ML)	N209604 004	Aug 03, 2017
+	!		200MG/2ML (100MG/ML)	N209604 001	Aug 03, 2017

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

<u>AB</u>	APOTEX	<u>600MG</u>	<u>A075034 001</u>	Jul 20, 1998		
<u>AB</u>	ASCENT PHARMS INC	<u>600MG</u>	<u>A214603 001</u>	Jan 13, 2021		
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A202726 001</u>	Sep 16, 2015		
<u>AB</u>	CADILA PHARMS LTD	<u>600MG</u>	<u>A203266 001</u>	Jun 17, 2016		
<u>AB</u>	CARIBE HOLDINGS	<u>600MG</u>	<u>A078012 001</u>	Mar 26, 2007		
<u>AB</u>	CHARTWELL MOLECULES	<u>600MG</u>	<u>A074270 001</u>	Sep 27, 1993		
<u>AB</u>	IMPAX PHARMS	<u>600MG</u>	<u>A078207 001</u>	Jun 01, 2007		
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A077836 001</u>	Jul 27, 2006		
<u>AB</u>	NORTHSTAR HLTHCARE	<u>600MG</u>	<u>A079072 001</u>	Sep 13, 2010		
<u>LOPID</u>						
<u>AB</u>	+	!	PFIZER PHARMS	<u>600MG</u>	<u>N018422 003</u>	Nov 20, 1986

GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN SULFATE

<u>AT</u>	COSETTE	<u>EQ 0.1% BASE</u>	<u>A064056 001</u>	Apr 29, 1994
<u>AT</u>	!	PADAGIS US	<u>EQ 0.1% BASE</u>	<u>A062307 001</u>

INJECTABLE; INJECTION

GENTAMICIN SULFATE

<u>AP</u>	EUGIA PHARMA	<u>EQ 10MG BASE/ML</u>	<u>A215236 001</u>	Jan 08, 2024	
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>A215237 001</u>	Jan 08, 2024	
<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 10MG BASE/ML</u>	<u>A062366 002</u>	Feb 06, 1986
<u>AP</u>	!		<u>EQ 40MG BASE/ML</u>	<u>A062366 001</u>	Aug 04, 1983
<u>AP</u>	HIKMA	<u>EQ 10MG BASE/ML</u>	<u>A062251 002</u>		
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>A062251 001</u>		
<u>AP</u>	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A062420 001</u>	Aug 15, 1983	
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>A062420 002</u>	Aug 15, 1983	

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 1.2MG BASE/ML</u>	<u>A062373 007</u>	Sep 07, 1982	
<u>AP</u>		<u>EQ 1.6MG BASE/ML</u>	<u>A062373 008</u>	Sep 07, 1982	
<u>AP</u>		<u>EQ 80MG BASE/100ML</u>	<u>A062373 002</u>	Sep 07, 1982	
<u>AP</u>		<u>EQ 100MG BASE/100ML</u>	<u>A062373 005</u>	Sep 07, 1982	
<u>AP</u>	HOSPIRA	<u>EQ 1.2MG BASE/ML</u>	<u>A062414 001</u>	Aug 15, 1983	
<u>AP</u>		<u>EQ 1.6MG BASE/ML</u>	<u>A062414 003</u>	Aug 15, 1983	
<u>AP</u>		<u>EQ 80MG BASE/100ML</u>	<u>A062414 008</u>	Aug 15, 1983	
<u>AP</u>		<u>EQ 100MG BASE/100ML</u>	<u>A062414 010</u>	Aug 15, 1983	
	!	BAXTER HLTHCARE	EQ 2MG BASE/ML	A062373 009	Sep 07, 1982
	!		EQ 120MG BASE/100ML	A062373 006	Sep 07, 1982

OINTMENT; OPHTHALMIC

GENTAMICIN SULFATE

!	!	FERA PHARMS LLC	EQ 0.3% BASE	A065024 001	Jul 30, 2004
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PRESCRIPTION DRUG PRODUCT LIST

GENTAMICIN SULFATE

OINTMENT; TOPICAL

GENTAMICIN SULFATE

AT	COSETTE	EQ 0.1% BASE	A064054 001	Apr 29, 1994
AT	FOUGERA PHARMS INC	EQ 0.1% BASE	A062533 001	Oct 05, 1984
AT	! PADAGIS US	EQ 0.1% BASE	A062351 001	Feb 18, 1982
AT	TARO	EQ 0.1% BASE	A062477 001	Dec 23, 1983

SOLUTION/DROPS; OPHTHALMIC

GENTAMICIN SULFATE

AT	BAUSCH AND LOMB	EQ 0.3% BASE	A064048 001	May 11, 1994
AT	PADAGIS US	EQ 0.3% BASE	A065121 001	Jan 30, 2004
AT	! SANDOZ	EQ 0.3% BASE	A062196 001	

GEPİRONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EXXUA

+	FABRE KRAMER	EQ 18.2MG BASE	N021164 001	Sep 22, 2023
+		EQ 36.3MG BASE	N021164 002	Sep 22, 2023
+		EQ 54.5MG BASE	N021164 003	Sep 22, 2023
+	!	EQ 72.6MG BASE	N021164 004	Sep 22, 2023

GILTERITINIB FUMARATE

TABLET; ORAL

XOSPATA

+	! ASTELLAS	EQ 40MG BASE	N211349 001	Nov 28, 2018
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GIVOSIRAN SODIUM

SOLUTION; SUBCUTANEOUS

GIVLAARI

+	! ALNYLAM PHARMS INC	EQ 189MG BASE/ML (EQ 189MG BASE/ML)	N212194 001	Nov 20, 2019
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GLASDEGIB MALEATE

TABLET; ORAL

DAURISMO

+	PFIZER	EQ 25MG BASE	N210656 001	Nov 21, 2018
+	!	EQ 100MG BASE	N210656 002	Nov 21, 2018

GLATIRAMER ACETATE

INJECTABLE; SUBCUTANEOUS

COPAXONE

AP	! TEVA PHARMS USA	20MG/ML	N020622 002	Feb 12, 2002
AP	!	40MG/ML	N020622 003	Jan 28, 2014

GLATIRAMER ACETATE

AP	MYLAN	20MG/ML	A091646 001	Oct 03, 2017
AP		40MG/ML	A206936 001	Oct 03, 2017

GLATOPIA

AP	SANDOZ	20MG/ML	A090218 001	Apr 16, 2015
AP		40MG/ML	A206921 001	Feb 12, 2018

GLECAPREVIR; PIBRENTASVIR

PELLETS; ORAL

MAVYRET

+	! ABBVIE	50MG; 20MG/PACKET	N215110 001	Jun 10, 2021
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TABLET; ORAL

MAVYRET

+	! ABBVIE	100MG; 40MG	N209394 001	Aug 03, 2017
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GLIMEPIRIDE

TABLET; ORAL

AMARYL

AB	! SANOFI AVENTIS US	1MG	N020496 001	Nov 30, 1995
AB	+	2MG	N020496 002	Nov 30, 1995
AB	+	4MG	N020496 003	Nov 30, 1995

GLIMEPIRIDE

AB	ACCORD HLTHCARE	1MG	A078181 001	Aug 23, 2007
AB		2MG	A078181 002	Aug 23, 2007
AB		4MG	A078181 003	Aug 23, 2007
AB	AUROBINDO PHARMA LTD	1MG	A202759 001	Jun 29, 2012
AB		2MG	A202759 002	Jun 29, 2012
AB		4MG	A202759 003	Jun 29, 2012
AB	CARLSBAD	1MG	A077911 001	Sep 22, 2009
AB		2MG	A077911 002	Sep 22, 2009
AB		4MG	A077911 003	Sep 22, 2009
AB	CHARTWELL MOLECULAR	1MG	A077295 001	Oct 06, 2005
AB		2MG	A077295 002	Oct 06, 2005

PRESCRIPTION DRUG PRODUCT LIST

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

<u>AB</u>		<u>4MG</u>	<u>A077295</u>	<u>003</u>	Oct 06, 2005
<u>AB</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A077091</u>	<u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A077091</u>	<u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A077091</u>	<u>003</u>	Oct 06, 2005
<u>AB</u>	INDOCO REMEDIES	<u>1MG</u>	<u>A202112</u>	<u>001</u>	Apr 17, 2013
<u>AB</u>		<u>2MG</u>	<u>A202112</u>	<u>002</u>	Apr 17, 2013
<u>AB</u>		<u>4MG</u>	<u>A202112</u>	<u>003</u>	Apr 17, 2013
<u>AB</u>	MICRO LABS	<u>1MG</u>	<u>A091220</u>	<u>001</u>	Jun 29, 2012
<u>AB</u>		<u>2MG</u>	<u>A091220</u>	<u>002</u>	Jun 29, 2012
<u>AB</u>		<u>4MG</u>	<u>A091220</u>	<u>004</u>	Jun 29, 2012
<u>AB</u>		<u>8MG</u>	<u>A091220</u>	<u>006</u>	Jun 29, 2012
<u>AB</u>	PRINSTON INC	<u>1MG</u>	<u>A077370</u>	<u>001</u>	Dec 23, 2005
<u>AB</u>		<u>2MG</u>	<u>A077370</u>	<u>002</u>	Dec 23, 2005
<u>AB</u>		<u>4MG</u>	<u>A077370</u>	<u>003</u>	Dec 23, 2005
<u>AB</u>		<u>8MG</u>	<u>A077370</u>	<u>004</u>	Dec 23, 2005
	MICRO LABS	3MG	A091220	003	Jun 29, 2012
		6MG	A091220	005	Jun 29, 2012

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACT

<u>AB</u>	+	TAKEDA PHARMS USA	<u>2MG;30MG</u>	<u>N021925</u>	<u>001</u>	Jul 28, 2006
<u>AB</u>	+		<u>4MG;30MG</u>	<u>N021925</u>	<u>002</u>	Jul 28, 2006

PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE

<u>AB</u>		SANDOZ	<u>2MG;30MG</u>	<u>A201049</u>	<u>001</u>	Jan 04, 2013
<u>AB</u>			<u>4MG;30MG</u>	<u>A201049</u>	<u>002</u>	Jan 04, 2013

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A074550</u>	<u>001</u>	Sep 11, 1997
<u>AB</u>			<u>10MG</u>	<u>A074550</u>	<u>002</u>	Sep 11, 1997
<u>AB</u>		ANI PHARMS	<u>5MG</u>	<u>A074497</u>	<u>001</u>	Aug 31, 1995
<u>AB</u>			<u>10MG</u>	<u>A074497</u>	<u>002</u>	Aug 31, 1995
<u>AB</u>		APOTEX	<u>5MG</u>	<u>A075795</u>	<u>001</u>	Jun 13, 2001
<u>AB</u>	!		<u>10MG</u>	<u>A075795</u>	<u>002</u>	Jun 13, 2001
<u>AB</u>		AUROBINDO PHARMA USA	<u>5MG</u>	<u>A074226</u>	<u>001</u>	May 10, 1994
<u>AB</u>			<u>10MG</u>	<u>A074226</u>	<u>002</u>	May 10, 1994
<u>AB</u>		RUBICON	<u>5MG</u>	<u>A214874</u>	<u>002</u>	Oct 03, 2023
<u>AB</u>			<u>10MG</u>	<u>A214874</u>	<u>003</u>	Oct 03, 2023
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A074305</u>	<u>001</u>	Apr 07, 1995
<u>AB</u>			<u>10MG</u>	<u>A074305</u>	<u>002</u>	Apr 07, 1995
<u>AB</u>		WATSON LABS TEVA	<u>5MG</u>	<u>A074223</u>	<u>001</u>	Feb 27, 1995
<u>AB</u>			<u>10MG</u>	<u>A074223</u>	<u>002</u>	Feb 27, 1995
		RUBICON	2.5MG	A214874	001	Oct 03, 2023

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

<u>AB</u>		AUROBINDO PHARMA	<u>2.5MG</u>	<u>A206928</u>	<u>001</u>	May 12, 2017
<u>AB</u>			<u>5MG</u>	<u>A206928</u>	<u>002</u>	May 12, 2017
<u>AB</u>			<u>10MG</u>	<u>A206928</u>	<u>003</u>	May 12, 2017
<u>AB</u>		UNIQUE	<u>2.5MG</u>	<u>A204720</u>	<u>001</u>	Dec 29, 2016
<u>AB</u>			<u>5MG</u>	<u>A204720</u>	<u>002</u>	Dec 29, 2016
<u>AB</u>			<u>10MG</u>	<u>A204720</u>	<u>003</u>	Dec 29, 2016
<u>AB</u>		WATSON LABS	<u>2.5MG</u>	<u>A076467</u>	<u>003</u>	Mar 27, 2006
<u>AB</u>			<u>5MG</u>	<u>A076467</u>	<u>001</u>	Sep 08, 2003
<u>AB</u>			<u>10MG</u>	<u>A076467</u>	<u>002</u>	Nov 07, 2003
<u>AB</u>		ZYDUS PHARMS	<u>2.5MG</u>	<u>A203499</u>	<u>001</u>	Jul 16, 2018
<u>AB</u>			<u>5MG</u>	<u>A203499</u>	<u>002</u>	Jul 16, 2018
<u>AB</u>			<u>10MG</u>	<u>A203499</u>	<u>003</u>	Jul 16, 2018

GLUCOTROL XL

<u>AB</u>	+	PFIZER	<u>2.5MG</u>	<u>N020329</u>	<u>003</u>	Aug 10, 1999
<u>AB</u>	+		<u>5MG</u>	<u>N020329</u>	<u>001</u>	Apr 26, 1994
<u>AB</u>	+		<u>10MG</u>	<u>N020329</u>	<u>002</u>	Apr 26, 1994

PRESCRIPTION DRUG PRODUCT LIST

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

AB	EPIC PHARMA LLC	<u>2.5MG;250MG</u>	<u>A077507 001</u>	Oct 27, 2005
AB		<u>2.5MG;500MG</u>	<u>A077507 002</u>	Oct 27, 2005
AB		<u>5MG;500MG</u>	<u>A077507 003</u>	Oct 27, 2005
AB	HERITAGE PHARMS	<u>2.5MG;250MG</u>	<u>A078728 001</u>	Jun 23, 2010
AB		<u>2.5MG;500MG</u>	<u>A078728 002</u>	Jun 23, 2010
AB		<u>5MG;500MG</u>	<u>A078728 003</u>	Jun 23, 2010
AB	TEVA PHARMS	<u>2.5MG;250MG</u>	<u>A077270 001</u>	Oct 28, 2005
AB		<u>2.5MG;500MG</u>	<u>A077270 002</u>	Oct 28, 2005
AB	!	<u>5MG;500MG</u>	<u>A077270 003</u>	Oct 28, 2005
AB	ZYDUS PHARMS USA INC	<u>2.5MG;250MG</u>	<u>A078905 001</u>	Jan 31, 2011
AB		<u>2.5MG;500MG</u>	<u>A078905 002</u>	Jan 31, 2011
AB		<u>5MG;500MG</u>	<u>A078905 003</u>	Jan 31, 2011

GLUCAGON

INJECTABLE; INJECTION

GLUCAGON

!	AMPHASTAR PHARMS INC	1MG/VIAL	A208086 001	Dec 28, 2020
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POWDER; NASAL

BAQSIMI

+	AMPHASTAR PHARMS INC	3MG	N210134 001	Jul 24, 2019
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SOLUTION; SUBCUTANEOUS

GVOKE HYPOPEN

+	XERIS	0.5MG/0.1ML (0.5MG/0.1ML)	N212097 003	Sep 10, 2019
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+	XERIS	1MG/0.2ML (1MG/0.2ML)	N212097 004	Sep 10, 2019
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GVOKE KIT

+	XERIS	1MG/0.2ML (1MG/0.2ML)	N212097 005	Aug 20, 2021
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GVOKE PFS

+	XERIS	1MG/0.2ML (1MG/0.2ML)	N212097 002	Sep 10, 2019
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GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGEN

+	NOVO NORDISK	EQ 1MG BASE/VIAL	N020918 001	Jun 22, 1998
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POWDER; INTRAMUSCULAR, INTRAVENOUS

GLUCAGON

+	FRESENIUS KABI USA	EQ 1MG BASE/VIAL	N201849 001	May 08, 2015
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GLYBURIDE

TABLET; ORAL

GLYBURIDE (MICRONIZED)

AB	TEVA	<u>1.5MG</u>	<u>A074686 001</u>	Apr 20, 1999
AB		<u>3MG</u>	<u>A074686 002</u>	Apr 20, 1999
AB		<u>6MG</u>	<u>A074686 004</u>	Apr 20, 1999

GLYNASE

AB	+	PFIZER	<u>1.5MG</u>	<u>N020051 001</u>	Mar 04, 1992
AB	+		<u>3MG</u>	<u>N020051 002</u>	Mar 04, 1992
AB	+		<u>6MG</u>	<u>N020051 004</u>	Sep 24, 1993

GLYBURIDE

AB1	CADILA PHARMS LTD	<u>1.25MG</u>	<u>A203379 001</u>	Jan 04, 2019
AB1		<u>2.5MG</u>	<u>A203379 002</u>	Jan 04, 2019
AB1		<u>5MG</u>	<u>A203379 003</u>	Jan 04, 2019
AB1	EPIC PHARMA LLC	<u>1.25MG</u>	<u>A076257 001</u>	Jun 27, 2002
AB1		<u>2.5MG</u>	<u>A076257 002</u>	Jun 27, 2002
AB1		<u>5MG</u>	<u>A076257 003</u>	Jun 27, 2002
AB1	HERITAGE PHARMS	<u>1.25MG</u>	<u>A090937 001</u>	Feb 28, 2011
AB1		<u>2.5MG</u>	<u>A090937 002</u>	Feb 28, 2011
AB1		<u>5MG</u>	<u>A090937 003</u>	Feb 28, 2011
AB1	ORIENT PHARMA CO LTD	<u>1.25MG</u>	<u>A206483 001</u>	Feb 22, 2019
AB1		<u>2.5MG</u>	<u>A206483 002</u>	Feb 22, 2019
AB1		<u>5MG</u>	<u>A206483 003</u>	Feb 22, 2019
AB1	TEVA	<u>1.25MG</u>	<u>A074388 001</u>	Aug 29, 1995
AB1		<u>2.5MG</u>	<u>A074388 002</u>	Aug 29, 1995
AB1	!	<u>5MG</u>	<u>A074388 003</u>	Aug 29, 1995
AB1	ZYDUS PHARMS	<u>1.25MG</u>	<u>A206749 001</u>	May 10, 2016
AB1		<u>2.5MG</u>	<u>A206749 002</u>	May 10, 2016
AB1		<u>5MG</u>	<u>A206749 003</u>	May 10, 2016

PRESCRIPTION DRUG PRODUCT LIST

GLYBURIDE

TABLET; ORAL

DIABETA

AB2	+	SANOFI AVENTIS US	<u>1.25MG</u>	<u>N017532</u>	<u>001</u>	May 01, 1984
AB2	+		<u>2.5MG</u>	<u>N017532</u>	<u>002</u>	May 01, 1984
AB2	+	!	<u>5MG</u>	<u>N017532</u>	<u>003</u>	May 01, 1984

GLYBURIDE

AB2		IMPAX LABS INC	<u>1.25MG</u>	<u>A206079</u>	<u>001</u>	Sep 30, 2015
AB2			<u>2.5MG</u>	<u>A206079</u>	<u>002</u>	Sep 30, 2015
AB2			<u>5MG</u>	<u>A206079</u>	<u>003</u>	Sep 30, 2015
		GLYBURIDE (MICRONIZED)				
		TEVA	4.5MG	A074686	003	Apr 20, 1999

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	<u>1.25MG;250MG</u>	<u>A076716</u>	<u>001</u>	Jun 28, 2005
AB			<u>2.5MG;500MG</u>	<u>A076716</u>	<u>002</u>	Jun 28, 2005
AB			<u>5MG;500MG</u>	<u>A076716</u>	<u>003</u>	Jun 28, 2005
AB		AUROBINDO PHARMA	<u>1.25MG;250MG</u>	<u>A077870</u>	<u>001</u>	Nov 14, 2007
AB		!	<u>2.5MG;500MG</u>	<u>A077870</u>	<u>002</u>	Nov 14, 2007
AB			<u>5MG;500MG</u>	<u>A077870</u>	<u>003</u>	Nov 14, 2007
AB		IMPAX LABS INC	<u>1.25MG;250MG</u>	<u>A076345</u>	<u>001</u>	Feb 18, 2004
AB			<u>2.5MG;500MG</u>	<u>A076345</u>	<u>002</u>	Feb 18, 2004
AB			<u>5MG;500MG</u>	<u>A076345</u>	<u>003</u>	Feb 18, 2004
AB		ZYDUS PHARMS	<u>1.25MG;250MG</u>	<u>A206748</u>	<u>001</u>	Feb 29, 2016
AB			<u>2.5MG;500MG</u>	<u>A206748</u>	<u>002</u>	Feb 29, 2016
AB			<u>5MG;500MG</u>	<u>A206748</u>	<u>003</u>	Feb 29, 2016

GLYCEROL PHENYLBUTYRATE

LIQUID; ORAL

RAVICTI

+	!	HORIZON THERAP US	1.1GM/ML	N203284	001	Feb 01, 2013
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GLYCINE

SOLUTION; IRRIGATION

AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER

AT	+	BAXTER HLTHCARE	<u>1.5GM/100ML</u>	<u>N017865</u>	<u>001</u>	
AT		B BRAUN	<u>1.5GM/100ML</u>	<u>N016784</u>	<u>001</u>	
AT		ICU MEDICAL INC	<u>1.5GM/100ML</u>	<u>N018315</u>	<u>001</u>	

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

AP		ACCORD HLTHCARE	<u>0.2MG/ML</u>	<u>A213238</u>	<u>001</u>	Jul 08, 2020
AP		ALEMBIC	<u>0.2MG/ML</u>	<u>A214635</u>	<u>001</u>	Oct 26, 2022
AP		AM REGENT	<u>0.2MG/ML</u>	<u>A089335</u>	<u>001</u>	Jul 23, 1986
AP		AMNEAL	<u>0.2MG/ML</u>	<u>A208973</u>	<u>001</u>	Jun 15, 2017
AP			<u>0.2MG/ML</u>	<u>A215333</u>	<u>001</u>	Oct 21, 2022
AP		APOTEX	<u>0.2MG/ML</u>	<u>A210246</u>	<u>001</u>	Oct 29, 2019
AP		ASPEN	<u>0.2MG/ML</u>	<u>A212871</u>	<u>001</u>	Nov 29, 2022
AP		CAPLIN	<u>0.2MG/ML</u>	<u>A211705</u>	<u>001</u>	Mar 20, 2019
AP		FRESENIUS KABI USA	<u>0.2MG/ML</u>	<u>A209024</u>	<u>001</u>	Oct 31, 2018
AP			<u>0.2MG/ML</u>	<u>A209328</u>	<u>001</u>	Oct 27, 2017
AP		GLAND PHARMA LTD	<u>0.2MG/ML</u>	<u>A212612</u>	<u>001</u>	Sep 30, 2019
AP	!	HIKMA FARMACEUTICA	<u>0.2MG/ML</u>	<u>A090963</u>	<u>001</u>	Sep 21, 2011
AP		MEITHEAL	<u>0.2MG/ML</u>	<u>A212802</u>	<u>001</u>	Jul 06, 2021
AP		PIRAMAL CRITICAL	<u>0.2MG/ML</u>	<u>A210842</u>	<u>001</u>	Oct 25, 2018
AP		PRINSTON INC	<u>0.2MG/ML</u>	<u>A210927</u>	<u>001</u>	Oct 31, 2018
AP		SAGENT	<u>0.2MG/ML</u>	<u>A210083</u>	<u>001</u>	Feb 21, 2020
AP		SANDOZ	<u>0.2MG/ML</u>	<u>A211334</u>	<u>001</u>	May 14, 2019
AP		SOMERSET THERAPS	<u>0.2MG/ML</u>	<u>A207639</u>	<u>001</u>	Jun 23, 2017
		LLC				
AP		UMEDICA	<u>0.2MG/ML</u>	<u>A212591</u>	<u>001</u>	Oct 13, 2021
AP		XIROMED	<u>0.2MG/ML</u>	<u>A212227</u>	<u>001</u>	Mar 04, 2021

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

GLYCOPYRROLATE

+	!	FRESENIUS KABI USA	0.6MG/3ML (0.2MG/ML)	N214919	001	Apr 21, 2022
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GLYRX-PF

+	!	EXELA PHARMA	0.2MG/ML (0.2MG/ML)	N210997	001	Jul 11, 2018
+	!		0.4MG/2ML (0.2MG/ML)	N210997	002	Jul 11, 2018
+	!		0.6MG/3ML (0.2MG/ML)	N210997	004	Dec 14, 2020
+	!		1MG/5ML (0.2MG/ML)	N210997	003	Apr 09, 2020

PRESCRIPTION DRUG PRODUCT LIST

GLYCOPYRROLATE

SOLUTION; ORAL

CUVPOSA

<u>AA</u>	<u>+</u> !	MERZ PHARMS	<u>1MG/5ML</u>	<u>N022571</u>	<u>001</u>	Jul 28, 2010
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GLYCOPYRROLATE

<u>AA</u>		ANNORA PHARMA	<u>1MG/5ML</u>	<u>A213698</u>	<u>001</u>	Jul 05, 2022
<u>AA</u>		PAR PHARM INC	<u>1MG/5ML</u>	<u>A204438</u>	<u>001</u>	Aug 09, 2021
<u>AA</u>		SUVEN PHARMS	<u>1MG/5ML</u>	<u>A212467</u>	<u>001</u>	Jul 05, 2022

TABLET; ORAL

GLYCOPYRROLATE

<u>AA</u>		ALEMBIC	<u>1MG</u>	<u>A203657</u>	<u>001</u>	Nov 30, 2018
<u>AA</u>			<u>2MG</u>	<u>A203657</u>	<u>002</u>	Nov 30, 2018
<u>AA</u>		AUROBINDO PHARMA	<u>1MG</u>	<u>A202675</u>	<u>001</u>	Apr 15, 2013
<u>AA</u>			<u>2MG</u>	<u>A202675</u>	<u>002</u>	Oct 30, 2018
<u>AA</u>		DR REDDYS LABS LTD	<u>1MG</u>	<u>A040847</u>	<u>001</u>	Mar 21, 2008
<u>AA</u>			<u>2MG</u>	<u>A040847</u>	<u>002</u>	Mar 21, 2008
<u>AA</u>		HERITAGE PHARMS INC	<u>1MG</u>	<u>A207201</u>	<u>001</u>	Jan 03, 2017
<u>AA</u>			<u>2MG</u>	<u>A207201</u>	<u>002</u>	Jan 03, 2017
<u>AA</u>		INDOCO	<u>1MG</u>	<u>A091182</u>	<u>001</u>	Feb 03, 2014
<u>AA</u>			<u>2MG</u>	<u>A091182</u>	<u>002</u>	Feb 03, 2014
<u>AA</u>		LEADING	<u>1MG</u>	<u>A090195</u>	<u>001</u>	Sep 21, 2012
<u>AA</u>			<u>2MG</u>	<u>A090195</u>	<u>002</u>	Sep 21, 2012
<u>AA</u>		NATCO	<u>1MG</u>	<u>A091413</u>	<u>001</u>	Jun 20, 2016
<u>AA</u>			<u>2MG</u>	<u>A091413</u>	<u>002</u>	Jun 20, 2016
<u>AA</u>		OXFORD PHARMS	<u>1MG</u>	<u>A090020</u>	<u>001</u>	Oct 19, 2011
<u>AA</u>			<u>2MG</u>	<u>A090020</u>	<u>002</u>	Oct 19, 2011
<u>AA</u>	<u>!</u>	PAR PHARM	<u>1MG</u>	<u>A040653</u>	<u>001</u>	Aug 31, 2006
<u>AA</u>	<u>!</u>		<u>2MG</u>	<u>A040653</u>	<u>002</u>	Aug 31, 2006
<u>AA</u>		RISING	<u>1MG</u>	<u>A040821</u>	<u>001</u>	Dec 29, 2008
<u>AA</u>			<u>2MG</u>	<u>A040821</u>	<u>002</u>	Dec 29, 2008
<u>AA</u>		SUN PHARM INDS LTD	<u>1MG</u>	<u>A040844</u>	<u>001</u>	Aug 18, 2009
<u>AA</u>			<u>2MG</u>	<u>A040844</u>	<u>002</u>	Aug 18, 2009

ROBINUL

<u>AA</u>	<u>+</u>	CASPER PHARMA LLC	<u>1MG</u>	<u>N012827</u>	<u>001</u>	
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ROBINUL FORTE

<u>AA</u>	<u>+</u>	CASPER PHARMA LLC	<u>2MG</u>	<u>N012827</u>	<u>002</u>	
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GLYCOPYRROLATE

		LGM PHARMA	1.5MG	A091522	001	Mar 12, 2012
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GLYCOPYRROLATE; NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

PREVDUO

<u>+</u> !	SLAYBACK PHARMA LLC	0.6MG/3ML (0.2MG/ML); 3MG/3ML (1MG/ML)	N216903	001	Feb 23, 2023
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GLYCOPYRROLATE; NEOSTIGMINE METHYLSULFATE

CLOTH; TOPICAL

QBREXZA

<u>+</u> !	JOURNEY	EQ 2.4% BASE	N210361	001	Jun 28, 2018
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GOLODIRSEN

SOLUTION; INTRAVENOUS

VYONDYS 53

<u>+</u> !	SAREPTA THERAPS INC	100MG/2ML (50MG/ML)	N211970	001	Dec 12, 2019
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GOSERELIN ACETATE

IMPLANT; IMPLANTATION

ZOLADEX

<u>+</u> !	TERSERA	EQ 3.6MG BASE	N019726	001	Dec 29, 1989
<u>+</u> !		EQ 10.8MG BASE	N020578	001	Jan 11, 1996

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

<u>AT</u>		AMRING PHARMS	<u>0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML</u>	<u>A065187</u>	<u>001</u>	Oct 28, 2005
<u>AT</u>	<u>!</u>	BAUSCH AND LOMB	<u>0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML</u>	<u>A064047</u>	<u>001</u>	Jan 31, 1996

NEOSPORIN

<u>AT</u>	<u>!</u>	MONARCH PHARMS	<u>0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML</u>	<u>A060582</u>	<u>001</u>	
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PRESCRIPTION DRUG PRODUCT LIST

GRANISETRON

FILM, EXTENDED RELEASE;TRANSDERMAL

SANCUSO

+! CUMBERLAND

3.1MG/24HR

N022198 001 Sep 12, 2008

INJECTION, EXTENDED RELEASE;SUBCUTANEOUS

SUSTOL

+! HERON THERAPS INC

10MG/0.4ML (10MG/0.4ML)

N022445 001 Aug 09, 2016

GRANISETRON HYDROCHLORIDE

INJECTABLE;INJECTION

GRANISETRON HYDROCHLORIDE

<u>AP</u>	AMNEAL	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078262 001</u>	Dec 31, 2007
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078258 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078258 002</u>	Jun 30, 2008
<u>AP</u>	BIONPHARMA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078863 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078880 001</u>	Jun 30, 2008
<u>AP</u>	! DR REDDYS	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078392 001</u>	Dec 31, 2007
<u>AP</u>	!	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077297 001</u>	Jun 30, 2008
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078522 001</u>	Dec 31, 2007
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078090 001</u>	Jun 30, 2008
<u>AP</u>	HIKMA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A077913 001</u>	Jun 26, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077186 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A077187 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077177 001</u>	Dec 31, 2007
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078629 001</u>	Dec 23, 2009
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078629 002</u>	Dec 23, 2009
<u>AP</u>	MYLAN ASI	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A091136 001</u>	Apr 09, 2010
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A091136 002</u>	Apr 09, 2010
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A091137 002</u>	Apr 09, 2010
<u>AP</u>	RISING	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A079078 001</u>	Sep 14, 2009
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A079078 002</u>	Sep 14, 2009
<u>AP</u>	SANDOZ	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078534 001</u>	Apr 30, 2009
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078531 001</u>	Apr 30, 2009
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078531 002</u>	Apr 30, 2009
<u>AP</u>	SANDOZ INC	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078835 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078835 002</u>	Jun 30, 2008

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	BIONPHARMA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078863 002</u>	Jun 30, 2008
<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078096 001</u>	Jun 30, 2008

TABLET;ORAL

GRANISETRON HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>EQ 1MG BASE</u>	<u>A078843 001</u>	Feb 27, 2008
<u>AB</u>	CHARTWELL MOLECULAR	<u>EQ 1MG BASE</u>	<u>A078037 001</u>	Feb 27, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 1MG BASE</u>	<u>A078846 001</u>	Feb 27, 2009
<u>AB</u>	NATCO PHARMA	<u>EQ 1MG BASE</u>	<u>A078969 001</u>	Jun 22, 2009
<u>AB</u>	! ORBION PHARMS	<u>EQ 1MG BASE</u>	<u>A078678 001</u>	Feb 13, 2008
<u>AB</u>	TARO	<u>EQ 1MG BASE</u>	<u>A090817 001</u>	May 28, 2010

GRISEOFULVIN, MICROCRYSTALLINE

TABLET;ORAL

FULVICIN-U/F

CHARTWELL RX

250MG

A060569 002

500MG

A060569 001

GRISEOFULVIN, MICROSIZE

SUSPENSION;ORAL

GRISEOFULVIN

<u>AB</u>	! ACTAVIS MID	<u>125MG/5ML</u>	<u>A065394 001</u>	Jul 06, 2007
	ATLANTIC			
<u>AB</u>	CHARTWELL RX	<u>125MG/5ML</u>	<u>A065200 001</u>	Mar 02, 2005
<u>AB</u>	CIPLA	<u>125MG/5ML</u>	<u>A065354 001</u>	Sep 10, 2007
<u>AB</u>	COSETTE	<u>125MG/5ML</u>	<u>A065438 001</u>	Oct 08, 2010

TABLET;ORAL

GRISEOFULVIN

<u>AB</u>	! SANDOZ	<u>500MG</u>	<u>A091592 002</u>	Aug 07, 2013
<u>AB</u>	SIGMAPHARM LABS LLC	<u>500MG</u>	<u>A202482 001</u>	Oct 22, 2012
	SANDOZ	250MG	A091592 001	Aug 07, 2013

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET;ORAL

FULVICIN P/G

CHARTWELL RX

125MG

A061996 001

250MG

A061996 002

PRESCRIPTION DRUG PRODUCT LIST

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET;ORAL

FULVICIN P/G 165					
CHARTWELL RX	165MG		A061996	003	Apr 06, 1982
FULVICIN P/G 330					
CHARTWELL RX	330MG		A061996	004	Apr 06, 1982

GRISEOFULVIN, ULTRAMICROSIZED

TABLET;ORAL

GRIS-PEG

AB	+	BAUSCH	125MG	N050475	001	
AB	+		250MG	N050475	002	
<u>GRISEOFULVIN, ULTRAMICROSIZED</u>						
AB		MOUNTAIN	125MG	A204371	001	Jan 09, 2014
AB	!		250MG	A204371	002	Jan 09, 2014
AB		SANDOZ	125MG	A202805	001	Dec 26, 2018
AB			250MG	A202805	002	Dec 26, 2018
<u>GRISEOFULVIN, ULTRAMICROSIZED</u>						
AB		SIGMAPHARM LABS LLC	125MG	A202545	001	Oct 22, 2012
AB			250MG	A202545	002	Oct 22, 2012

GUANFACINE HYDROCHLORIDE

TABLET;ORAL

GUANFACINE HYDROCHLORIDE

AB		AJANTA PHARMA LTD	EQ 1MG BASE	A217293	001	Apr 27, 2023
AB			EQ 2MG BASE	A217293	002	Apr 27, 2023
AB		AMNEAL PHARM	EQ 1MG BASE	A075109	001	Nov 25, 1998
AB	!		EQ 2MG BASE	A075109	002	Nov 25, 1998
AB		AUROBINDO PHARMA USA	EQ 1MG BASE	A074796	001	Jan 27, 1997
AB			EQ 2MG BASE	A074796	002	Jan 27, 1997
AB		EPIC PHARMA LLC	EQ 1MG BASE	A074673	001	Feb 28, 1997
AB			EQ 2MG BASE	A074673	002	Feb 28, 1997
AB		I 3 PHARMS	EQ 1MG BASE	A216828	001	Oct 05, 2023
AB			EQ 2MG BASE	A216828	002	Oct 05, 2023
AB		RUBICON	EQ 1MG BASE	A216762	001	Oct 17, 2023
AB			EQ 2MG BASE	A216762	002	Oct 17, 2023
AB		TWI PHARMS	EQ 1MG BASE	A216399	001	Jun 08, 2022
AB			EQ 2MG BASE	A216399	002	Jun 08, 2022
AB		UNICHEM	EQ 1MG BASE	A214689	001	Mar 03, 2021
AB			EQ 2MG BASE	A214689	002	Mar 03, 2021
AB		WATSON LABS	EQ 1MG BASE	A074145	001	Oct 17, 1995
AB			EQ 2MG BASE	A074145	002	Oct 17, 1995

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	EQ 1MG BASE	A200881	001	Oct 05, 2012
AB			EQ 2MG BASE	A200881	002	Oct 05, 2012
AB			EQ 3MG BASE	A200881	003	Oct 05, 2012
AB			EQ 4MG BASE	A200881	004	Oct 05, 2012
AB		ALEMBIC	EQ 1MG BASE	A217269	001	Aug 07, 2023
AB			EQ 2MG BASE	A217269	002	Aug 07, 2023
AB			EQ 3MG BASE	A217269	003	Aug 07, 2023
AB			EQ 4MG BASE	A217269	004	Aug 07, 2023
AB		APOTEX	EQ 1MG BASE	A205430	001	Jul 25, 2018
AB			EQ 2MG BASE	A205430	002	Jul 25, 2018
AB			EQ 3MG BASE	A205430	003	Jul 25, 2018
AB			EQ 4MG BASE	A205430	004	Jul 25, 2018
AB		SANDOZ	EQ 1MG BASE	A202568	001	Jun 03, 2015
AB			EQ 2MG BASE	A202568	002	Jun 03, 2015
AB			EQ 3MG BASE	A202568	003	Jun 03, 2015
AB			EQ 4MG BASE	A202568	004	Jun 03, 2015
AB		SUN PHARM	EQ 1MG BASE	A205689	001	Nov 16, 2017
AB			EQ 2MG BASE	A205689	002	Nov 16, 2017
AB			EQ 3MG BASE	A205689	003	Nov 16, 2017
AB			EQ 4MG BASE	A205689	004	Nov 16, 2017
AB		TEVA PHARMS USA	EQ 1MG BASE	A201382	001	Jun 02, 2015
AB			EQ 2MG BASE	A201382	002	Jun 02, 2015
AB			EQ 3MG BASE	A201382	003	Jun 02, 2015
AB			EQ 4MG BASE	A201382	004	Jun 02, 2015
AB		TWI PHARMS	EQ 1MG BASE	A201408	001	Jun 02, 2015
AB			EQ 2MG BASE	A201408	002	Jun 02, 2015
AB			EQ 3MG BASE	A201408	003	Jun 02, 2015
AB			EQ 4MG BASE	A201408	004	Jun 02, 2015
AB		YICHANG HUMANWELL	EQ 1MG BASE	A213428	001	Nov 25, 2020

PRESCRIPTION DRUG PRODUCT LIST

GUANFACINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A213428 002</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A213428 003</u>	Nov 25, 2020
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A213428 004</u>	Nov 25, 2020
<u>INTUNIV</u>				
<u>AB</u>	+	TAKEDA PHARMS USA	<u>EQ 1MG BASE</u>	<u>N022037 001</u> Sep 02, 2009
<u>AB</u>	+		<u>EQ 2MG BASE</u>	<u>N022037 002</u> Sep 02, 2009
<u>AB</u>	+		<u>EQ 3MG BASE</u>	<u>N022037 003</u> Sep 02, 2009
<u>AB</u>	+	!	<u>EQ 4MG BASE</u>	<u>N022037 004</u> Sep 02, 2009

HALCINONIDE

CREAM;TOPICAL

HALCINONIDE

<u>AB</u>		CHARTWELL RX	<u>0.1%</u>	<u>A214723 001</u> Sep 08, 2021
<u>AB</u>		MYLAN	<u>0.1%</u>	<u>A211027 001</u> Aug 12, 2019
<u>HALOG</u>				
<u>AB</u>	+	!	SUN PHARM INDS INC	<u>0.1%</u> <u>N017556 001</u>
OINTMENT;TOPICAL				
HALOG				
	+	!	SUN PHARM INDS INC	0.1% N017824 001
SOLUTION;TOPICAL				
HALOG				
	+	!	SUN PHARM INDS INC	0.1% N017823 001

HALOBETASOL PROPIONATE

AEROSOL, FOAM;TOPICAL

HALOBETASOL PROPIONATE

<u>AB</u>		PADAGIS ISRAEL	<u>0.05%</u>	<u>A215266 001</u> Aug 11, 2023
<u>LEXETTE</u>				
<u>AB</u>	+	!	MAYNE PHARMA	<u>0.05%</u> <u>N210566 001</u> May 24, 2018

CREAM;TOPICAL

HALOBETASOL PROPIONATE

<u>AB</u>	!	COSETTE	<u>0.05%</u>	<u>A078162 001</u> Apr 24, 2007
<u>AB</u>		FOUGERA PHARMS	<u>0.05%</u>	<u>A077001 001</u> Dec 16, 2004
<u>AB</u>		PADAGIS ISRAEL	<u>0.05%</u>	<u>A077123 001</u> Dec 16, 2004
<u>AB</u>		TARO	<u>0.05%</u>	<u>A077227 001</u> Aug 04, 2005
LOTION;TOPICAL				
BRYHALI				
	+	!	BAUSCH	0.01% N209355 001 Nov 06, 2018
ULTRAVATE				
	+	!	SUN PHARM INDUSTRIES	0.05% N208183 001 Nov 06, 2015
OINTMENT;TOPICAL				
<u>HALOBETASOL PROPIONATE</u>				
<u>AB</u>		COSETTE	<u>0.05%</u>	<u>A077109 001</u> Jun 14, 2005
<u>AB</u>	!	PADAGIS ISRAEL	<u>0.05%</u>	<u>A076872 001</u> Dec 16, 2004
<u>AB</u>		QUAGEN	<u>0.05%</u>	<u>A213560 001</u> Oct 06, 2020
<u>AB</u>		TARO	<u>0.05%</u>	<u>A076994 001</u> Dec 16, 2004

HALOBETASOL PROPIONATE; TAZAROTENE

LOTION;TOPICAL

DUOBRII

	+	!	BAUSCH	0.01%;0.045% N209354 001 Apr 25, 2019
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HALOPERIDOL

TABLET;ORAL

HALOPERIDOL

<u>AB</u>		INNOGENIX	<u>0.5MG</u>	<u>A071173 002</u> Jan 02, 1987
<u>AB</u>			<u>1MG</u>	<u>A071173 003</u> Jan 02, 1987
<u>AB</u>			<u>2MG</u>	<u>A071173 004</u> Jan 02, 1987
<u>AB</u>			<u>5MG</u>	<u>A071173 005</u> Jan 07, 1988
<u>AB</u>			<u>10MG</u>	<u>A071173 001</u> Jan 07, 1988
<u>AB</u>			<u>20MG</u>	<u>A071173 006</u> Jan 07, 1988
<u>AB</u>		MSN	<u>0.5MG</u>	<u>A216004 001</u> Nov 18, 2022
<u>AB</u>			<u>1MG</u>	<u>A216004 002</u> Nov 18, 2022
<u>AB</u>			<u>2MG</u>	<u>A216004 003</u> Nov 18, 2022
<u>AB</u>			<u>5MG</u>	<u>A216004 004</u> Nov 18, 2022
<u>AB</u>			<u>10MG</u>	<u>A216004 005</u> Nov 18, 2022
<u>AB</u>			<u>20MG</u>	<u>A216004 006</u> Nov 18, 2022
<u>AB</u>		MYLAN	<u>0.5MG</u>	<u>A070278 006</u> Jun 10, 1986
<u>AB</u>			<u>1MG</u>	<u>A070278 004</u> Jun 10, 1986
<u>AB</u>	!		<u>2MG</u>	<u>A070278 001</u> Jun 10, 1986
<u>AB</u>			<u>5MG</u>	<u>A070278 005</u> Jun 10, 1986

PRESCRIPTION DRUG PRODUCT LIST

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

<u>AB</u>		<u>10MG</u>	<u>A070278 002</u>	Jul 16, 2009
<u>AB</u>		<u>20MG</u>	<u>A070278 003</u>	Jul 16, 2009
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A071209 002</u>	Nov 17, 1986
<u>AB</u>		<u>1MG</u>	<u>A071209 003</u>	Nov 17, 1986
<u>AB</u>		<u>5MG</u>	<u>A071209 001</u>	Nov 17, 1986
<u>AB</u>		<u>10MG</u>	<u>A071210 001</u>	Mar 11, 1988
<u>AB</u>		<u>20MG</u>	<u>A071211 001</u>	Mar 11, 1988
<u>AB</u>	SCIEGEN PHARMS INC	<u>2MG</u>	<u>A071130 001</u>	Feb 17, 1987
<u>AB</u>		<u>5MG</u>	<u>A071131 001</u>	Feb 17, 1987
<u>AB</u>		<u>10MG</u>	<u>A071132 001</u>	May 12, 1987
<u>AB</u>	UPSHER SMITH LABS	<u>0.5MG</u>	<u>A211061 001</u>	Jan 08, 2020
<u>AB</u>		<u>1MG</u>	<u>A211061 002</u>	Jan 08, 2020
<u>AB</u>		<u>2MG</u>	<u>A211061 003</u>	Jan 08, 2020
<u>AB</u>		<u>5MG</u>	<u>A211061 004</u>	Jan 08, 2020
<u>AB</u>		<u>10MG</u>	<u>A211061 005</u>	Jan 08, 2020
<u>AB</u>		<u>20MG</u>	<u>A211061 006</u>	Jan 08, 2020
<u>AB</u>	ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A077580 001</u>	Jan 17, 2023
<u>AB</u>		<u>1MG</u>	<u>A077580 002</u>	Jan 17, 2023
<u>AB</u>		<u>2MG</u>	<u>A077580 006</u>	Jan 17, 2023
<u>AB</u>		<u>5MG</u>	<u>A077580 003</u>	Nov 29, 2007
<u>AB</u>		<u>10MG</u>	<u>A077580 004</u>	Nov 29, 2007
<u>AB</u>		<u>20MG</u>	<u>A077580 005</u>	Nov 29, 2007

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALDOL

<u>AO</u>	+	JANSSEN PHARMS	<u>EQ 50MG BASE/ML</u>	<u>N018701 001</u>	Jan 14, 1986
<u>AO</u>	+		<u>EQ 100MG BASE/ML</u>	<u>N018701 002</u>	Jan 31, 1997

HALOPERIDOL DECANOATE

<u>AO</u>		FRESENIUS KABI USA	<u>EQ 50MG BASE/ML</u>	<u>A074893 001</u>	Dec 19, 1997
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A074893 002</u>	Dec 19, 1997
<u>AO</u>		GLAND PHARMA LTD	<u>EQ 50MG BASE/ML</u>	<u>A205241 001</u>	May 12, 2017
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A205241 002</u>	May 12, 2017
<u>AO</u>		HIKMA	<u>EQ 50MG BASE/ML</u>	<u>A074811 001</u>	Jan 30, 1998
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A075305 001</u>	Sep 28, 1998
<u>AO</u>		MANKIND PHARMA	<u>EQ 50MG BASE/ML</u>	<u>A216730 001</u>	May 23, 2023
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A216730 002</u>	May 23, 2023
<u>AO</u>		MEITHEAL	<u>EQ 50MG BASE/ML</u>	<u>A214507 001</u>	Jul 26, 2021
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A214507 002</u>	Jul 26, 2021
<u>AO</u>		MYLAN LABS LTD	<u>EQ 50MG BASE/ML</u>	<u>A075440 001</u>	Feb 28, 2000
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A075440 002</u>	Feb 28, 2000
<u>AO</u>		SOMERSET THERAPS LLC	<u>EQ 50MG BASE/ML</u>	<u>A209101 001</u>	Jul 03, 2018
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A209101 002</u>	Jul 03, 2018
<u>AO</u>		TEVA PHARMS USA	<u>EQ 50MG BASE/ML</u>	<u>A075393 001</u>	May 11, 1999
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A075393 002</u>	May 11, 1999
<u>AO</u>		ZYDUS PHARMS	<u>EQ 50MG BASE/ML</u>	<u>A211180 001</u>	Oct 22, 2019
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A211180 002</u>	Oct 22, 2019

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALOPERIDOL

<u>AA</u>		LANNETT CO INC	<u>EQ 2MG BASE/ML</u>	<u>A073364 001</u>	Sep 28, 1993
<u>AA</u>	!	PHARM ASSOC	<u>EQ 2MG BASE/ML</u>	<u>A073037 001</u>	Feb 26, 1993

INJECTABLE; INJECTION

HALOPERIDOL

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A075689 001</u>	Mar 09, 2001
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A210356 001</u>	Jul 01, 2019
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A076774 001</u>	Aug 25, 2004
<u>AP</u>		HIKMA	<u>EQ 5MG BASE/ML</u>	<u>A075858 001</u>	Jun 18, 2001
<u>AP</u>	!	MYLAN LABS LTD	<u>EQ 5MG BASE/ML</u>	<u>A078347 001</u>	Sep 14, 2009
<u>AP</u>		SAGENT PHARMS	<u>EQ 5MG BASE/ML</u>	<u>A091637 001</u>	Sep 02, 2011
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A200742 001</u>	Sep 02, 2011

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>		B BRAUN MEDICAL INC	<u>5,000 UNITS/0.5ML</u>	<u>A208827 001</u>	Nov 19, 2018
<u>AP</u>		BE PHARMS	<u>1,000 UNITS/ML</u>	<u>A214804 001</u>	Dec 29, 2020
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A214839 001</u>	Dec 29, 2020
<u>AP</u>	+	FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<u>N017029 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A206552 001</u>	Jun 10, 2016
<u>AP</u>	<u>+</u>	<u>5,000 UNITS/ML</u>	<u>N017651 006</u>	
<u>AP</u>	<u>+</u>	<u>10,000 UNITS/ML</u>	<u>N017029 003</u>	
<u>AP</u>	<u>+</u>	<u>20,000 UNITS/ML</u>	<u>N017029 004</u>	
<u>AP</u>	GLAND	<u>1,000 UNITS/ML</u>	<u>A205323 002</u>	Nov 18, 2019
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A205323 001</u>	Feb 06, 2017
<u>AP</u>	<u>+</u>	<u>1,000 UNITS/ML</u>	<u>N017037 001</u>	
<u>AP</u>	<u>+</u>	<u>5,000 UNITS/ML</u>	<u>N017037 002</u>	
<u>AP</u>		<u>5,000 UNITS/0.5ML</u>	<u>N017037 013</u>	Apr 07, 1986
<u>AP</u>	<u>+</u>	<u>10,000 UNITS/ML</u>	<u>N017037 003</u>	
<u>AP</u>	HOSPIRA INC	<u>1,000 UNITS/ML</u>	<u>A090571 001</u>	Aug 31, 2009
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A090571 002</u>	Aug 31, 2009
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A090571 003</u>	Aug 31, 2009
<u>AP</u>	MYLAN LABS LTD	<u>1,000 UNITS/ML</u>	<u>A203851 001</u>	Nov 30, 2017
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A203851 002</u>	Nov 30, 2017
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A203851 003</u>	Nov 30, 2017
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A203852 001</u>	Nov 30, 2017
<u>AP</u>	NANJING KING-FRIEND	<u>1,000 UNITS/ML</u>	<u>A211005 001</u>	Dec 14, 2018
<u>AP</u>		<u>1,000 UNITS/ML</u>	<u>A211007 001</u>	May 28, 2019
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A211007 002</u>	May 28, 2019
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A212061 001</u>	Jul 15, 2020
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A211007 003</u>	May 28, 2019
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A211004 001</u>	Feb 24, 2020
<u>AP</u>	SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090808 001</u>	Jun 30, 2010
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A090808 002</u>	Jun 30, 2010
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A090808 003</u>	Jun 30, 2010
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A090809 001</u>	Jun 30, 2010
<u>AP</u>	SANDOZ	<u>1,000 UNITS/ML</u>	<u>A091682 001</u>	Jun 08, 2011
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A091682 002</u>	Jun 08, 2011
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A201002 001</u>	Jun 08, 2011
<u>AP</u>	SHENZHEN TECHDOW	<u>1,000 UNITS/ML</u>	<u>A202957 001</u>	Jun 12, 2014
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A202733 001</u>	Jun 12, 2014
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A202957 002</u>	Jun 12, 2014
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A203198 001</u>	Jun 12, 2014
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A203198 002</u>	Jun 12, 2014
	<u>HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>	BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 001</u>	Apr 28, 1982
	<u>HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>	<u>+</u>	<u>200 UNITS/100ML</u>	<u>N019953 001</u>	Jul 20, 1992
<u>AP</u>	FRESENIUS KABI USA	<u>200 UNITS/100ML</u>	<u>A212441 001</u>	Jul 24, 2020
<u>AP</u>	<u>+</u>	<u>200 UNITS/100ML</u>	<u>N018916 010</u>	Jun 23, 1989
	<u>HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	HOSPIRA	<u>10,000 UNITS/100ML</u>	<u>N019339 003</u>	Mar 27, 1985
	<u>HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>	BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 002</u>	Apr 28, 1982
	<u>HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>	FRESENIUS KABI USA	<u>200 UNITS/100ML</u>	<u>A212441 002</u>	Jul 24, 2020
<u>AP</u>	<u>+</u>	<u>200 UNITS/100ML</u>	<u>N018916 011</u>	Jun 23, 1989
	<u>HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	<u>+</u>	<u>5,000 UNITS/100ML</u>	<u>N019952 004</u>	Jul 20, 1992
<u>AP</u>	<u>+</u>	<u>10,000 UNITS/100ML</u>	<u>N019952 005</u>	Jul 20, 1992
<u>AP</u>	HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N019339 004</u>	Mar 27, 1985
<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N019339 002</u>	Mar 27, 1985
	<u>HEPARIN SODIUM IN PLASTIC CONTAINER</u>			
<u>AP</u>	<u>+</u>	<u>1,000 UNITS/ML</u>	<u>N017029 013</u>	Dec 05, 1985
<u>AP</u>	<u>+</u>	<u>5,000 UNITS/ML</u>	<u>N017029 014</u>	Dec 05, 1985
<u>AP</u>	<u>+</u>	<u>10,000 UNITS/ML</u>	<u>N017029 015</u>	Dec 05, 1985
<u>AP</u>	<u>+</u>	<u>20,000 UNITS/ML</u>	<u>N017029 016</u>	Dec 05, 1985
	<u>HEPARIN SODIUM PRESERVATIVE FREE</u>			
<u>AP</u>	<u>+</u>	<u>1,000 UNITS/ML</u>	<u>N017029 010</u>	Apr 28, 1986
<u>AP</u>	<u>+</u>	<u>10,000 UNITS/ML</u>	<u>N017029 019</u>	Nov 22, 2010
<u>AP</u>	HOSPIRA	<u>10,000 UNITS/ML</u>	<u>A089522 001</u>	May 04, 1987
<u>AP</u>	NANJING KING-FRIEND	<u>10,000 UNITS/ML</u>	<u>A212060 001</u>	Apr 02, 2020
<u>AP</u>	SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090810 001</u>	Jun 30, 2010
<u>AP</u>	SHENZHEN TECHDOW	<u>1,000 UNITS/ML</u>	<u>A202732 001</u>	Jun 12, 2014
	<u>HEPARIN SODIUM</u>			
	<u>+</u>	<u>10,000 UNITS/ML</u>	<u>N017029 020</u>	Mar 31, 2011
	<u>!</u>	<u>5,000 UNITS/ML</u>	<u>A088100 001</u>	Apr 28, 1983
	<u>+</u>	<u>1,000 UNITS/ML</u>	<u>N201370 001</u>	Jul 21, 2011

PRESCRIPTION DRUG PRODUCT LIST

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

+	!		5,000 UNITS/ML	N201370	002	Jul 21, 2011
		HEPARIN SODIUM 12,500 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER			
		HOSPIRA	5,000 UNITS/100ML	N019339	001	Mar 27, 1985
		HEPARIN SODIUM 12,500 UNITS	IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
		HOSPIRA	5,000 UNITS/100ML	N018916	006	Jan 31, 1984
		HEPARIN SODIUM 2,000 UNITS	AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
		BAXTER HLTHCARE	2,000 UNITS/1000ML	N018609	004	Apr 23, 2020
		HEPARIN SODIUM 2,000 UNITS	IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+	!	B BRAUN	200 UNITS/100ML	N019953	002	Dec 15, 2023
		HEPARIN SODIUM 20,000 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER			
+	!	B BRAUN	4,000 UNITS/100ML	N019952	001	Jul 20, 1992
		HEPARIN SODIUM 25,000 UNITS	IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
		HOSPIRA	5,000 UNITS/100ML	N018916	007	Jan 31, 1984
			10,000 UNITS/100ML	N018916	008	Jan 31, 1984
		HEPARIN SODIUM PRESERVATIVE	FREE			
+	!	PFIZER	1,000 UNITS/ML	N201370	004	Jul 21, 2011

HEPARIN SODIUM; TAUROLIDINE

SOLUTION; N/A

DEFENCATH

+	!	CORMEDIX	3,000 UNITS/3ML (1,000 UNITS/ML); 40.5MG/3ML (13.5MG/ML)	N214520	001	Nov 15, 2023
+	!		5,000 UNITS/5ML (1,000 UNITS/ML); 67.5MG/5ML (13.5MG/ML)	N214520	002	Nov 15, 2023

HEXACHLOROPHENE

SPONGE; TOPICAL

PRE-OP**AT** +! DAVIS AND GECK **480MG** **N017433 001**PRE-OP II**AT** + DAVIS AND GECK **480MG** **N017433 002**HEXAMINOLEVULINATE HYDROCHLORIDE

FOR SOLUTION; INTRAVESICAL

CYSVIEW KIT

+	!	PHOTOCURE ASA	100MG/VIAL	N022555	001	May 28, 2010
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HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

SUPPRELIN LA

+	!	ENDO PHARM	50MG	N022058	001	May 03, 2007
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HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODAN**AA** + GENUS **1.5MG/5ML; 5MG/5ML** **N005213 002** Jul 26, 1988HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE**AA** ABHAI LLC **1.5MG/5ML; 5MG/5ML** **A207487 001** Feb 21, 2017**AA** ACTAVIS MID ATLANTIC **1.5MG/5ML; 5MG/5ML** **A088017 001** Jul 05, 1983**AA** NOVEL LABS INC **1.5MG/5ML; 5MG/5ML** **A203535 001** Feb 13, 2017**AA** PADAGIS US **1.5MG/5ML; 5MG/5ML** **A205731 001** Feb 15, 2017**AA** ! PAI HOLDINGS PHARM **1.5MG/5ML; 5MG/5ML** **A040613 001** Feb 08, 2008**AA** WOCKHARDT BIO AG **1.5MG/5ML; 5MG/5ML** **A088008 001** Mar 03, 1983

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE**AA** ! AVANTHI INC **1.5MG; 5MG** **A207176 001** Aug 07, 2017HYCODAN**AA** + GENUS **1.5MG; 5MG** **N005213 001** Jul 26, 1988HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

AP		AM REGENT	20MG/ML	A040136 001		Jun 30, 1997
AP		BEOWULF ASSET	20MG/ML	A203110 001		Jun 29, 2015
AP		EUGIA PHARMA	20MG/ML	A215147 001		Feb 13, 2023
AP		FRESENIUS KABI USA	20MG/ML	A040388 001		Mar 13, 2001
AP		HIKMA	20MG/ML	A213667 001		Dec 18, 2020
AP		NAVINTA LLC	20MG/ML	A202938 001		Mar 28, 2013
AP	!	RISING	20MG/ML	A040730 001		Apr 21, 2009

PRESCRIPTION DRUG PRODUCT LISTHYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

<u>AA</u>	ALKEM LABS LTD	<u>10MG</u>	<u>A200737 001</u>	Dec 07, 2012
<u>AA</u>		<u>25MG</u>	<u>A200737 002</u>	Dec 07, 2012
<u>AA</u>		<u>50MG</u>	<u>A200737 003</u>	Dec 07, 2012
<u>AA</u>		<u>100MG</u>	<u>A200737 004</u>	Dec 07, 2012
<u>AA</u>	CADILA PHARMS LTD	<u>25MG</u>	<u>A203845 001</u>	Sep 18, 2014
<u>AA</u>		<u>50MG</u>	<u>A203845 002</u>	Sep 18, 2014
<u>AA</u>		<u>100MG</u>	<u>A203845 003</u>	Sep 18, 2014
<u>AA</u>	GLENMARK PHARMS LTD	<u>10MG</u>	<u>A090527 001</u>	May 27, 2009
<u>AA</u>		<u>25MG</u>	<u>A090527 002</u>	May 27, 2009
<u>AA</u>		<u>50MG</u>	<u>A090527 003</u>	May 27, 2009
<u>AA</u>		<u>100MG</u>	<u>A090527 004</u>	May 27, 2009
<u>AA</u>	HERITAGE PHARMS	<u>10MG</u>	<u>A086242 001</u>	Feb 04, 2010
<u>AA</u>		<u>25MG</u>	<u>A086242 003</u>	
<u>AA</u>		<u>50MG</u>	<u>A086242 002</u>	
<u>AA</u>		<u>100MG</u>	<u>A086242 004</u>	Feb 04, 2010
<u>AA</u>	HERITAGE PHARMS INC	<u>10MG</u>	<u>A040858 001</u>	Feb 26, 2010
<u>AA</u>		<u>25MG</u>	<u>A040858 002</u>	Feb 26, 2010
<u>AA</u>		<u>50MG</u>	<u>A040858 003</u>	Feb 26, 2010
<u>AA</u>		<u>100MG</u>	<u>A040858 004</u>	Feb 26, 2010
<u>AA</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A040901 001</u>	Sep 12, 2008
<u>AA</u>		<u>25MG</u>	<u>A040901 002</u>	Sep 12, 2008
<u>AA</u>		<u>50MG</u>	<u>A040901 003</u>	Sep 12, 2008
<u>AA</u>		<u>100MG</u>	<u>A040901 004</u>	Sep 12, 2008
<u>AA</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A090255 001</u>	Dec 15, 2008
<u>AA</u>		<u>25MG</u>	<u>A090255 002</u>	Dec 15, 2008
<u>AA</u>		<u>50MG</u>	<u>A090255 003</u>	Dec 15, 2008
<u>AA</u>		<u>100MG</u>	<u>A090255 004</u>	Dec 15, 2008
<u>AA</u>	! PLIVA	<u>10MG</u>	<u>A089097 001</u>	Dec 18, 1985
<u>AA</u>	+!	<u>25MG</u>	<u>A088467 001</u>	May 01, 1984
<u>AA</u>	+!	<u>50MG</u>	<u>A088468 001</u>	May 01, 1984
<u>AA</u>	!	<u>100MG</u>	<u>A089098 001</u>	Dec 18, 1985
<u>AA</u>	SCIEGEN PHARMS INC	<u>10MG</u>	<u>A205236 001</u>	May 26, 2017
<u>AA</u>		<u>25MG</u>	<u>A205236 002</u>	May 26, 2017
<u>AA</u>		<u>50MG</u>	<u>A205236 003</u>	May 26, 2017
<u>AA</u>		<u>100MG</u>	<u>A205236 004</u>	May 26, 2017
<u>AA</u>	STRIDES PHARMA	<u>10MG</u>	<u>A087836 001</u>	Oct 05, 1982
<u>AA</u>		<u>25MG</u>	<u>A086961 002</u>	
<u>AA</u>		<u>25MG</u>	<u>A200770 001</u>	May 03, 2013
<u>AA</u>		<u>50MG</u>	<u>A086962 001</u>	
<u>AA</u>		<u>50MG</u>	<u>A200770 002</u>	May 03, 2013
<u>AA</u>		<u>100MG</u>	<u>A088391 001</u>	Sep 27, 1983
<u>AA</u>		<u>100MG</u>	<u>A200770 003</u>	May 03, 2013

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRA-ZIDE

	STRIDES PHARMA	25MG; 25MG	A088957 001	Oct 21, 1985
	!	50MG; 50MG	A088946 001	Oct 21, 1985

HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE

TABLET; ORAL

BIDIL

<u>AB</u>	+! AZURITY	<u>37.5MG; 20MG</u>	<u>N020727 001</u>	Jun 23, 2005
	<u>ISOSORBIDE DINITRATE AND HYDRALAZINE HYDROCHLORIDE</u>			
<u>AB</u>	RICONPHARMA LLC	<u>37.5MG; 20MG</u>	<u>A215586 001</u>	Apr 06, 2022

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG</u>	<u>A078164 001</u>	Sep 18, 2007
<u>AB</u>	IPCA LABS LTD	<u>12.5MG</u>	<u>A079237 001</u>	Apr 02, 2009
<u>AB</u>	JUBILANT CADISTA	<u>12.5MG</u>	<u>A078391 001</u>	Feb 11, 2008
<u>AB</u>	PRINSTON INC	<u>12.5MG</u>	<u>A075907 001</u>	Sep 17, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>12.5MG</u>	<u>A203561 001</u>	Jan 14, 2019
<u>AB</u>	UNICHEM	<u>12.5MG</u>	<u>A090510 001</u>	Jan 19, 2010

MICROZIDE

<u>AB</u>	+! TEVA BRANDED PHARM	<u>12.5MG</u>	<u>N020504 001</u>	Dec 27, 1996
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TABLET; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>	! ACCORD HLTHCARE	<u>12.5MG</u>	<u>A202556 001</u>	Sep 24, 2012
<u>AB</u>		<u>25MG</u>	<u>A202556 002</u>	Sep 24, 2012

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>50MG</u>	<u>A202556</u>	<u>003</u>	Sep 24, 2012
<u>AB</u>	ACTAVIS ELIZABETH	<u>12.5MG</u>	<u>A040707</u>	<u>001</u>	Feb 27, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A040780</u>	<u>001</u>	Jul 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040780</u>	<u>002</u>	Jul 20, 2007
<u>AB</u>	HERITAGE PHARMS	<u>12.5MG</u>	<u>A085182</u>	<u>003</u>	May 02, 2023
<u>AB</u>		<u>25MG</u>	<u>A085182</u>	<u>002</u>	
<u>AB</u>		<u>50MG</u>	<u>A085182</u>	<u>001</u>	
<u>AB</u>	IPCA LABS LTD	<u>12.5MG</u>	<u>A040807</u>	<u>001</u>	Jul 20, 2007
<u>AB</u>		<u>25MG</u>	<u>A040807</u>	<u>002</u>	Jul 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040807</u>	<u>003</u>	Jul 20, 2007
<u>AB</u>	+ IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A083177</u>	<u>001</u>	
<u>AB</u>	+!	<u>50MG</u>	<u>A083177</u>	<u>002</u>	
<u>AB</u>	LEADING	<u>12.5MG</u>	<u>A040702</u>	<u>003</u>	May 10, 2017
<u>AB</u>		<u>25MG</u>	<u>A040702</u>	<u>001</u>	Mar 16, 2007
<u>AB</u>		<u>50MG</u>	<u>A040702</u>	<u>002</u>	Mar 16, 2007
<u>AB</u>	OXFORD PHARMS	<u>25MG</u>	<u>A087059</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A087068</u>	<u>001</u>	
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A040412</u>	<u>001</u>	Mar 29, 2002
<u>AB</u>		<u>50MG</u>	<u>A040412</u>	<u>002</u>	Mar 29, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A203018</u>	<u>001</u>	Jul 23, 2014
<u>AB</u>		<u>50MG</u>	<u>A203018</u>	<u>002</u>	Jul 23, 2014
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A040907</u>	<u>001</u>	Aug 15, 2008
<u>AB</u>		<u>50MG</u>	<u>A040907</u>	<u>002</u>	Aug 15, 2008

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

<u>AB</u>	+!	SANOFI AVENTIS US	<u>12.5MG; 150MG</u>	<u>N020758</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+!		<u>12.5MG; 300MG</u>	<u>N020758</u>	<u>003</u>	Aug 31, 1998

IRBESARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMBIC	<u>12.5MG; 150MG</u>	<u>A091370</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A091370</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>		<u>25MG; 300MG</u>	<u>A091370</u>	<u>003</u>	Oct 12, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>12.5MG; 150MG</u>	<u>A203630</u>	<u>001</u>	Feb 22, 2013
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A203630</u>	<u>002</u>	Feb 22, 2013
<u>AB</u>		<u>25MG; 300MG</u>	<u>A203630</u>	<u>003</u>	Mar 31, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>12.5MG; 150MG</u>	<u>A203500</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A203500</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>	HIKMA	<u>12.5MG; 150MG</u>	<u>A090351</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A090351</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>		<u>25MG; 300MG</u>	<u>A090351</u>	<u>003</u>	Jun 08, 2017
<u>AB</u>	HISUN PHARM HANGZHOU	<u>12.5MG; 150MG</u>	<u>A207896</u>	<u>001</u>	Oct 14, 2016
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A207896</u>	<u>002</u>	Oct 14, 2016
<u>AB</u>	LUPIN LTD	<u>12.5MG; 150MG</u>	<u>A201524</u>	<u>001</u>	Feb 27, 2013
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A201524</u>	<u>002</u>	Feb 27, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG; 150MG</u>	<u>A202414</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A202414</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG; 150MG</u>	<u>A203072</u>	<u>001</u>	May 09, 2014
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A203072</u>	<u>002</u>	May 09, 2014
<u>AB</u>	SANDOZ	<u>12.5MG; 150MG</u>	<u>A077446</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A077446</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>	TEVA	<u>12.5MG; 150MG</u>	<u>A077369</u>	<u>001</u>	Mar 30, 2012
<u>AB</u>		<u>12.5MG; 300MG</u>	<u>A077369</u>	<u>002</u>	Mar 30, 2012

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	AUROBINDO	<u>12.5MG; 10MG</u>	<u>A077606</u>	<u>001</u>	Mar 14, 2006
<u>AB</u>		<u>12.5MG; 20MG</u>	<u>A077606</u>	<u>002</u>	Mar 14, 2006
<u>AB</u>		<u>25MG; 20MG</u>	<u>A077606</u>	<u>003</u>	Mar 14, 2006
<u>AB</u>	INVAGEN PHARMS	<u>12.5MG; 10MG</u>	<u>A204058</u>	<u>001</u>	May 23, 2017
<u>AB</u>	!	<u>12.5MG; 20MG</u>	<u>A204058</u>	<u>002</u>	May 23, 2017
<u>AB</u>	!	<u>25MG; 20MG</u>	<u>A204058</u>	<u>003</u>	May 23, 2017
<u>AB</u>	LUPIN	<u>12.5MG; 10MG</u>	<u>A077912</u>	<u>001</u>	Sep 27, 2006
<u>AB</u>		<u>12.5MG; 20MG</u>	<u>A077912</u>	<u>002</u>	Sep 27, 2006
<u>AB</u>		<u>25MG; 20MG</u>	<u>A077912</u>	<u>003</u>	Sep 27, 2006
<u>AB</u>	PRINSTON INC	<u>12.5MG; 10MG</u>	<u>A076230</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG; 20MG</u>	<u>A076230</u>	<u>002</u>	Jul 01, 2002

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>25MG; 20MG</u>	<u>A076230 003</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>12.5MG; 10MG</u>	<u>A076262 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG; 20MG</u>	<u>A076262 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG; 20MG</u>	<u>A076262 003</u>	Jul 01, 2002
<u>AB</u>	SUN PHARM INDS LTD	<u>12.5MG; 10MG</u>	<u>A076007 001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG; 20MG</u>	<u>A076007 002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG; 20MG</u>	<u>A076007 003</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>12.5MG; 10MG</u>	<u>A076194 003</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG; 20MG</u>	<u>A076194 001</u>	Jul 01, 2002
<u>AB</u>		<u>25MG; 20MG</u>	<u>A076194 002</u>	Jul 01, 2002
<u>ZESTORETIC</u>				
<u>AB</u>	+ ALMATICA	<u>12.5MG; 10MG</u>	<u>N019888 003</u>	Nov 18, 1993
<u>AB</u>	+	<u>12.5MG; 20MG</u>	<u>N019888 001</u>	Sep 20, 1990
<u>AB</u>	+	<u>25MG; 20MG</u>	<u>N019888 002</u>	Jul 20, 1989

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

HYZAAR

<u>AB</u>	+ ORGANON	<u>12.5MG; 50MG</u>	<u>N020387 001</u>	Apr 28, 1995
<u>AB</u>	+	<u>12.5MG; 100MG</u>	<u>N020387 003</u>	Oct 20, 2005
<u>AB</u>	+	<u>25MG; 100MG</u>	<u>N020387 002</u>	Nov 10, 1998

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMbic PHARMS LTD	<u>12.5MG; 50MG</u>	<u>A091617 001</u>	Feb 17, 2012
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A091617 002</u>	Feb 17, 2012
<u>AB</u>		<u>25MG; 100MG</u>	<u>A091617 003</u>	Feb 17, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG; 50MG</u>	<u>A091629 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A091629 002</u>	Oct 06, 2010
<u>AB</u>	!	<u>25MG; 100MG</u>	<u>A091629 003</u>	Jan 06, 2010
<u>AB</u>	GRANULES	<u>12.5MG; 50MG</u>	<u>A218015 001</u>	Sep 29, 2023
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A218015 002</u>	Sep 29, 2023
<u>AB</u>		<u>25MG; 100MG</u>	<u>A218015 003</u>	Sep 29, 2023
<u>AB</u>	IPCA LABS LTD	<u>12.5MG; 50MG</u>	<u>A201682 001</u>	Mar 01, 2013
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A201682 002</u>	Mar 01, 2013
<u>AB</u>		<u>25MG; 100MG</u>	<u>A201682 003</u>	Mar 01, 2013
<u>AB</u>	JUBILANT CADISTA	<u>12.5MG; 50MG</u>	<u>A201845 001</u>	Sep 18, 2012
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A201845 002</u>	Sep 18, 2012
<u>AB</u>		<u>25MG; 100MG</u>	<u>A201845 003</u>	Sep 18, 2012
<u>AB</u>	LUPIN LTD	<u>12.5MG; 50MG</u>	<u>A078245 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A078245 002</u>	May 21, 2010
<u>AB</u>		<u>25MG; 100MG</u>	<u>A078245 003</u>	Oct 06, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG; 50MG</u>	<u>A202289 001</u>	Aug 09, 2012
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A202289 002</u>	Aug 09, 2012
<u>AB</u>		<u>25MG; 100MG</u>	<u>A202289 003</u>	Aug 09, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG; 50MG</u>	<u>A204901 001</u>	Nov 06, 2017
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A204901 002</u>	Nov 06, 2017
<u>AB</u>		<u>25MG; 100MG</u>	<u>A204901 003</u>	Nov 06, 2017
<u>AB</u>	SANDOZ	<u>12.5MG; 50MG</u>	<u>A077948 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A077948 003</u>	Aug 19, 2010
<u>AB</u>		<u>25MG; 100MG</u>	<u>A077948 002</u>	Oct 06, 2010
<u>AB</u>	TEVA PHARMS	<u>12.5MG; 50MG</u>	<u>A077157 001</u>	Apr 06, 2010
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A077157 002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG; 100MG</u>	<u>A077157 003</u>	Apr 06, 2010
<u>AB</u>	UNICHEM	<u>12.5MG; 50MG</u>	<u>A204832 001</u>	Jul 21, 2017
<u>AB</u>		<u>12.5MG; 100MG</u>	<u>A204832 002</u>	Jul 21, 2017
<u>AB</u>		<u>25MG; 100MG</u>	<u>A204832 003</u>	Jul 21, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>12.5MG; 50MG</u>	<u>A078385 001</u>	Oct 06, 2010
<u>AB</u>		<u>25MG; 100MG</u>	<u>A078385 002</u>	Oct 06, 2010

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

<u>AB</u>	+ VALIDUS PHARMS	<u>25MG; 50MG</u>	<u>N018303 001</u>	Dec 31, 1984
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METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMbic	<u>25MG; 50MG</u>	<u>A202870 001</u>	Nov 06, 2013
<u>AB</u>	!	<u>25MG; 100MG</u>	<u>A202870 002</u>	Nov 06, 2013
<u>AB</u>		<u>50MG; 100MG</u>	<u>A202870 003</u>	Nov 06, 2013
<u>AB</u>	MYLAN	<u>25MG; 50MG</u>	<u>A076792 001</u>	Aug 20, 2004
<u>AB</u>		<u>25MG; 100MG</u>	<u>A076792 002</u>	Aug 20, 2004
<u>AB</u>		<u>50MG; 100MG</u>	<u>A076792 003</u>	Aug 20, 2004
<u>AB</u>	SUN PHARM INDS	<u>25MG; 50MG</u>	<u>A090654 001</u>	Jan 19, 2012

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

AB		<u>25MG;10MG</u>	<u>A090654 002</u>	Jan 19, 2012
AB		<u>50MG;10MG</u>	<u>A090654 003</u>	Jan 19, 2012

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB	GLENMARK PHARMS	<u>12.5MG;7.5MG</u>	<u>A090718 001</u>	Mar 17, 2010
AB		<u>12.5MG;15MG</u>	<u>A090718 002</u>	Mar 17, 2010
AB		<u>25MG;15MG</u>	<u>A090718 003</u>	Mar 17, 2010
AB	TEVA	<u>12.5MG;7.5MG</u>	<u>A076980 001</u>	Mar 07, 2007
AB		<u>12.5MG;15MG</u>	<u>A076980 003</u>	Mar 07, 2007
AB	!	<u>25MG;15MG</u>	<u>A076980 002</u>	Mar 07, 2007

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR HCT

AB	+ COSETTE	<u>12.5MG;20MG</u>	<u>N021532 002</u>	Jun 05, 2003
AB	+	<u>12.5MG;40MG</u>	<u>N021532 003</u>	Jun 05, 2003
AB	+	<u>25MG;40MG</u>	<u>N021532 005</u>	Jun 05, 2003

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

AB	ACCORD HLTHCARE	<u>12.5MG;20MG</u>	<u>A209281 001</u>	Feb 07, 2019
AB		<u>12.5MG;40MG</u>	<u>A209281 002</u>	Feb 07, 2019
AB		<u>25MG;40MG</u>	<u>A209281 003</u>	Feb 07, 2019
AB	ALEMBIC	<u>12.5MG;20MG</u>	<u>A204233 001</u>	Apr 24, 2017
AB		<u>12.5MG;40MG</u>	<u>A204233 002</u>	Apr 24, 2017
AB		<u>25MG;40MG</u>	<u>A204233 003</u>	Apr 24, 2017
AB	AUROBINDO PHARMA	<u>12.5MG;20MG</u>	<u>A205391 001</u>	Apr 24, 2017
AB		<u>12.5MG;40MG</u>	<u>A205391 002</u>	Apr 24, 2017
AB		<u>25MG;40MG</u>	<u>A205391 003</u>	Apr 24, 2017
AB	MACLEODS PHARMS LTD	<u>12.5MG;20MG</u>	<u>A204801 001</u>	Jun 01, 2023
AB		<u>12.5MG;40MG</u>	<u>A204801 002</u>	Jun 01, 2023
AB		<u>25MG;40MG</u>	<u>A204801 003</u>	Jun 01, 2023
AB	PRINSTON INC	<u>12.5MG;20MG</u>	<u>A207804 001</u>	Apr 24, 2017
AB		<u>12.5MG;40MG</u>	<u>A207804 002</u>	Apr 24, 2017
AB		<u>25MG;40MG</u>	<u>A207804 003</u>	Apr 24, 2017
AB	TEVA PHARMS USA	<u>12.5MG;40MG</u>	<u>A200532 002</u>	Apr 24, 2017
AB	UMEDICA	<u>12.5MG;20MG</u>	<u>A208847 001</u>	Sep 17, 2019
AB		<u>12.5MG;40MG</u>	<u>A208847 003</u>	Sep 17, 2019
AB		<u>25MG;40MG</u>	<u>A208847 002</u>	Sep 17, 2019

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB	APOTEX	<u>12.5MG;EQ 10MG BASE</u>	<u>A091524 001</u>	Mar 12, 2013
AB		<u>12.5MG;EQ 20MG BASE</u>	<u>A091524 002</u>	Mar 12, 2013
AB		<u>25MG;EQ 20MG BASE</u>	<u>A091524 003</u>	Mar 12, 2013
AB	AUROBINDO PHARMA	<u>12.5MG;EQ 10MG BASE</u>	<u>A078450 001</u>	Aug 24, 2007
AB		<u>12.5MG;EQ 20MG BASE</u>	<u>A078450 002</u>	Aug 24, 2007
AB	!	<u>25MG;EQ 20MG BASE</u>	<u>A078450 003</u>	Aug 24, 2007
AB	INVAGEN PHARMS	<u>12.5MG;EQ 10MG BASE</u>	<u>A201356 001</u>	Apr 20, 2011
AB		<u>12.5MG;EQ 20MG BASE</u>	<u>A201356 002</u>	Apr 20, 2011
AB		<u>25MG;EQ 20MG BASE</u>	<u>A201356 003</u>	Apr 20, 2011

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

ALDACTAZIDE

AB	+ PFIZER	<u>25MG;25MG</u>	<u>N012616 004</u>	Dec 30, 1982
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SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

AB	MYLAN	<u>25MG;25MG</u>	<u>A086513 001</u>	
AB	SUN PHARM INDUSTRIES	<u>25MG;25MG</u>	<u>A089534 001</u>	Jul 02, 1987

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

MICARDIS HCT

AB	+ BOEHRINGER INGELHEIM	<u>12.5MG;40MG</u>	<u>N021162 001</u>	Nov 17, 2000
AB	+	<u>12.5MG;80MG</u>	<u>N021162 002</u>	Nov 17, 2000
AB	+	<u>25MG;80MG</u>	<u>N021162 003</u>	Apr 19, 2004

TELMISARTAN AND HYDROCHLOROTHIAZIDE

AB	ALEMBIC	<u>12.5MG;40MG</u>	<u>A203010 001</u>	Feb 25, 2014
AB		<u>12.5MG;80MG</u>	<u>A203010 002</u>	Feb 25, 2014

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>25MG; 80MG</u>	<u>A203010 003</u>	Feb 25, 2014
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG; 40MG</u>	<u>A208727 001</u>	Dec 15, 2016
<u>AB</u>		<u>12.5MG; 80MG</u>	<u>A208727 002</u>	Dec 15, 2016
<u>AB</u>		<u>25MG; 80MG</u>	<u>A208727 003</u>	Dec 15, 2016
<u>AB</u>	GLENMARK PHARMS LTD	<u>12.5MG; 40MG</u>	<u>A202544 001</u>	Mar 04, 2019
<u>AB</u>		<u>12.5MG; 80MG</u>	<u>A202544 002</u>	Mar 04, 2019
<u>AB</u>		<u>25MG; 80MG</u>	<u>A202544 003</u>	Mar 04, 2019
<u>AB</u>	LUPIN LTD	<u>12.5MG; 40MG</u>	<u>A091351 001</u>	Aug 07, 2014
<u>AB</u>		<u>12.5MG; 80MG</u>	<u>A091351 002</u>	Aug 07, 2014
<u>AB</u>		<u>25MG; 80MG</u>	<u>A091351 003</u>	Aug 07, 2014
<u>AB</u>	PRINSTON INC	<u>12.5MG; 40MG</u>	<u>A209028 001</u>	Nov 06, 2017
<u>AB</u>		<u>12.5MG; 80MG</u>	<u>A209028 002</u>	Nov 06, 2017
<u>AB</u>		<u>25MG; 80MG</u>	<u>A209028 003</u>	Nov 06, 2017
<u>AB</u>	ZYDUS PHARMS	<u>12.5MG; 40MG</u>	<u>A204221 001</u>	Aug 15, 2017
<u>AB</u>		<u>12.5MG; 80MG</u>	<u>A204221 002</u>	Aug 15, 2017
<u>AB</u>		<u>25MG; 80MG</u>	<u>A204221 003</u>	Aug 15, 2017

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	CADILA	<u>25MG; 37.5MG</u>	<u>A208358 001</u>	Feb 11, 2019
<u>AB</u>	LANNETT CO INC	<u>25MG; 37.5MG</u>	<u>A201407 001</u>	Dec 09, 2011
<u>AB</u>	! SANDOZ	<u>25MG; 37.5MG</u>	<u>A074821 001</u>	Jun 05, 1997

TABLET; ORAL

MAXZIDE

<u>AB</u>	+! AUROBINDO PHARMA USA	<u>50MG; 75MG</u>	<u>N019129 001</u>	Oct 22, 1984
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MAXZIDE-25

<u>AB</u>	+ AUROBINDO PHARMA USA	<u>25MG; 37.5MG</u>	<u>N019129 003</u>	May 13, 1988
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TRIAMTERENE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX INC	<u>25MG; 37.5MG</u>	<u>A071251 002</u>	May 05, 1998
<u>AB</u>		<u>50MG; 75MG</u>	<u>A071251 001</u>	Apr 17, 1988
<u>AB</u>	RUBICON	<u>25MG; 37.5MG</u>	<u>A216211 001</u>	Feb 23, 2022
<u>AB</u>		<u>50MG; 75MG</u>	<u>A216211 002</u>	Feb 23, 2022
<u>AB</u>	SANDOZ	<u>25MG; 37.5MG</u>	<u>A073281 001</u>	Apr 30, 1992
<u>AB</u>		<u>50MG; 75MG</u>	<u>A072011 001</u>	Jun 17, 1988
<u>AB</u>	WATSON LABS	<u>25MG; 37.5MG</u>	<u>A073449 001</u>	Sep 23, 1993
<u>AB</u>		<u>50MG; 75MG</u>	<u>A071851 001</u>	Nov 30, 1988
<u>AB</u>	ZYDUS PHARMS	<u>25MG; 37.5MG</u>	<u>A208360 001</u>	Jun 29, 2018
<u>AB</u>		<u>50MG; 75MG</u>	<u>A208360 002</u>	Jun 29, 2018

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

<u>AB</u>	+ NOVARTIS	<u>12.5MG; 80MG</u>	<u>N020818 001</u>	Mar 06, 1998
<u>AB</u>	+	<u>12.5MG; 160MG</u>	<u>N020818 002</u>	Mar 06, 1998
<u>AB</u>	+	<u>12.5MG; 320MG</u>	<u>N020818 004</u>	Apr 28, 2006
<u>AB</u>	+	<u>25MG; 160MG</u>	<u>N020818 003</u>	Jan 17, 2002
<u>AB</u>	+!	<u>25MG; 320MG</u>	<u>N020818 005</u>	Apr 28, 2006

VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMBIC	<u>12.5MG; 80MG</u>	<u>A201662 001</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A201662 002</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A201662 003</u>	Mar 21, 2013
<u>AB</u>		<u>25MG; 160MG</u>	<u>A201662 004</u>	Mar 21, 2013
<u>AB</u>		<u>25MG; 320MG</u>	<u>A201662 005</u>	Mar 21, 2013
<u>AB</u>	AMNEAL PHARMS	<u>12.5MG; 80MG</u>	<u>A204382 001</u>	Aug 11, 2023
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A204382 002</u>	Aug 11, 2023
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A204382 004</u>	Aug 11, 2023
<u>AB</u>		<u>25MG; 160MG</u>	<u>A204382 003</u>	Aug 11, 2023
<u>AB</u>		<u>25MG; 320MG</u>	<u>A204382 005</u>	Aug 11, 2023
<u>AB</u>	AUROBINDO PHARMA LTD	<u>12.5MG; 80MG</u>	<u>A202519 001</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A202519 002</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A202519 003</u>	Mar 21, 2013
<u>AB</u>		<u>25MG; 160MG</u>	<u>A202519 004</u>	Mar 21, 2013
<u>AB</u>		<u>25MG; 320MG</u>	<u>A202519 005</u>	Mar 21, 2013
<u>AB</u>	LUPIN LTD	<u>12.5MG; 80MG</u>	<u>A078946 003</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A078946 004</u>	Mar 21, 2013
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A078946 001</u>	Mar 21, 2013
<u>AB</u>		<u>25MG; 160MG</u>	<u>A078946 005</u>	Mar 21, 2013

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>25MG; 320MG</u>	<u>A078946 002</u>	Mar 21, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG; 80MG</u>	<u>A203145 001</u>	Apr 19, 2013
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A203145 002</u>	Apr 19, 2013
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A203145 003</u>	Apr 19, 2013
<u>AB</u>		<u>25MG; 160MG</u>	<u>A203145 004</u>	Apr 19, 2013
<u>AB</u>		<u>25MG; 320MG</u>	<u>A203145 005</u>	Apr 19, 2013
<u>AB</u>	MYLAN PHARMS INC	<u>12.5MG; 80MG</u>	<u>A078020 001</u>	Sep 21, 2012
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A078020 002</u>	Sep 21, 2012
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A078020 004</u>	Sep 21, 2012
<u>AB</u>		<u>25MG; 160MG</u>	<u>A078020 003</u>	Sep 21, 2012
<u>AB</u>		<u>25MG; 320MG</u>	<u>A078020 005</u>	Sep 21, 2012
<u>AB</u>	PRINSTON INC	<u>12.5MG; 80MG</u>	<u>A206083 001</u>	Feb 08, 2016
<u>AB</u>		<u>12.5MG; 160MG</u>	<u>A206083 002</u>	Feb 08, 2016
<u>AB</u>		<u>12.5MG; 320MG</u>	<u>A206083 003</u>	Feb 08, 2016
<u>AB</u>		<u>25MG; 160MG</u>	<u>A206083 004</u>	Feb 08, 2016
<u>AB</u>		<u>25MG; 320MG</u>	<u>A206083 005</u>	Feb 08, 2016

HYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

HYDROCODONE BITARTRATE

! ALVOGEN

10MG

A206986 001 Jan 21, 2020

15MG

A206986 002 Jan 21, 2020

20MG

A206986 003 Jan 21, 2020

30MG

A206986 004 Jan 21, 2020

40MG

A206986 005 Jan 21, 2020

50MG

A206986 006 Jan 21, 2020

TABLET, EXTENDED RELEASE; ORAL

HYDROCODONE BITARTRATE

<u>AB</u>	ALVOGEN	<u>20MG</u>	<u>A208269 001</u>	Mar 01, 2021
<u>AB</u>		<u>30MG</u>	<u>A208269 002</u>	Mar 01, 2021
<u>AB</u>		<u>40MG</u>	<u>A208269 003</u>	Mar 01, 2021
<u>AB</u>		<u>60MG</u>	<u>A208269 004</u>	Mar 01, 2021
<u>AB</u>		<u>80MG</u>	<u>A208269 005</u>	Mar 01, 2021
<u>AB</u>		<u>100MG</u>	<u>A208269 006</u>	Mar 01, 2021
<u>AB</u>		<u>120MG</u>	<u>A208269 007</u>	Mar 01, 2021

HYSINGLA ER

<u>AB</u>	+! PURDUE PHARMA LP	<u>20MG</u>	<u>N206627 001</u>	Nov 20, 2014
<u>AB</u>	+	<u>30MG</u>	<u>N206627 002</u>	Nov 20, 2014
<u>AB</u>	+	<u>40MG</u>	<u>N206627 003</u>	Nov 20, 2014
<u>AB</u>	+	<u>60MG</u>	<u>N206627 004</u>	Nov 20, 2014
<u>AB</u>	+	<u>80MG</u>	<u>N206627 005</u>	Nov 20, 2014
<u>AB</u>	+	<u>100MG</u>	<u>N206627 006</u>	Nov 20, 2014
<u>AB</u>	+	<u>120MG</u>	<u>N206627 007</u>	Nov 20, 2014

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE

<u>AB</u>	NOSTRUM LABS INC	<u>5MG; 200MG</u>	<u>A077723 003</u>	Nov 06, 2006
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HYDROCODONE BITARTRATE AND IBUPROFEN

<u>AB</u>	ACTAVIS LABS FL INC	<u>7.5MG; 200MG</u>	<u>A076604 001</u>	Dec 31, 2003
<u>AB</u>	AMNEAL PHARMS NY	<u>5MG; 200MG</u>	<u>A076642 002</u>	Mar 18, 2004
<u>AB</u>	!	<u>7.5MG; 200MG</u>	<u>A076642 001</u>	Oct 12, 2004
<u>AB</u>	AUROLIFE PHARMA LLC	<u>7.5MG; 200MG</u>	<u>A204575 001</u>	Jun 02, 2016
<u>AB</u>	NOSTRUM LABS INC	<u>7.5MG; 200MG</u>	<u>A077723 001</u>	Nov 06, 2006
		10MG; 200MG	A077723 002	Nov 06, 2006

HYDROCORTISONE

CREAM; TOPICAL

ALA-CORT

<u>AT</u>	CROWN LABS	<u>2.5%</u>	<u>A080706 007</u>	Jan 05, 2016
<u>AT</u>		<u>1%</u>	<u>A080706 006</u>	

ANUSOL HC

<u>AT</u>	SALIX PHARMS	<u>2.5%</u>	<u>A088250 001</u>	Jun 06, 1984
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HYDROCORTISONE

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>1%</u>	<u>A087795 001</u>	May 03, 1983
<u>AT</u>		<u>2.5%</u>	<u>A089682 001</u>	Mar 10, 1988
<u>AT</u>	+! FOUGERA PHARMS INC	<u>1%</u>	<u>A080693 003</u>	
<u>AT</u>	!	<u>2.5%</u>	<u>A089414 001</u>	Dec 16, 1986
<u>AT</u>	PADAGIS US	<u>2.5%</u>	<u>A085025 001</u>	
<u>AT</u>	RISING	<u>2.5%</u>	<u>A040879 001</u>	Aug 20, 2010

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE

CREAM; TOPICAL

HYDROCORTISONE

AT TARO **2.5%** **A088799 001** Nov 09, 1984
ENEMA; RECTAL

COLOCORT

AB CHARTWELL **100MG/60ML** **A075172 001** Dec 03, 1999

CORTENEMA

AB +! ANI PHARMS **100MG/60ML** **N016199 001**

GRANULE; ORAL

ALKINDI SPRINKLE

+ ETON

0.5MG

N213876 001 Sep 29, 2020

+

1MG

N213876 002 Sep 29, 2020

+

2MG

N213876 003 Sep 29, 2020

+!

5MG

N213876 004 Sep 29, 2020

LOTION; TOPICAL

HYDROCORTISONE

AT TARO **2.5%** **A040247 001** Jul 23, 1999

STIE-CORT

AT ! PADAGIS US **2.5%** **A089074 001** Nov 26, 1985

ALA-SCALP

LEGACY PHARMA

2%

A083231 001

OINTMENT; TOPICAL

HYDROCORTISONE

AT +! FOUGERA PHARMS **1%** **A080692 001**

AT ! FOUGERA PHARMS INC **2.5%** **A081203 001** May 28, 1993

AT PADAGIS US **2.5%** **A085027 001**

AT TARO **1%** **A086257 001**

HYDROCORTISONE IN ABSORBASE

AT CMP PHARMA INC **1%** **A088138 001** Sep 06, 1985

SOLUTION; TOPICAL

TEXACORT

! MISSION PHARMA

2.5%

A081271 001 Apr 17, 1992

TABLET; ORAL

CORTEF

AB + PHARMACIA AND **5MG** **N008697 003**
UPJOHN

AB + **10MG** **N008697 001**

AB +! **20MG** **N008697 002**

HYDROCORTISONE

AB AUROBINDO PHARMA **5MG** **A214649 001** Jul 17, 2023
LTD

AB **10MG** **A214649 002** Jul 17, 2023

AB **20MG** **A214649 003** Jul 17, 2023

AB IMPAX LABS INC **5MG** **A040646 001** Mar 30, 2007

AB **10MG** **A040646 002** Mar 30, 2007

AB **20MG** **A040646 003** Mar 30, 2007

AB STRIDES PHARMA **5MG** **A207029 001** Apr 27, 2017

AB **10MG** **A207029 002** Apr 27, 2017

AB **20MG** **A207029 003** Apr 27, 2017

HYDROCORTISONE ACETATE

AEROSOL, METERED; RECTAL

CORTIFOAM

+! MYLAN SPECIALITY LP 10%

N017351 001 Feb 10, 1982

CREAM; TOPICAL

MICORT-HC

SEBELA IRELAND LTD

2.5%

A040396 001 Feb 27, 2001

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED; TOPICAL

EPIFOAM

BX MYLAN SPECIALITY LP **1%;1%** **A086457 001**
PROCTOFOAM HC

BX +! MYLAN SPECIALITY LP **1%;1%** **A086195 001**

CREAM; TOPICAL

PRAMOSONE

SEBELA IRELAND LTD

0.5%;1%

A083778 001

1%;1%

A085368 001

LOTION; TOPICAL

PRAMOSONE

SEBELA IRELAND LTD

1%;1%

A085980 001

2.5%;1%

A085979 001

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

HYDROCORTISONE BUTYRATE**AB1** TARO PHARM INDS **0.1%** **A076654 001** Aug 03, 2005LOCOID**AB1** +! BAUSCH **0.1%** **N018514 001** Mar 31, 1982HYDROCORTISONE BUTYRATE**AB2** ACTAVIS MID **0.1%** **A205134 001** Dec 08, 2017

ATLANTIC

AB2 GLENMARK GENERICS **0.1%** **A202145 001** Sep 27, 2013LOCOID LIPOCREAM**AB2** +! PRECISION DERMAT **0.1%** **N020769 001** Sep 08, 1997

LOTION; TOPICAL

HYDROCORTISONE BUTYRATE**AB** LUPIN LTD **0.1%** **A210209 001** Aug 17, 2018**AB** THE J MOLNER **0.1%** **A209556 001** Nov 21, 2017LOCOID**AB** +! BAUSCH **0.1%** **N022076 001** May 18, 2007

OINTMENT; TOPICAL

HYDROCORTISONE BUTYRATE**AB** TARO **0.1%** **A076842 001** Dec 27, 2004LOCOID**AB** +! PRECISION DERMAT **0.1%** **N018652 001** Oct 29, 1982

SOLUTION; TOPICAL

HYDROCORTISONE BUTYRATE**AT** TARO PHARM INDS **0.1%** **A076364 001** Jan 14, 2004LOCOID**AT** +! BAUSCH **0.1%** **N019116 001** Feb 25, 1987HYDROCORTISONE PROBUTATE

CREAM; TOPICAL

PANDEL

+! ANI PHARMS **0.1%** **N020453 001** Feb 28, 1997HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

SOLU-CORTEF

+! PHARMACIA AND **EQ 100MG BASE/VIAL** **N009866 001**

UPJOHN

+! **EQ 250MG BASE/VIAL** **N009866 002**+! **EQ 500MG BASE/VIAL** **N009866 003**+! **EQ 1GM BASE/VIAL** **N009866 004**HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE**AB** COSETTE **0.2%** **A213724 001** Feb 11, 2021**AB** ENCUBE ETHICALS **0.2%** **A211047 001** Nov 04, 2021**AB** GLENMARK PHARMS LTD **0.2%** **A211129 001** Oct 12, 2018**AB** LUPIN LTD **0.2%** **A210307 001** Aug 15, 2019**AB** PADAGIS ISRAEL **0.2%** **A075666 001** May 24, 2000**AB** ! TARO **0.2%** **A075042 001** Aug 25, 1998

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE**AB** COSETTE **0.2%** **A211764 001** Mar 04, 2020**AB** GLENMARK PHARMS LTD **0.2%** **A211750 001** Dec 14, 2018**AB** ! TARO **0.2%** **A075043 001** Aug 25, 1998HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE**AT** ! BAUSCH AND LOMB **1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML** **A064053 001** Dec 29, 1995**AT** SANDOZ **1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML** **A062423 001** Aug 25, 1983

SUSPENSION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE! SANDOZ **1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML** **A062874 001** May 11, 1988

SUSPENSION/DROPS; OTIC

CASPORYN HC**AT** +! CASPER PHARMA LLC **1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML** **N060613 001**NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE**AT** AMRING PHARMS **1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML** **A065219 001** May 01, 2006**AT** SANDOZ **1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML** **A062488 001** Nov 06, 1985OTICAIR**AT** BAUSCH AND LOMB **1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML** **A064065 001** Aug 28, 1996

PRESCRIPTION DRUG PRODUCT LIST

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

DILAUDID

<u>AP</u>	+	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>N019034</u>	<u>003</u>	Apr 30, 2009
<u>AP</u>	+		<u>2MG/ML</u>	<u>N019034</u>	<u>004</u>	Apr 30, 2009

HYDROMORPHONE HYDROCHLORIDE

<u>AP</u>		HIKMA	<u>2MG/ML</u>	<u>A202159</u>	<u>001</u>	Apr 27, 2018
<u>AP</u>		HOSPIRA INC	<u>1MG/ML</u>	<u>N200403</u>	<u>001</u>	Dec 01, 2011
<u>AP</u>			<u>2MG/ML</u>	<u>N200403</u>	<u>002</u>	Dec 01, 2011
<u>AP</u>			<u>10MG/ML</u>	<u>A078591</u>	<u>001</u>	Jun 17, 2008
<u>AP</u>		RISING	<u>10MG/ML</u>	<u>A078228</u>	<u>001</u>	Apr 14, 2010
<u>AP</u>			<u>10MG/ML</u>	<u>A078261</u>	<u>001</u>	Apr 14, 2010

DILAUDID

+	FRESENIUS KABI USA	0.2MG/ML	N019034	006	Jan 16, 2020
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HYDROMORPHONE HYDROCHLORIDE

	HOSPIRA INC	0.25MG/0.5ML	N200403	005	Mar 03, 2023
		0.5MG/0.5ML	N200403	004	Jun 13, 2022
		4MG/ML	N200403	003	Dec 01, 2011

SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

HYDROMORPHONE HYDROCHLORIDE

+	HIKMA	40MG/20ML (2MG/ML)	N217812	001	Dec 14, 2023
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SOLUTION; ORAL

DILAUDID

<u>AA</u>	+	RHODES PHARMS	<u>5MG/5ML</u>	<u>N019891</u>	<u>001</u>	Dec 07, 1992
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HYDROMORPHONE HYDROCHLORIDE

<u>AA</u>		ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A210176</u>	<u>001</u>	Oct 27, 2017
<u>AA</u>		HIKMA	<u>5MG/5ML</u>	<u>A074653</u>	<u>001</u>	Jul 29, 1998

TABLET; ORAL

DILAUDID

<u>AB</u>	+	RHODES PHARMS	<u>2MG</u>	<u>N019892</u>	<u>003</u>	Nov 09, 2007
<u>AB</u>	+		<u>4MG</u>	<u>N019892</u>	<u>002</u>	Nov 09, 2007
<u>AB</u>	+		<u>8MG</u>	<u>N019892</u>	<u>001</u>	Dec 07, 1992

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>		ASCENT PHARMS INC	<u>2MG</u>	<u>A210506</u>	<u>001</u>	Jan 17, 2018
<u>AB</u>			<u>4MG</u>	<u>A210506</u>	<u>002</u>	Jan 17, 2018
<u>AB</u>			<u>8MG</u>	<u>A210506</u>	<u>003</u>	Jan 17, 2018
<u>AB</u>		AUROLIFE PHARMA LLC	<u>2MG</u>	<u>A205814</u>	<u>001</u>	May 13, 2016
<u>AB</u>			<u>4MG</u>	<u>A205814</u>	<u>002</u>	May 13, 2016
<u>AB</u>			<u>8MG</u>	<u>A205814</u>	<u>003</u>	May 13, 2016
<u>AB</u>		SPECGX LLC	<u>2MG</u>	<u>A076855</u>	<u>002</u>	Sep 19, 2007
<u>AB</u>			<u>4MG</u>	<u>A076855</u>	<u>003</u>	Sep 19, 2007
<u>AB</u>			<u>8MG</u>	<u>A076855</u>	<u>001</u>	Dec 23, 2004

TABLET, EXTENDED RELEASE; ORAL

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>		ASCENT PHARMS INC	<u>8MG</u>	<u>A212133</u>	<u>001</u>	Sep 23, 2020
<u>AB</u>			<u>12MG</u>	<u>A212133</u>	<u>002</u>	Sep 23, 2020
<u>AB</u>			<u>16MG</u>	<u>A212133</u>	<u>003</u>	Sep 23, 2020
<u>AB</u>			<u>32MG</u>	<u>A212133</u>	<u>004</u>	Sep 23, 2020
<u>AB</u>		OSMOTICA PHARM US	<u>8MG</u>	<u>A205629</u>	<u>001</u>	Jul 07, 2016
<u>AB</u>			<u>12MG</u>	<u>A205629</u>	<u>002</u>	Jul 07, 2016
<u>AB</u>			<u>16MG</u>	<u>A205629</u>	<u>003</u>	Jul 07, 2016
<u>AB</u>	!		<u>32MG</u>	<u>A205629</u>	<u>004</u>	Jul 07, 2016
<u>AB</u>		PADAGIS US	<u>8MG</u>	<u>A204278</u>	<u>001</u>	Apr 06, 2015
<u>AB</u>			<u>12MG</u>	<u>A204278</u>	<u>002</u>	Apr 06, 2015
<u>AB</u>			<u>16MG</u>	<u>A204278</u>	<u>003</u>	Apr 06, 2015
<u>AB</u>			<u>32MG</u>	<u>A204278</u>	<u>004</u>	Sep 20, 2017

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

CYANOKIT

+	BTG INTL	5GM/VIAL (5GM/KIT)	N022041	001	Apr 08, 2011
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HYDROXOCOBALAMIN

!	ACTAVIS	1MG/ML	A085998	001	
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HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

<u>AB</u>		ACCORD HLTHCARE	<u>100MG</u>	<u>A213342</u>	<u>002</u>	Aug 18, 2021
<u>AB</u>			<u>200MG</u>	<u>A213342</u>	<u>001</u>	Apr 07, 2020
<u>AB</u>			<u>300MG</u>	<u>A213342</u>	<u>003</u>	Aug 18, 2021
<u>AB</u>	!		<u>400MG</u>	<u>A213342</u>	<u>004</u>	Aug 18, 2021
<u>AB</u>		ALKALOIDA ZRT	<u>200MG</u>	<u>A201691</u>	<u>001</u>	May 08, 2018
<u>AB</u>		AMNEAL PHARMS CO	<u>200MG</u>	<u>A210577</u>	<u>001</u>	May 15, 2018

PRESCRIPTION DRUG PRODUCT LIST

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

AB	APPCO	100MG	A210441 002	Sep 19, 2022
AB		200MG	A210441 001	May 01, 2018
AB		300MG	A210441 003	Sep 19, 2022
AB		400MG	A210441 004	Sep 19, 2022
AB	AUROBINDO PHARMA USA	200MG	A040274 001	May 29, 1998
AB	CHARTWELL RX	200MG	A210543 001	Jul 06, 2018
AB	IPCA LABS LTD	200MG	A040766 001	Jun 14, 2007
AB	LAURUS	200MG	A210959 001	Jan 15, 2019
AB	RISE PHARMA	200MG	A212902 001	May 14, 2020
AB	SANDOZ	200MG	A040104 001	Nov 30, 1995
AB	ZYDUS PHARMS USA INC	200MG	A040657 001	Sep 21, 2007

PLAQUENIL

AB	+ ! CONCORDIA	200MG	N009768 001	
	SOVUNA			
	+ NOVITIUM PHARMA	200MG	N214581 001	Jan 14, 2022
	+ !	300MG	N214581 002	Jan 14, 2022

HYDROXYPROPYL CELLULOSE

INSERT; OPHTHALMIC

LACRISERT

+ !	BAUSCH AND LOMB INC	5MG	N018771 001	
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HYDROXYUREA

CAPSULE; ORAL

HYDREA

AB	+ ! CHEPLAPHARM	500MG	N016295 001	
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HYDROXYUREA

AB	BARR	500MG	A075143 001	Oct 16, 1998
AB	LEADING	500MG	A213438 001	Apr 08, 2020
AB	PAR PHARM	500MG	A075340 001	Feb 24, 1999
	DROXIA			
	+ CHEPLAPHARM	200MG	N016295 002	Feb 25, 1998
	+	300MG	N016295 003	Feb 25, 1998
	+	400MG	N016295 004	Feb 25, 1998

TABLET; ORAL

SIKLOS

+	THERAVIA	100MG	N208843 001	Dec 21, 2017
+ !		1GM	N208843 002	Dec 21, 2017

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

+ !	AM REGENT	25MG/ML	A087408 001	
+ !		50MG/ML	A087408 002	

SYRUP; ORAL

HYDROXYZINE HYDROCHLORIDE

AA	+ ! CHARTWELL RX	10MG/5ML	A087294 001	Apr 12, 1982
AA	LANNETT CO INC	10MG/5ML	A201674 001	Aug 21, 2013

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

AB	AMNEAL PHARM	10MG	A040808 001	Sep 24, 2008
AB		25MG	A040808 002	Sep 24, 2008
AB		50MG	A040808 003	Sep 24, 2008
AB	AUROBINDO PHARMA LTD	10MG	A087871 002	Dec 20, 1982
AB		25MG	A087871 003	Dec 20, 1982
AB		50MG	A087871 001	Dec 20, 1982
AB	CHARTWELL RX	10MG	A040804 001	Jun 30, 2008
AB		25MG	A040804 002	Jun 30, 2008
AB		50MG	A040804 003	Jun 30, 2008
AB	EPIC PHARMA LLC	10MG	A040604 002	Dec 28, 2004
AB		25MG	A040604 003	Dec 28, 2004
AB		50MG	A040604 001	Dec 28, 2004
AB	GRAVITI PHARMS	10MG	A217652 001	Aug 17, 2023
AB		25MG	A217652 002	Aug 17, 2023
AB		50MG	A217652 003	Aug 17, 2023
AB	HERITAGE PHARMA	10MG	A204279 001	Aug 20, 2014
AB		25MG	A204279 002	Aug 20, 2014
AB		50MG	A204279 003	Aug 20, 2014

PRESCRIPTION DRUG PRODUCT LIST

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u>		HETERO LABS LTD III	<u>10MG</u>	<u>A040805</u>	<u>001</u>	May 29, 2008
<u>AB</u>			<u>25MG</u>	<u>A040805</u>	<u>002</u>	May 29, 2008
<u>AB</u>			<u>50MG</u>	<u>A040805</u>	<u>003</u>	May 29, 2008
<u>AB</u>		INVAGEN PHARMS	<u>10MG</u>	<u>A040812</u>	<u>001</u>	Mar 12, 2008
<u>AB</u>			<u>25MG</u>	<u>A040812</u>	<u>002</u>	Mar 12, 2008
<u>AB</u>			<u>50MG</u>	<u>A040812</u>	<u>003</u>	Mar 12, 2008
<u>AB</u>		KVK TECH	<u>10MG</u>	<u>A040786</u>	<u>001</u>	Mar 20, 2007
<u>AB</u>			<u>25MG</u>	<u>A040786</u>	<u>002</u>	Mar 20, 2007
<u>AB</u>			<u>50MG</u>	<u>A040786</u>	<u>003</u>	Mar 20, 2007
<u>AB</u>		NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A040840</u>	<u>002</u>	Mar 31, 2008
<u>AB</u>			<u>25MG</u>	<u>A040840</u>	<u>003</u>	Mar 31, 2008
<u>AB</u>			<u>50MG</u>	<u>A040840</u>	<u>001</u>	Mar 31, 2008
<u>AB</u>		NUVO PHARMS INC	<u>10MG</u>	<u>A207121</u>	<u>002</u>	Mar 29, 2017
<u>AB</u>			<u>25MG</u>	<u>A207121</u>	<u>001</u>	Mar 29, 2017
<u>AB</u>			<u>50MG</u>	<u>A207121</u>	<u>003</u>	Mar 29, 2017
<u>AB</u>	+	PLIVA	<u>10MG</u>	<u>A088617</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	+		<u>25MG</u>	<u>A088618</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	+		<u>50MG</u>	<u>A088619</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>		PRINSTON INC	<u>10MG</u>	<u>A040580</u>	<u>003</u>	May 27, 2005
<u>AB</u>			<u>25MG</u>	<u>A040580</u>	<u>002</u>	May 27, 2005
<u>AB</u>			<u>50MG</u>	<u>A040580</u>	<u>001</u>	May 27, 2005

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

<u>AB</u>		BARR	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A088496</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>			<u>EQ 50MG HYDROCHLORIDE</u>	<u>A088487</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>		HERITAGE PHARMA	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A201507</u>	<u>001</u>	Jun 03, 2013
<u>AB</u>			<u>EQ 50MG HYDROCHLORIDE</u>	<u>A201507</u>	<u>002</u>	Jun 03, 2013
<u>AB</u>		IMPAX LABS INC	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>001</u>	Jul 15, 1996
<u>AB</u>			<u>EQ 50MG HYDROCHLORIDE</u>	<u>A040156</u>	<u>002</u>	Jul 15, 1996
<u>AB</u>		SANDOZ	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A087479</u>	<u>001</u>	
<u>AB</u>	!		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A086183</u>	<u>001</u>	
		<u>VISTARIL</u>				
<u>AB</u>	+	PFIZER	<u>EQ 25MG HYDROCHLORIDE</u>	<u>N011459</u>	<u>002</u>	
<u>AB</u>	+		<u>EQ 50MG HYDROCHLORIDE</u>	<u>N011459</u>	<u>004</u>	

HYDROXYZINE PAMOATE

BARR

EQ 100MG HYDROCHLORIDE

A088488 001 Jun 15, 1984

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

IBANDRONATE SODIUM

<u>AP</u>		ACCORD HLTHCARE	<u>EQ 3MG BASE/3ML</u>	<u>A206058</u>	<u>001</u>	Feb 05, 2016
<u>AP</u>	!	APOTEX	<u>EQ 3MG BASE/3ML</u>	<u>A204222</u>	<u>001</u>	Oct 16, 2015
<u>AP</u>		EUGIA PHARMA	<u>EQ 3MG BASE/3ML</u>	<u>A205332</u>	<u>001</u>	Aug 19, 2015
<u>AP</u>		MYLAN LABS LTD	<u>EQ 3MG BASE/3ML</u>	<u>A202671</u>	<u>001</u>	Sep 02, 2014
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 3MG BASE/3ML</u>	<u>A202235</u>	<u>001</u>	Sep 02, 2014
<u>AP</u>		SUN PHARM	<u>EQ 3MG BASE/3ML</u>	<u>A090853</u>	<u>001</u>	Feb 14, 2014

TABLET; ORAL

IBANDRONATE SODIUM

<u>AB</u>		APOTEX INC	<u>EQ 150MG BASE</u>	<u>A078948</u>	<u>001</u>	Mar 19, 2012
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 150MG BASE</u>	<u>A204502</u>	<u>001</u>	Mar 11, 2016
<u>AB</u>	!	DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A078997</u>	<u>001</u>	Apr 30, 2012
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A206887</u>	<u>001</u>	Oct 31, 2017
<u>AB</u>		ORBION PHARMS	<u>EQ 150MG BASE</u>	<u>A078998</u>	<u>001</u>	Mar 19, 2012
<u>AB</u>		WATSON LABS TEVA	<u>EQ 150MG BASE</u>	<u>A079003</u>	<u>001</u>	Mar 20, 2012

IBREXAFUNGERP CITRATE

TABLET; ORAL

BREXAFEMME

+! SCYNEXIS

EQ 150MG BASE

N214900 001 Jun 01, 2021

IBRUTINIB

CAPSULE; ORAL

IMBRUVICA

+ PHARMACYCLICS LLC

70MG

N205552 002 Dec 20, 2017

+!

140MG

N205552 001 Nov 13, 2013

SUSPENSION; ORAL

IMBRUVICA

+! PHARMACYCLICS LLC

70MG/ML

N217003 001 Aug 24, 2022

PRESCRIPTION DRUG PRODUCT LIST

IBRUTINIB

TABLET; ORAL

IMBRUVICA

+	PHARMACYCLICS LLC	140MG	N210563	001	Feb 16, 2018
+		280MG	N210563	002	Feb 16, 2018
+		420MG	N210563	003	Feb 16, 2018

IBUPROFEN

SOLUTION; INTRAVENOUS

CALDOLOR

+	CUMBERLAND PHARMS	800MG/8ML (100MG/ML)	N022348	002	Jun 11, 2009
+		800MG/200ML (4MG/ML)	N022348	003	Jan 25, 2019

SUSPENSION; ORAL

IBUPROFEN

<u>AB</u>	!	ACTAVIS MID ATLANTIC	<u>100MG/5ML</u>	<u>A074978</u>	<u>001</u>	Mar 25, 1998
<u>AB</u>		AUROBINDO PHARMA LTD	<u>100MG/5ML</u>	<u>A209178</u>	<u>001</u>	Feb 16, 2018
<u>AB</u>		PADAGIS US	<u>100MG/5ML</u>	<u>A076925</u>	<u>001</u>	Sep 23, 2004
<u>AB</u>		STRIDES PHARMA	<u>100MG/5ML</u>	<u>A215311</u>	<u>001</u>	May 27, 2022
<u>AB</u>		TARO	<u>100MG/5ML</u>	<u>A209204</u>	<u>001</u>	Jun 23, 2017

TABLET; ORAL

IBUPROFEN

<u>AB</u>		ALKEM LABS LTD	<u>400MG</u>	<u>A214699</u>	<u>001</u>	Sep 13, 2021
<u>AB</u>			<u>600MG</u>	<u>A214699</u>	<u>002</u>	Sep 13, 2021
<u>AB</u>			<u>800MG</u>	<u>A214699</u>	<u>003</u>	Sep 13, 2021
<u>AB</u>		AMNEAL PHARMS NY	<u>400MG</u>	<u>A071334</u>	<u>001</u>	Nov 25, 1986
<u>AB</u>			<u>400MG</u>	<u>A078558</u>	<u>001</u>	Jun 18, 2007
<u>AB</u>			<u>600MG</u>	<u>A071335</u>	<u>001</u>	Nov 25, 1986
<u>AB</u>			<u>600MG</u>	<u>A078558</u>	<u>002</u>	Jun 18, 2007
<u>AB</u>			<u>800MG</u>	<u>A071935</u>	<u>001</u>	Oct 13, 1987
<u>AB</u>			<u>800MG</u>	<u>A078558</u>	<u>003</u>	Jun 18, 2007
<u>AB</u>		AUROBINDO PHARMA LTD	<u>400MG</u>	<u>A213794</u>	<u>001</u>	May 08, 2020
<u>AB</u>			<u>600MG</u>	<u>A213794</u>	<u>002</u>	May 08, 2020
<u>AB</u>			<u>800MG</u>	<u>A213794</u>	<u>003</u>	May 08, 2020
<u>AB</u>		CONTRACT PHARMACAL	<u>400MG</u>	<u>A071268</u>	<u>002</u>	Oct 15, 1986
<u>AB</u>			<u>600MG</u>	<u>A071268</u>	<u>001</u>	Oct 15, 1986
<u>AB</u>			<u>800MG</u>	<u>A071268</u>	<u>003</u>	Jul 01, 1988
<u>AB</u>		DR REDDYS LA	<u>400MG</u>	<u>A075682</u>	<u>001</u>	Nov 14, 2001
<u>AB</u>			<u>600MG</u>	<u>A075682</u>	<u>002</u>	Nov 14, 2001
<u>AB</u>	!		<u>800MG</u>	<u>A075682</u>	<u>003</u>	Nov 14, 2001
<u>AB</u>		DR REDDYS LABS INC	<u>400MG</u>	<u>A076112</u>	<u>001</u>	Oct 31, 2001
<u>AB</u>			<u>600MG</u>	<u>A076112</u>	<u>002</u>	Oct 31, 2001
<u>AB</u>			<u>800MG</u>	<u>A076112</u>	<u>003</u>	Oct 31, 2001
<u>AB</u>		GRANULES	<u>400MG</u>	<u>A091625</u>	<u>001</u>	Sep 15, 2015
<u>AB</u>			<u>600MG</u>	<u>A091625</u>	<u>002</u>	Sep 15, 2015
<u>AB</u>			<u>800MG</u>	<u>A091625</u>	<u>003</u>	Sep 15, 2015
<u>AB</u>		MARKSANS PHARMA	<u>400MG</u>	<u>A090796</u>	<u>001</u>	Dec 21, 2010
<u>AB</u>			<u>600MG</u>	<u>A090796</u>	<u>002</u>	Dec 21, 2010
<u>AB</u>			<u>800MG</u>	<u>A090796</u>	<u>003</u>	Dec 21, 2010
<u>AB</u>		SHANDONG XINHUA	<u>400MG</u>	<u>A202413</u>	<u>001</u>	Nov 23, 2016
<u>AB</u>			<u>600MG</u>	<u>A202413</u>	<u>002</u>	Nov 23, 2016
<u>AB</u>			<u>800MG</u>	<u>A202413</u>	<u>003</u>	Nov 23, 2016
<u>AB</u>		STRIDES PHARMA	<u>400MG</u>	<u>A078329</u>	<u>001</u>	Feb 05, 2009
<u>AB</u>			<u>600MG</u>	<u>A078329</u>	<u>002</u>	Feb 05, 2009
<u>AB</u>			<u>800MG</u>	<u>A078329</u>	<u>003</u>	Feb 05, 2009
<u>AB</u>		YICHANG HUMANWELL	<u>400MG</u>	<u>A215318</u>	<u>001</u>	Mar 30, 2022
<u>AB</u>			<u>600MG</u>	<u>A215318</u>	<u>002</u>	Mar 30, 2022
<u>AB</u>			<u>800MG</u>	<u>A215318</u>	<u>003</u>	Mar 30, 2022

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS

IBUPROFEN LYSINE

<u>AP</u>		XGEN PHARMS	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>A202402</u>	<u>001</u>	Mar 30, 2016
<u>AP</u>	+	RECORDATI RARE	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>	<u>N021903</u>	<u>001</u>	Apr 13, 2006

PRESCRIPTION DRUG PRODUCT LIST

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

CORVERT

<u>AP</u>	<u>+</u> !	PFIZER	<u>0.1MG/ML</u>	<u>N020491</u>	<u>001</u>	Dec 28, 1995
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IBUTILIDE FUMARATE

<u>AP</u>		MYLAN INSTITUTIONAL	<u>0.1MG/ML</u>	<u>A090643</u>	<u>001</u>	Jan 11, 2010
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ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

FIRAZYR

<u>AP</u>	<u>+</u> !	TAKEDA PHARMS USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>N022150</u>	<u>001</u>	Aug 25, 2011
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ICATIBANT ACETATE

<u>AP</u>		CIPLA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A212446</u>	<u>001</u>	Jul 13, 2020
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<u>AP</u>		EUGIA PHARMA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A213521</u>	<u>001</u>	Aug 14, 2023
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<u>AP</u>		FRESENIUS KABI USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A208317</u>	<u>001</u>	Jun 18, 2020
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<u>AP</u>		JIANGSU HANSOH PHARM	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A211021</u>	<u>001</u>	Mar 09, 2020
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<u>AP</u>		NANG KUANG PHARM CO	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A212081</u>	<u>001</u>	Dec 16, 2020
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<u>AP</u>		SLAYBACK PHARMA LLC	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A211501</u>	<u>001</u>	Sep 01, 2020
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<u>AP</u>		TEVA PHARMS USA	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A210118</u>	<u>001</u>	Jul 15, 2019
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ICODEXTRIN

SOLUTION; INTRAPERITONEAL

EXTRANEAL

<u>+</u> !	BAXTER HLTHCARE	7.5GM/100ML	<u>N021321</u>	<u>001</u>	Dec 20, 2002
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ICOSAPENT ETHYL

CAPSULE; ORAL

ICOSAPENT ETHYL

<u>AB</u>		APOTEX	<u>1GM</u>	<u>A209437</u>	<u>001</u>	Jun 30, 2021
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<u>AB</u>		ASCENT PHARMS INC	<u>500MG</u>	<u>A216811</u>	<u>001</u>	Dec 07, 2023
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<u>AB</u>		DR REDDYS	<u>500MG</u>	<u>A209499</u>	<u>002</u>	Mar 08, 2023
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<u>AB</u>			<u>1GM</u>	<u>A209499</u>	<u>001</u>	Aug 07, 2020
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<u>AB</u>		HIKMA	<u>500MG</u>	<u>A209457</u>	<u>002</u>	Mar 08, 2023
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<u>AB</u>			<u>1GM</u>	<u>A209457</u>	<u>001</u>	May 21, 2020
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<u>AB</u>		HUMANWELL PURACAP	<u>500MG</u>	<u>A217919</u>	<u>001</u>	Dec 22, 2023
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<u>AB</u>			<u>1GM</u>	<u>A217919</u>	<u>002</u>	Dec 22, 2023
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<u>AB</u>		STRIDES PHARMA	<u>500MG</u>	<u>A217844</u>	<u>001</u>	Sep 22, 2023
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<u>AB</u>			<u>1GM</u>	<u>A217844</u>	<u>002</u>	Sep 22, 2023
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<u>AB</u>		TEVA PHARMS USA	<u>500MG</u>	<u>A209525</u>	<u>001</u>	Sep 11, 2020
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<u>AB</u>			<u>1GM</u>	<u>A209525</u>	<u>002</u>	Sep 11, 2020
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<u>AB</u>		ZYDUS	<u>500MG</u>	<u>A217656</u>	<u>001</u>	Apr 20, 2023
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<u>AB</u>			<u>1GM</u>	<u>A217656</u>	<u>002</u>	Apr 20, 2023
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VASCEPA

<u>AB</u>	<u>+</u>	AMARIN PHARMS	<u>500MG</u>	<u>N202057</u>	<u>002</u>	Feb 16, 2017
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<u>AB</u>	<u>+</u> !		<u>1GM</u>	<u>N202057</u>	<u>001</u>	Jul 26, 2012
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IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFS

<u>AP</u>	<u>+</u> !	PFIZER	<u>1MG/ML</u>	<u>N050734</u>	<u>001</u>	Feb 17, 1997
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IDARUBICIN HYDROCHLORIDE

<u>AP</u>		HIKMA	<u>1MG/ML</u>	<u>A065275</u>	<u>001</u>	Dec 14, 2006
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<u>AP</u>			<u>1MG/ML</u>	<u>A065288</u>	<u>001</u>	May 15, 2007
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<u>AP</u>		MEITHEAL	<u>1MG/ML</u>	<u>A065036</u>	<u>001</u>	May 01, 2002
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IDELALISIB

TABLET; ORAL

ZYDELIG

<u>+</u>	GILEAD SCIENCES INC	100MG	<u>N205858</u>	<u>001</u>	Jul 23, 2014
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<u>+</u> !		150MG	<u>N205858</u>	<u>002</u>	Jul 23, 2014
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IFOSFAMIDE

INJECTABLE; INJECTION

IFEX

<u>AP</u>	<u>+</u>	BAXTER HLTHCARE	<u>1GM/VIAL</u>	<u>N019763</u>	<u>001</u>	Dec 30, 1988
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<u>AP</u>	<u>+</u>		<u>3GM/VIAL</u>	<u>N019763</u>	<u>002</u>	Dec 30, 1988
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IFOSFAMIDE

<u>AP</u>	<u>!</u>	FRESENIUS KABI USA	<u>1GM/VIAL</u>	<u>A076078</u>	<u>001</u>	May 28, 2002
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<u>AP</u>	<u>!</u>		<u>3GM/VIAL</u>	<u>A076078</u>	<u>002</u>	May 28, 2002
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<u>AP</u>		HIKMA	<u>1GM/20ML (50MG/ML)</u>	<u>A076619</u>	<u>001</u>	Jun 29, 2011
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<u>AP</u>			<u>3GM/60ML (50MG/ML)</u>	<u>A076619</u>	<u>002</u>	Jun 29, 2011
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<u>AP</u>	<u>!</u>	MEITHEAL	<u>1GM/20ML (50MG/ML)</u>	<u>A076657</u>	<u>001</u>	Apr 04, 2007
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<u>AP</u>	<u>!</u>		<u>3GM/60ML (50MG/ML)</u>	<u>A076657</u>	<u>002</u>	Apr 04, 2007
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PRESCRIPTION DRUG PRODUCT LIST

ILOPERIDONE

TABLET; ORAL

FANAPT

+	!	VANDA PHARMS INC	1MG	N022192	001	May 06, 2009
+			2MG	N022192	002	May 06, 2009
+			4MG	N022192	003	May 06, 2009
+			6MG	N022192	004	May 06, 2009
+			8MG	N022192	005	May 06, 2009
+			10MG	N022192	006	May 06, 2009
+			12MG	N022192	007	May 06, 2009

ILOPROST

SOLUTION; INHALATION

VENTAVIS

+	!	ACTELION	10MCG/ML (10MCG/ML)	N021779	002	Dec 08, 2005
+	!		20MCG/ML (20MCG/ML)	N021779	003	Aug 07, 2009

IMATINIB MESYLATE

TABLET; ORAL

GLEEVEC

AB	+	NOVARTIS	<u>EQ 100MG BASE</u>	<u>N021588</u>	<u>001</u>	Apr 18, 2003
AB	+	!	<u>EQ 400MG BASE</u>	<u>N021588</u>	<u>002</u>	Apr 18, 2003

IMATINIB MESYLATE

AB		APOTEX	<u>EQ 100MG BASE</u>	<u>A079179</u>	<u>001</u>	Aug 05, 2016
AB			<u>EQ 400MG BASE</u>	<u>A079179</u>	<u>002</u>	Aug 05, 2016
AB		CHARTWELL RX	<u>EQ 100MG BASE</u>	<u>A208429</u>	<u>001</u>	Jan 17, 2019
AB			<u>EQ 400MG BASE</u>	<u>A208429</u>	<u>002</u>	Jan 17, 2019
AB		DR REDDYS	<u>EQ 100MG BASE</u>	<u>A206547</u>	<u>001</u>	Aug 13, 2018
AB			<u>EQ 400MG BASE</u>	<u>A206547</u>	<u>002</u>	Aug 13, 2018
AB		EUGIA PHARMA	<u>EQ 100MG BASE</u>	<u>A212773</u>	<u>001</u>	Jul 23, 2020
AB			<u>EQ 400MG BASE</u>	<u>A212773</u>	<u>002</u>	Jul 23, 2020
AB		MYLAN	<u>EQ 100MG BASE</u>	<u>A204644</u>	<u>001</u>	Jun 21, 2017
AB			<u>EQ 400MG BASE</u>	<u>A204644</u>	<u>002</u>	Jun 21, 2017
AB		NATCO PHARMA LTD	<u>EQ 100MG BASE</u>	<u>A207818</u>	<u>001</u>	Mar 01, 2019
AB			<u>EQ 400MG BASE</u>	<u>A207818</u>	<u>002</u>	Mar 01, 2019
AB		QILU PHARM HAINAN	<u>EQ 100MG BASE</u>	<u>A212135</u>	<u>001</u>	Jun 21, 2022
AB			<u>EQ 400MG BASE</u>	<u>A212135</u>	<u>002</u>	Jun 21, 2022
AB		SHILPA	<u>EQ 100MG BASE</u>	<u>A208302</u>	<u>001</u>	Jan 17, 2019
AB			<u>EQ 400MG BASE</u>	<u>A208302</u>	<u>002</u>	Jan 17, 2019
AB		SUN PHARM	<u>EQ 100MG BASE</u>	<u>A078340</u>	<u>001</u>	Dec 03, 2015
AB			<u>EQ 400MG BASE</u>	<u>A078340</u>	<u>002</u>	Dec 03, 2015
AB		TEVA PHARMS USA	<u>EQ 100MG BASE</u>	<u>A204285</u>	<u>001</u>	Aug 04, 2016
AB			<u>EQ 400MG BASE</u>	<u>A204285</u>	<u>002</u>	Aug 04, 2016
AB		ZYDUS PHARMS	<u>EQ 100MG BASE</u>	<u>A210658</u>	<u>001</u>	Apr 08, 2020
AB			<u>EQ 400MG BASE</u>	<u>A210658</u>	<u>002</u>	Apr 08, 2020

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

AB		LEADING	<u>10MG</u>	<u>A040903</u>	<u>001</u>	Oct 24, 2012
AB			<u>25MG</u>	<u>A040903</u>	<u>002</u>	Oct 24, 2012
AB			<u>50MG</u>	<u>A040903</u>	<u>003</u>	Oct 24, 2012
AB		OXFORD PHARMS	<u>10MG</u>	<u>A040751</u>	<u>003</u>	Feb 28, 2008
AB			<u>25MG</u>	<u>A040751</u>	<u>002</u>	Feb 28, 2008
AB			<u>50MG</u>	<u>A040751</u>	<u>001</u>	Feb 28, 2008
AB		SANDOZ	<u>10MG</u>	<u>A084936</u>	<u>002</u>	
AB			<u>25MG</u>	<u>A083745</u>	<u>001</u>	
AB			<u>50MG</u>	<u>A084937</u>	<u>001</u>	
AB	+	SPECGX LLC	<u>10MG</u>	<u>A087846</u>	<u>002</u>	May 22, 1984
AB	+		<u>25MG</u>	<u>A087846</u>	<u>003</u>	May 22, 1984
AB		STRIDES PHARMA	<u>10MG</u>	<u>A088292</u>	<u>001</u>	Oct 21, 1983
AB			<u>25MG</u>	<u>A088262</u>	<u>001</u>	Oct 21, 1983
AB			<u>50MG</u>	<u>A088276</u>	<u>001</u>	Oct 21, 1983
AB		SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A081049</u>	<u>001</u>	Jun 05, 1990
AB			<u>50MG</u>	<u>A081050</u>	<u>001</u>	Jun 05, 1990

TOFRANIL

AB	+	!	SPECGX LLC	<u>50MG</u>	<u>A087846</u>	<u>001</u>	May 22, 1984
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PRESCRIPTION DRUG PRODUCT LIST

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

<u>AB</u>	!	HIKMA	<u>EQ 75MG HYDROCHLORIDE</u>	<u>A091099 001</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 100MG HYDROCHLORIDE</u>	<u>A091099 002</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 125MG HYDROCHLORIDE</u>	<u>A091099 003</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 150MG HYDROCHLORIDE</u>	<u>A091099 004</u>	Apr 16, 2010
<u>AB</u>		LUPIN LTD	<u>EQ 75MG HYDROCHLORIDE</u>	<u>A090444 001</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 100MG HYDROCHLORIDE</u>	<u>A090444 002</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 125MG HYDROCHLORIDE</u>	<u>A090444 003</u>	Apr 16, 2010
<u>AB</u>			<u>EQ 150MG HYDROCHLORIDE</u>	<u>A090444 004</u>	Apr 16, 2010

IMIQUIMOD

CREAM; TOPICAL

IMIQUIMOD

<u>AB</u>		APOTEX INC	<u>5%</u>	<u>A091308 001</u>	Apr 06, 2012
<u>AB</u>		FOUGERA PHARMS	<u>5%</u>	<u>A078548 001</u>	Feb 25, 2010
<u>AB</u>	!	GLENMARK GENERICS	<u>5%</u>	<u>A201994 001</u>	Mar 06, 2012
<u>AB</u>		PADAGIS ISRAEL	<u>5%</u>	<u>A078837 001</u>	Sep 07, 2010
<u>AB</u>		TARO	<u>3.75%</u>	<u>A205971 001</u>	Jan 26, 2021
<u>AB</u>			<u>5%</u>	<u>A200173 001</u>	Apr 15, 2011

ZYCLARA

<u>AB</u>	+	BAUSCH	<u>3.75%</u>	<u>N022483 001</u>	Mar 25, 2010
	+		<u>2.5%</u>	<u>N022483 002</u>	Jul 15, 2011

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

!	HIKMA	EQ 5MG BASE/ML	A075513 001	May 09, 2000
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INCLISIRAN SODIUM

SOLUTION; SUBCUTANEOUS

LEQVIO

+	NOVARTIS	EQ 284MG BASE/1.5ML (EQ 189MG BASE/ML)	N214012 001	Dec 22, 2021
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INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>1.25MG</u>	<u>A074722 001</u>	Jun 17, 1996
<u>AB</u>			<u>2.5MG</u>	<u>A074722 002</u>	Jun 17, 1996
<u>AB</u>		ANI PHARMS	<u>1.25MG</u>	<u>A074299 002</u>	Apr 29, 1996
<u>AB</u>	!		<u>2.5MG</u>	<u>A074299 001</u>	Jul 27, 1995
<u>AB</u>		RISING	<u>1.25MG</u>	<u>A074461 002</u>	Mar 26, 1997
<u>AB</u>			<u>2.5MG</u>	<u>A074461 001</u>	Mar 27, 1996

INDIGOTINDISULFONATE SODIUM

SOLUTION; INTRAVENOUS

BLUDIGO

+	PROVEPHARM SAS	40MG/5ML (8MG/ML)	N216264 001	Jul 08, 2022
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INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDIUM IN 111 CHLORIDE

+	CURIUM	5mCi/0.5ML	N019841 001	Sep 27, 1994
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INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN 111 OXYQUINOLINE

<u>AP</u>		BWXT ITG	<u>1mCi/ML</u>	<u>A202586 001</u>	Jul 25, 2018
<u>AP</u>	+	GE HEALTHCARE	<u>1mCi/ML</u>	<u>N019044 001</u>	Dec 24, 1985

INDIUM IN-111 PENTETATE DISODIUM

INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

+	GE HEALTHCARE	1mCi/ML	N017707 001	Feb 18, 1982
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INDIUM IN-111 PENTETREOTIDE KIT

INJECTABLE; INJECTION

OCTREOSCAN

+	CURIUM	3mCi/ML	N020314 001	Jun 02, 1994
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PRESCRIPTION DRUG PRODUCT LIST

INDOCYANINE GREEN

INJECTABLE; INJECTION

INDOCYANINE GREEN

!	RENEW PHARMS	25MG/VIAL	A040811	001	Nov 21, 2007
POWDER; INTRAVENOUS, INTERSTITIAL					
SPY AGENT GREEN KIT					
+	NOVADAQ TECH	25MG/VIAL	N211580	001	Nov 21, 2018

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

AB	CADILA	25MG	A090403	001	Nov 15, 2010
AB		50MG	A090403	002	Nov 15, 2010
AB	CHARTWELL MOLECULES	25MG	N018829	002	Aug 06, 1984
AB		50MG	A070651	001	Mar 05, 1986
AB		50MG	N018829	001	Aug 06, 1984
AB	GLENMARK GENERICS	25MG	A091276	001	Dec 22, 2010
AB	!	50MG	A091276	002	Dec 22, 2010
AB	HETERO LABS LTD III	25MG	A091240	001	Apr 12, 2011
AB		50MG	A091240	002	Apr 12, 2011
AB	SANDOZ	25MG	A070673	001	Apr 29, 1987
AB		50MG	A070674	001	Apr 29, 1987

CAPSULE, EXTENDED RELEASE; ORAL

INDOMETHACIN

AB	AMNEAL PHARMS	75MG	A091549	001	Dec 01, 2010
AB	AVANTHI INC	75MG	A079175	001	Mar 06, 2009
AB	CHARTWELL RX	75MG	A200529	001	Nov 30, 2010
AB	GLENMARK PHARMS LTD	75MG	A203501	001	Jun 22, 2017
AB	!	75MG	A201807	001	Sep 28, 2012
AB	NOVAST LABS	75MG	A204853	001	May 08, 2017
AB	ZYDUS PHARMS	75MG	A202711	001	Sep 25, 2017

INJECTABLE; INJECTION

INDOMETHACIN

+	FRESENIUS KABI USA	EQ 1MG BASE/VIAL	N022536	001	Mar 17, 2010
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SUPPOSITORY; RECTAL

INDOMETHACIN

AB	!	COSETTE	50MG	A073314	001	Aug 31, 1992
AB		ZYDUS	50MG	A216184	001	Aug 02, 2023

SUSPENSION; ORAL

INDOCIN

+	ZYLA	25MG/5ML	N018332	001	Oct 10, 1985
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INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOMETHACIN SODIUM

AP	HOSPIRA	EQ 1MG BASE/VIAL	A204118	001	Apr 19, 2016	
AP	!	NAVINTA LLC	EQ 1MG BASE/VIAL	A206561	001	Jul 19, 2017
AP	WEST-WARD PHARMS INT	EQ 1MG BASE/VIAL	A078713	001	Jul 16, 2008	

INOTERSEN SODIUM

SOLUTION; SUBCUTANEOUS

TEGSEDI

+	AKCEA THERAPS	EQ 284MG BASE/1.5ML (EQ 189.3MG BASE/ML)	N211172	001	Oct 05, 2018
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IOBENGUANE I-131

SOLUTION; INTRAVENOUS

AZEDRA

+	PROGENICS PHARMS INC	15mCi/ML	N209607	001	Jul 30, 2018
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IOBENGUANE SULFATE I-123

SOLUTION; INTRAVENOUS

ADREVIEW

+	GE HEALTHCARE	10mCi/5ML (2mCi/ML)	N022290	001	Sep 19, 2008
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IODIXANOL

INJECTABLE; INJECTION

IODIXANOL

AP	HENGRUI PHARMA	55%	A214271	001	May 19, 2022	
AP		65.2%	A214271	002	May 19, 2022	
VISIPAQUE 270						
AP	+	GE HEALTHCARE	55%	N020351	001	Mar 22, 1996

PRESCRIPTION DRUG PRODUCT LIST

IODIXANOL

INJECTABLE; INJECTION

VISIPAQUE 320

AP	+ !	GE HEALTHCARE	65.2%	N020351 002	Mar 22, 1996
			65.2%	N020808 002	Aug 29, 1997

IOFLUPANE I-123

SOLUTION; INTRAVENOUS

DATSCAN

AP	+ !	GE HLTHCARE INC	5mCi/2.5ML (2mCi/ML)	N022454 001	Jan 14, 2011
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IOFLUPANE I-123

AP		CURIUM	5mCi/2.5ML (2mCi/ML)	A213792 001	Mar 30, 2022
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IOHEXOL

INJECTABLE; INJECTION

OMNIPAQUE 140

+ !	GE HEALTHCARE	30.2%	N018956 005	Nov 30, 1988
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SOLUTION; INJECTION, ORAL

OMNIPAQUE 350

+ !	GE HEALTHCARE	75.5%	N018956 004	Dec 26, 1985
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		75.5%	N020608 003	Oct 24, 1995
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SOLUTION; INJECTION, ORAL, RECTAL

OMNIPAQUE 180

+ !	GE HEALTHCARE	38.8%	N018956 001	Dec 26, 1985
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OMNIPAQUE 240

+ !	GE HEALTHCARE	51.8%	N018956 002	Dec 26, 1985
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OMNIPAQUE 300

+ !	GE HEALTHCARE	64.7%	N018956 003	Dec 26, 1985
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		64.7%	N020608 002	Oct 24, 1995
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SOLUTION; ORAL

OMNIPAQUE 12

+ !	GE HEALTHCARE	2.6%	N018956 009	Apr 17, 2018
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OMNIPAQUE 9

+ !	GE HEALTHCARE	1.9%	N018956 008	Apr 17, 2018
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IOPAMIDOL

INJECTABLE; INJECTION

IOPAMIDOL

AP		HAINAN POLY	61%	A215382 002	Feb 27, 2023
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ISOVUE-M 300

AP	+ !	BRACCO	61%	N018735 004	Dec 31, 1985
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IOPAMIDOL

AP1		HAINAN POLY	41%	A215382 001	Feb 27, 2023
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ISOVUE-M 200

AP1	+ !	BRACCO	41%	N018735 001	Dec 31, 1985
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IOPAMIDOL

AP2		HAINAN POLY	41%	A217134 001	Sep 27, 2023
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ISOVUE-200

AP2	+ !	BRACCO	41%	N018735 006	Jul 07, 1987
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ISOVUE-250

+ !	BRACCO	51%	N018735 007	Jul 06, 1992
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ISOVUE-300

+ !	BRACCO	61%	N018735 002	Dec 31, 1985
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+ !		61%	N020327 003	Oct 12, 1994
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ISOVUE-370

+ !	BRACCO	76%	N018735 003	Dec 31, 1985
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+ !		76%	N020327 004	Oct 12, 1994
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IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+ !	BAYER HLTHCARE	62.3%	N021425 001	Sep 20, 2002
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+ !		76.9%	N021425 002	Sep 20, 2002
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ULTRAVIST 300

+ !	BAYER HLTHCARE	62.3%	N020220 002	May 10, 1995
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ULTRAVIST 370

+ !	BAYER HLTHCARE	76.9%	N020220 001	May 10, 1995
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IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY

+ !	LIEBEL-FLARSHEIM	60%	N013295 001	
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PRESCRIPTION DRUG PRODUCT LIST

IOTHALAMATE MEGLUMINESOLUTION; INTRAVESICAL
CYSTO-CONRAY II

LIEBEL-FLARSHEIM 17.2% N017057 002

IOTHALAMATE SODIUM I-125INJECTABLE; INJECTION
GLOFIL-125

ISOTEX 250-300uCi/ML N017279 001

IOVERSOLINJECTABLE; INJECTION
OPTIRAY 300+! LIEBEL-FLARSHEIM 64% N019710 004 Jan 22, 1992
+! 64% N020923 004 May 13, 1999

OPTIRAY 320

+! LIEBEL-FLARSHEIM 68% N019710 001 Dec 30, 1988
+! 68% N020923 002 May 29, 1998

OPTIRAY 350

+! LIEBEL-FLARSHEIM 74% N019710 005 Jan 22, 1992
+! 74% N020923 003 May 28, 1998IPRATROPIUM BROMIDEAEROSOL, METERED; INHALATION
ATROVENT HFA+! BOEHRINGER 0.021MG/INH N021527 001 Nov 27, 2004
INGELHEIM

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

<u>AN</u>	LUOXIN AUROVITAS	<u>0.02%</u>	<u>A206543</u>	<u>001</u>	Oct 27, 2016
<u>AN</u>	NEPHRON	<u>0.02%</u>	<u>A075562</u>	<u>001</u>	Sep 27, 2001
<u>AN</u>	! RITEDOSE CORP	<u>0.02%</u>	<u>A075693</u>	<u>001</u>	Jan 26, 2001
<u>AN</u>	SUN PHARM	<u>0.02%</u>	<u>A207903</u>	<u>001</u>	Jan 03, 2017

SPRAY, METERED; NASAL

IPRATROPIUM BROMIDE

<u>AB</u>	AMNEAL	<u>0.021MG/SPRAY</u>	<u>A215104</u>	<u>001</u>	Aug 16, 2022
<u>AB</u>		<u>0.042MG/SPRAY</u>	<u>A215105</u>	<u>001</u>	Aug 16, 2022
<u>AB</u>	APOTEX INC	<u>0.021MG/SPRAY</u>	<u>A076156</u>	<u>001</u>	Apr 18, 2003
<u>AB</u>		<u>0.042MG/SPRAY</u>	<u>A076155</u>	<u>001</u>	Apr 18, 2003
<u>AB</u>	BAUSCH	<u>0.021MG/SPRAY</u>	<u>A076025</u>	<u>001</u>	Mar 31, 2003
<u>AB</u>		<u>0.042MG/SPRAY</u>	<u>A076103</u>	<u>001</u>	Mar 31, 2003
<u>AB</u>	! HIKMA	<u>0.021MG/SPRAY</u>	<u>A076664</u>	<u>001</u>	Nov 05, 2003
<u>AB</u>	!	<u>0.042MG/SPRAY</u>	<u>A076598</u>	<u>001</u>	Nov 05, 2003

IPTACOPAN HYDROCHLORIDE

CAPSULE; ORAL

FABHALTA

+! NOVARTIS EQ 200MG BASE N218276 001 Dec 05, 2023

IRBESARTAN

TABLET; ORAL

AVAPRO

<u>AB</u>	+	SANOFI AVENTIS US	<u>75MG</u>	<u>N020757</u>	<u>001</u>	Sep 30, 1997
<u>AB</u>	+		<u>150MG</u>	<u>N020757</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+!		<u>300MG</u>	<u>N020757</u>	<u>003</u>	Sep 30, 1997

IRBESARTAN

<u>AB</u>		ALEMBIC PHARMS LTD	<u>75MG</u>	<u>A091236</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>			<u>150MG</u>	<u>A091236</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>			<u>300MG</u>	<u>A091236</u>	<u>003</u>	Oct 15, 2012
<u>AB</u>		AMNEAL PHARMS	<u>75MG</u>	<u>A204740</u>	<u>001</u>	Apr 17, 2018
<u>AB</u>			<u>150MG</u>	<u>A204740</u>	<u>002</u>	Apr 17, 2018
<u>AB</u>			<u>300MG</u>	<u>A204740</u>	<u>003</u>	Apr 17, 2018
<u>AB</u>		AUROBINDO PHARMA LTD	<u>75MG</u>	<u>A203081</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>150MG</u>	<u>A203081</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>			<u>300MG</u>	<u>A203081</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>		CHARTWELL MOLECULAR	<u>75MG</u>	<u>A077205</u>	<u>001</u>	Nov 14, 2012
<u>AB</u>			<u>150MG</u>	<u>A077205</u>	<u>002</u>	Nov 14, 2012
<u>AB</u>			<u>300MG</u>	<u>A077205</u>	<u>003</u>	Nov 14, 2012
<u>AB</u>		HETERO LABS LTD V	<u>75MG</u>	<u>A202910</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>			<u>150MG</u>	<u>A202910</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>			<u>300MG</u>	<u>A202910</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>		HISUN PHARM	<u>75MG</u>	<u>A206194</u>	<u>001</u>	Jun 14, 2016
<u>AB</u>		HANGZHOU	<u>150MG</u>	<u>A206194</u>	<u>002</u>	Jun 14, 2016

PRESCRIPTION DRUG PRODUCT LIST

IRBESARTAN

TABLET; ORAL

IRBESARTAN

<u>AB</u>		<u>300MG</u>	<u>A206194</u>	<u>003</u>	Jun 14, 2016
<u>AB</u>	JUBILANT GENERICS	<u>75MG</u>	<u>A203534</u>	<u>001</u>	Feb 23, 2015
<u>AB</u>		<u>150MG</u>	<u>A203534</u>	<u>002</u>	Feb 23, 2015
<u>AB</u>		<u>300MG</u>	<u>A203534</u>	<u>003</u>	Feb 23, 2015
<u>AB</u>	LUPIN LTD	<u>75MG</u>	<u>A201531</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>		<u>150MG</u>	<u>A201531</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>		<u>300MG</u>	<u>A201531</u>	<u>003</u>	Oct 15, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>75MG</u>	<u>A202254</u>	<u>001</u>	Oct 03, 2012
<u>AB</u>		<u>150MG</u>	<u>A202254</u>	<u>002</u>	Oct 03, 2012
<u>AB</u>		<u>300MG</u>	<u>A202254</u>	<u>003</u>	Oct 03, 2012
<u>AB</u>	PRINSTON INC	<u>75MG</u>	<u>A203071</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A203071</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A203071</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>	SANDOZ	<u>75MG</u>	<u>A077466</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A077466</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A077466</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>	SCIEGEN PHARMS INC	<u>75MG</u>	<u>A204774</u>	<u>001</u>	Dec 07, 2015
<u>AB</u>		<u>150MG</u>	<u>A204774</u>	<u>002</u>	Dec 07, 2015
<u>AB</u>		<u>300MG</u>	<u>A204774</u>	<u>003</u>	Dec 07, 2015
<u>AB</u>	TEVA PHARMS	<u>75MG</u>	<u>A077159</u>	<u>001</u>	Mar 30, 2012
<u>AB</u>		<u>150MG</u>	<u>A077159</u>	<u>002</u>	Mar 30, 2012
<u>AB</u>		<u>300MG</u>	<u>A077159</u>	<u>003</u>	Mar 30, 2012
<u>AB</u>	UNICHEM	<u>75MG</u>	<u>A203020</u>	<u>001</u>	Dec 07, 2015
<u>AB</u>		<u>150MG</u>	<u>A203020</u>	<u>002</u>	Dec 07, 2015
<u>AB</u>		<u>300MG</u>	<u>A203020</u>	<u>003</u>	Dec 07, 2015
<u>AB</u>	ZYDUS PHARMS USA INC	<u>75MG</u>	<u>A079213</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A079213</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A079213</u>	<u>003</u>	Sep 27, 2012

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

CAMPTOSAR

<u>AP</u>	<u>+</u> !	PFIZER INC	<u>40MG/2ML (20MG/ML)</u>	<u>N020571</u>	<u>001</u>	Jun 14, 1996
<u>AP</u>	<u>+</u> !		<u>100MG/5ML (20MG/ML)</u>	<u>N020571</u>	<u>002</u>	Jun 14, 1996
<u>AP</u>	<u>+</u> !		<u>300MG/15ML (20MG/ML)</u>	<u>N020571</u>	<u>003</u>	Aug 05, 2010

IRINOTECAN HYDROCHLORIDE

<u>AP</u>		ACCORD HLTHCARE	<u>40MG/2ML (20MG/ML)</u>	<u>A079068</u>	<u>001</u>	Nov 21, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A079068</u>	<u>002</u>	Nov 21, 2008
<u>AP</u>		ACTAVIS TOTOWA	<u>40MG/2ML (20MG/ML)</u>	<u>A078589</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A078589</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>			<u>500MG/25ML (20MG/ML)</u>	<u>A078589</u>	<u>003</u>	Nov 18, 2015
<u>AP</u>		AKORN	<u>40MG/2ML (20MG/ML)</u>	<u>A090726</u>	<u>001</u>	Sep 16, 2009
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A090726</u>	<u>002</u>	Sep 16, 2009
<u>AP</u>		EUGIA PHARMA	<u>40MG/2ML (20MG/ML)</u>	<u>A213278</u>	<u>001</u>	Nov 02, 2020
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A213278</u>	<u>002</u>	Nov 02, 2020
<u>AP</u>			<u>300MG/15ML (20MG/ML)</u>	<u>A213278</u>	<u>003</u>	Nov 02, 2020
<u>AP</u>			<u>500MG/25ML (20MG/ML)</u>	<u>A213278</u>	<u>004</u>	Apr 11, 2022
<u>AP</u>		FRESENIUS KABI USA	<u>40MG/2ML (20MG/ML)</u>	<u>A077776</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A077776</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>		GLAND PHARMA LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A212993</u>	<u>001</u>	Nov 18, 2019
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A212993</u>	<u>002</u>	Nov 18, 2019
<u>AP</u>		HENGRUI PHARMA	<u>40MG/2ML (20MG/ML)</u>	<u>A090675</u>	<u>002</u>	Dec 16, 2011
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A090675</u>	<u>001</u>	Dec 16, 2011
<u>AP</u>		HIKMA FARMACEUTICA	<u>40MG/2ML (20MG/ML)</u>	<u>A091032</u>	<u>001</u>	Dec 20, 2010
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A091032</u>	<u>002</u>	Dec 20, 2010
<u>AP</u>		HISUN PHARM HANGZHOU	<u>40MG/2ML (20MG/ML)</u>	<u>A090016</u>	<u>001</u>	Jan 28, 2009
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A090016</u>	<u>002</u>	Jan 28, 2009
<u>AP</u>		HOSPIRA	<u>40MG/2ML (20MG/ML)</u>	<u>A077915</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A077915</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>	<u>!</u>		<u>500MG/25ML (20MG/ML)</u>	<u>A078796</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>		INTAS PHARMS USA	<u>40MG/2ML (20MG/ML)</u>	<u>A203054</u>	<u>001</u>	Aug 31, 2017
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A203054</u>	<u>002</u>	Aug 31, 2017
<u>AP</u>		NOVAST LABS	<u>40MG/2ML (20MG/ML)</u>	<u>A206935</u>	<u>001</u>	May 26, 2017
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A206935</u>	<u>002</u>	May 26, 2017
<u>AP</u>		QILU PHARM HAINAN	<u>40MG/2ML (20MG/ML)</u>	<u>A203380</u>	<u>001</u>	May 03, 2016
<u>AP</u>			<u>100MG/5ML (20MG/ML)</u>	<u>A203380</u>	<u>002</u>	May 03, 2016
<u>AP</u>			<u>300MG/15ML (20MG/ML)</u>	<u>A203380</u>	<u>003</u>	May 03, 2016
<u>AP</u>		SHILPA	<u>40MG/2ML (20MG/ML)</u>	<u>A208718</u>	<u>001</u>	Dec 28, 2018

PRESCRIPTION DRUG PRODUCT LIST

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A208718 002</u>	Dec 28, 2018
<u>AP</u>		<u>300MG/15ML (20MG/ML)</u>	<u>A208718 003</u>	Aug 16, 2019
<u>AP</u>	TEVA PHARMS USA	<u>500MG/25ML (20MG/ML)</u>	<u>A090101 001</u>	Nov 26, 2008
<u>AP</u>	WEST-WARD PHARMS INT	<u>40MG/2ML (20MG/ML)</u>	<u>A078753 001</u>	Dec 24, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078753 002</u>	Dec 24, 2008
<u>AP</u>	ZENNOVA	<u>40MG/2ML (20MG/ML)</u>	<u>A090393 002</u>	May 13, 2011
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090393 003</u>	May 13, 2011

INJECTABLE, LIPOSOMAL; INTRAVENOUS

ONIVYDE

+	!	IPSEN	EQ 43MG BASE/10ML (EQ 4.3MG BASE/ML)	N207793 001	Oct 22, 2015
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ISAVUCONAZONIUM SULFATE

CAPSULE; ORAL

CRESEMBA

+		ASTELLAS	74.5MG	N207500 002	Nov 22, 2022
+	!		186MG	N207500 001	Mar 06, 2015

POWDER; INTRAVENOUS

CRESEMBA

+	!	ASTELLAS	372MG	N207501 001	Mar 06, 2015
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ISOCARBOXAZID

TABLET; ORAL

MARPLAN

+	!	VALIDUS PHARMS INC	10MG	N011961 001	
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ISOFLURANE

LIQUID; INHALATION

FORANE

<u>AN</u>	+	!	BAXTER HLTHCARE	<u>99.9%</u>	<u>N017624 001</u>
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ISOFLURANE

<u>AN</u>			HALOCARBON PRODS	<u>99.9%</u>	<u>A075225 001</u>	Oct 20, 1999
<u>AN</u>			PIRAMAL CRITICAL	<u>99.9%</u>	<u>A074416 001</u>	Sep 30, 1994
<u>AN</u>			PIRAMAL PHARMA	<u>99.9%</u>	<u>A074502 001</u>	Jun 27, 1995

ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

!		SANDOZ	100MG/ML	A040648 001	Jul 05, 2005
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SYRUP; ORAL

ISONIAZID

+	!	CMP PHARMA INC	50MG/5ML	A088235 001	Nov 10, 1983
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TABLET; ORAL

ISONIAZID

<u>AA</u>	!		BARR	<u>100MG</u>	<u>A080936 001</u>	
<u>AA</u>	!			<u>300MG</u>	<u>A080937 002</u>	
<u>AA</u>			OMNIVIUM PHARMS	<u>100MG</u>	<u>A040090 001</u>	Jun 26, 1997
<u>AA</u>				<u>300MG</u>	<u>A040090 002</u>	Jun 26, 1997
<u>AA</u>			THEPHARMANETWORK LLC	<u>100MG</u>	<u>A202610 001</u>	Oct 29, 2014
<u>AA</u>				<u>300MG</u>	<u>A202610 002</u>	Oct 29, 2014

ISOPROTERENOL HYDROCHLORIDE

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

<u>AP</u>			AMNEAL	<u>0.2MG/ML</u>	<u>A210576 001</u>	Oct 17, 2018
<u>AP</u>			AMPHASTAR PHARMS INC	<u>0.2MG/ML</u>	<u>A210106 001</u>	Jun 18, 2018
<u>AP</u>			AMRING PHARMS	<u>0.2MG/ML</u>	<u>A211237 001</u>	May 19, 2021
<u>AP</u>			AVET LIFESCIENCES	<u>0.2MG/ML</u>	<u>A212189 001</u>	Nov 19, 2021
<u>AP</u>			CIPLA	<u>0.2MG/ML</u>	<u>A210322 001</u>	Jun 12, 2018
<u>AP</u>			EUGIA PHARMA	<u>0.2MG/ML</u>	<u>A211864 001</u>	Feb 09, 2021
<u>AP</u>			HIKMA	<u>0.2MG/ML</u>	<u>A211703 001</u>	Apr 26, 2023
<u>AP</u>			MICRO LABS	<u>0.2MG/ML</u>	<u>A210845 001</u>	Feb 16, 2021
<u>AP</u>	!		NEXUS	<u>0.2MG/ML</u>	<u>A206961 001</u>	Aug 02, 2017
<u>AP</u>			PENN LIFE	<u>0.2MG/ML</u>	<u>A215542 001</u>	Sep 20, 2022

PRESCRIPTION DRUG PRODUCT LIST

ISOSORBIDE DINITRATE

TABLET; ORAL

ISORDIL

<u>AB</u>	+	BAUSCH	<u>5MG</u>	<u>N012093</u>	<u>007</u>	Jul 29, 1988
<u>AB</u>	+	!	<u>40MG</u>	<u>N012093</u>	<u>001</u>	Jul 29, 1988

ISOSORBIDE DINITRATE

<u>AB</u>		HIKMA INTL PHARMS	<u>5MG</u>	<u>A086067</u>	<u>001</u>	Oct 29, 1987
<u>AB</u>			<u>10MG</u>	<u>A086066</u>	<u>001</u>	Oct 29, 1987
<u>AB</u>			<u>20MG</u>	<u>A088088</u>	<u>001</u>	Nov 02, 1987
<u>AB</u>		PAR PHARM	<u>5MG</u>	<u>A086923</u>	<u>001</u>	Mar 12, 1987
<u>AB</u>			<u>10MG</u>	<u>A086925</u>	<u>001</u>	Mar 12, 1987
<u>AB</u>			<u>20MG</u>	<u>A087537</u>	<u>001</u>	Oct 02, 1987
<u>AB</u>	!		<u>30MG</u>	<u>A087946</u>	<u>001</u>	Jan 12, 1988
<u>AB</u>		PAR PHARM INC	<u>40MG</u>	<u>A211290</u>	<u>001</u>	Nov 23, 2021
<u>AB</u>		RUBICON	<u>5MG</u>	<u>A215723</u>	<u>001</u>	Jul 08, 2022
<u>AB</u>			<u>10MG</u>	<u>A215723</u>	<u>002</u>	Jul 08, 2022
<u>AB</u>			<u>20MG</u>	<u>A215723</u>	<u>003</u>	Jul 08, 2022
<u>AB</u>			<u>30MG</u>	<u>A215723</u>	<u>004</u>	Jul 08, 2022
<u>AB</u>			<u>40MG</u>	<u>A215723</u>	<u>005</u>	Jul 08, 2022
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A086221</u>	<u>001</u>	Jan 07, 1988
<u>AB</u>			<u>10MG</u>	<u>A086223</u>	<u>001</u>	Jan 07, 1988
<u>AB</u>			<u>20MG</u>	<u>A089367</u>	<u>001</u>	Apr 07, 1988
<u>AB</u>		ZYDUS	<u>5MG</u>	<u>A213057</u>	<u>001</u>	Nov 20, 2019
<u>AB</u>			<u>10MG</u>	<u>A213057</u>	<u>002</u>	Nov 20, 2019
<u>AB</u>			<u>20MG</u>	<u>A213057</u>	<u>003</u>	Nov 20, 2019
<u>AB</u>			<u>30MG</u>	<u>A213057</u>	<u>004</u>	Nov 20, 2019
<u>AB</u>			<u>40MG</u>	<u>A213057</u>	<u>005</u>	Nov 20, 2019

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>10MG</u>	<u>A075037</u>	<u>002</u>	Oct 30, 1998
<u>AB</u>			<u>20MG</u>	<u>A075037</u>	<u>001</u>	Oct 30, 1998

MONOKET

<u>AB</u>	+	GENUS LIFESCIENCES	<u>10MG</u>	<u>N020215</u>	<u>002</u>	Jun 30, 1993
<u>AB</u>	+	!	<u>20MG</u>	<u>N020215</u>	<u>001</u>	Jun 30, 1993

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>		AUROBINDO PHARMA	<u>30MG</u>	<u>A216557</u>	<u>001</u>	Nov 07, 2022
<u>AB</u>			<u>60MG</u>	<u>A216557</u>	<u>002</u>	Nov 07, 2022
<u>AB</u>			<u>120MG</u>	<u>A216557</u>	<u>003</u>	Nov 07, 2022
<u>AB</u>		CHARTWELL MOLECULAR	<u>30MG</u>	<u>A075155</u>	<u>002</u>	Jan 13, 2000
<u>AB</u>			<u>60MG</u>	<u>A075155</u>	<u>001</u>	Oct 30, 1998
<u>AB</u>	!		<u>120MG</u>	<u>A075155</u>	<u>003</u>	Aug 04, 2000
<u>AB</u>		DEXCEL LTD	<u>30MG</u>	<u>A075522</u>	<u>002</u>	Sep 20, 2016
<u>AB</u>			<u>60MG</u>	<u>A075522</u>	<u>001</u>	Apr 17, 2000
<u>AB</u>			<u>120MG</u>	<u>A210822</u>	<u>001</u>	Aug 29, 2018
<u>AB</u>		RICONPHARMA LLC	<u>30MG</u>	<u>A210918</u>	<u>001</u>	Nov 05, 2018
<u>AB</u>			<u>60MG</u>	<u>A210918</u>	<u>002</u>	Nov 05, 2018
<u>AB</u>			<u>120MG</u>	<u>A210918</u>	<u>003</u>	Nov 05, 2018
<u>AB</u>		TORRENT PHARMS	<u>30MG</u>	<u>A200270</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>			<u>60MG</u>	<u>A200495</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>			<u>120MG</u>	<u>A200495</u>	<u>002</u>	Jun 03, 2011
<u>AB</u>		ZYDUS HLTHCARE	<u>30MG</u>	<u>A075395</u>	<u>001</u>	Mar 16, 2000
<u>AB</u>			<u>60MG</u>	<u>A075395</u>	<u>002</u>	Mar 16, 2000
<u>AB</u>			<u>120MG</u>	<u>A075395</u>	<u>003</u>	Mar 16, 2000

ISOSULFAN BLUE

INJECTABLE; INJECTION

ISOSULFAN BLUE

<u>AP</u>		AM REGENT	<u>1%</u>	<u>A210294</u>	<u>001</u>	May 17, 2023
<u>AP</u>		MEITHEAL	<u>1%</u>	<u>A213130</u>	<u>001</u>	Nov 03, 2021
<u>AP</u>		MSN	<u>1%</u>	<u>A216090</u>	<u>001</u>	Sep 16, 2022
<u>AP</u>	!	MYLAN INSTITUTIONAL	<u>1%</u>	<u>A090874</u>	<u>001</u>	Jul 20, 2010

ISOTRETINOIN

CAPSULE; ORAL

AMNESTEEM

<u>AB1</u>		MYLAN PHARMS INC	<u>10MG</u>	<u>A075945</u>	<u>001</u>	Nov 08, 2002
<u>AB1</u>			<u>20MG</u>	<u>A075945</u>	<u>002</u>	Nov 08, 2002
<u>AB1</u>			<u>40MG</u>	<u>A075945</u>	<u>003</u>	Nov 08, 2002

CLARAVIS

<u>AB1</u>		TEVA PHARMS USA	<u>10MG</u>	<u>A076356</u>	<u>001</u>	Apr 11, 2003
<u>AB1</u>			<u>20MG</u>	<u>A076135</u>	<u>002</u>	Apr 11, 2003

PRESCRIPTION DRUG PRODUCT LIST

ISOTRETINOIN

CAPSULE; ORAL

CLARAVIS

<u>AB1</u>		<u>30MG</u>	<u>A076135</u>	<u>003</u>	May 11, 2006
<u>AB1</u>	!	<u>40MG</u>	<u>A076135</u>	<u>001</u>	Apr 11, 2003

ISOTRETINOIN

<u>AB1</u>	AMNEAL PHARMS NY	<u>10MG</u>	<u>A207792</u>	<u>001</u>	Sep 29, 2017
<u>AB1</u>		<u>20MG</u>	<u>A207792</u>	<u>002</u>	Sep 29, 2017
<u>AB1</u>		<u>30MG</u>	<u>A207792</u>	<u>003</u>	Sep 29, 2017
<u>AB1</u>		<u>40MG</u>	<u>A207792</u>	<u>004</u>	Sep 29, 2017
<u>AB1</u>	ZYDUS PHARMS	<u>10MG</u>	<u>A211568</u>	<u>001</u>	Aug 29, 2023
<u>AB1</u>		<u>20MG</u>	<u>A211568</u>	<u>002</u>	Aug 29, 2023
<u>AB1</u>		<u>30MG</u>	<u>A211568</u>	<u>003</u>	Aug 29, 2023
<u>AB1</u>		<u>40MG</u>	<u>A211568</u>	<u>004</u>	Aug 29, 2023

MYORISAN

<u>AB1</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A076485</u>	<u>001</u>	Jan 19, 2012
<u>AB1</u>		<u>20MG</u>	<u>A076485</u>	<u>002</u>	Jan 19, 2012
<u>AB1</u>		<u>30MG</u>	<u>A076485</u>	<u>004</u>	Aug 25, 2015
<u>AB1</u>		<u>40MG</u>	<u>A076485</u>	<u>003</u>	Jan 19, 2012

ZENATANE

<u>AB1</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A202099</u>	<u>001</u>	Mar 25, 2013
<u>AB1</u>		<u>20MG</u>	<u>A202099</u>	<u>002</u>	Mar 25, 2013
<u>AB1</u>		<u>30MG</u>	<u>A202099</u>	<u>004</u>	Feb 23, 2015
<u>AB1</u>		<u>40MG</u>	<u>A202099</u>	<u>003</u>	Mar 25, 2013

ABSORICA

<u>AB2</u>	+ SUN PHARM INDS INC	<u>10MG</u>	<u>N021951</u>	<u>001</u>	May 25, 2012
<u>AB2</u>	+	<u>20MG</u>	<u>N021951</u>	<u>002</u>	May 25, 2012
<u>AB2</u>	+	<u>25MG</u>	<u>N021951</u>	<u>005</u>	Aug 15, 2014
<u>AB2</u>	+	<u>30MG</u>	<u>N021951</u>	<u>003</u>	May 25, 2012
<u>AB2</u>	+	<u>35MG</u>	<u>N021951</u>	<u>006</u>	Aug 15, 2014
<u>AB2</u>	+	<u>40MG</u>	<u>N021951</u>	<u>004</u>	May 25, 2012

ISOTRETINOIN

<u>AB2</u>	ACTAVIS LABS FL	<u>10MG</u>	<u>A205063</u>	<u>001</u>	Mar 31, 2021
<u>AB2</u>		<u>20MG</u>	<u>A205063</u>	<u>002</u>	Mar 31, 2021
<u>AB2</u>		<u>25MG</u>	<u>A205063</u>	<u>003</u>	Mar 31, 2021
<u>AB2</u>		<u>30MG</u>	<u>A205063</u>	<u>004</u>	Mar 31, 2021
<u>AB2</u>		<u>35MG</u>	<u>A205063</u>	<u>005</u>	Mar 31, 2021
<u>AB2</u>		<u>40MG</u>	<u>A205063</u>	<u>006</u>	Mar 31, 2021
<u>AB2</u>	UPSHER SMITH LABS	<u>10MG</u>	<u>A212333</u>	<u>001</u>	Sep 21, 2021
<u>AB2</u>		<u>20MG</u>	<u>A212333</u>	<u>002</u>	Sep 21, 2021
<u>AB2</u>		<u>30MG</u>	<u>A212333</u>	<u>003</u>	Sep 21, 2021
<u>AB2</u>		<u>40MG</u>	<u>A213571</u>	<u>001</u>	Apr 12, 2021
	ABSORICA LD				
	+ SUN PHARM	8MG	N211913	001	Nov 05, 2019
	+	16MG	N211913	002	Nov 05, 2019
	+	24MG	N211913	004	Nov 05, 2019
	+	32MG	N211913	006	Nov 05, 2019

ISRADIPINE

CAPSULE; ORAL

ISRADIPINE

<u>AB</u>	ELITE LABS INC	<u>2.5MG</u>	<u>A077169</u>	<u>001</u>	Apr 24, 2006
<u>AB</u>		<u>5MG</u>	<u>A077169</u>	<u>002</u>	Apr 24, 2006
<u>AB</u>	WATSON LABS TEVA	<u>2.5MG</u>	<u>A077317</u>	<u>001</u>	Jan 05, 2006
<u>AB</u>	!	<u>5MG</u>	<u>A077317</u>	<u>002</u>	Jan 05, 2006

ISTRADIFYLLINE

TABLET; ORAL

NOURIANZ

	+ KYOWA KIRIN	20MG	N022075	001	Aug 27, 2019
	+	40MG	N022075	002	Aug 27, 2019

ITRACONAZOLE

CAPSULE; ORAL

ITRACONAZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>100MG</u>	<u>A205991</u>	<u>001</u>	May 26, 2016
<u>AB</u>	ALEMBIC	<u>100MG</u>	<u>A206741</u>	<u>001</u>	Dec 13, 2016
<u>AB</u>	ALKEM LABS LTD	<u>100MG</u>	<u>A208591</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>	AMNEAL PHARMS	<u>100MG</u>	<u>A205080</u>	<u>001</u>	Sep 26, 2016
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A076104</u>	<u>001</u>	May 28, 2004
<u>AB</u>	TORRENT	<u>100MG</u>	<u>A209460</u>	<u>001</u>	Aug 24, 2018
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A204672</u>	<u>001</u>	Sep 19, 2017

PRESCRIPTION DRUG PRODUCT LIST

ITRACONAZOLE

CAPSULE; ORAL

SPORANOX

AB	+!	JANSSEN PHARMS	100MG	N020083	001	Sep 11, 1992
		TOLSURA				
	+!	MAYNE PHARMA	65MG	N208901	001	Dec 11, 2018

SOLUTION; ORAL

ITRACONAZOLE

AA		AMNEAL PHARMS	10MG/ML	A205573	001	Oct 30, 2015
AA		ANNORA PHARMA	10MG/ML	A212239	001	Sep 01, 2020

SPORANOX

AA	+!	JANSSEN PHARMS	10MG/ML	N020657	001	Feb 21, 1997
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IVABRADINE

SOLUTION; ORAL

CORLANOR

+!	AMGEN INC	5MG/5ML (1MG/ML)	N209964	001	Apr 22, 2019
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IVABRADINE HYDROCHLORIDE

TABLET; ORAL

CORLANOR

AB	+	AMGEN INC	EQ 5MG BASE	N206143	001	Apr 15, 2015
AB	+!		EQ 7.5MG BASE	N206143	002	Apr 15, 2015

IVABRADINE HYDROCHLORIDE

AB		ANNORA PHARMA	EQ 5MG BASE	A213366	001	Oct 05, 2022
AB			EQ 7.5MG BASE	A213366	002	Oct 05, 2022
AB		ZYDUS PHARMS	EQ 5MG BASE	A213442	001	Nov 29, 2023
AB			EQ 7.5MG BASE	A213442	002	Nov 29, 2023

IVACAFTOR

GRANULE; ORAL

KALYDECO

+	VERTEX PHARMS INC	5.8MG/PACKET	N207925	004	May 03, 2023
+		13.4MG/PACKET	N207925	005	May 03, 2023
+		25MG/PACKET	N207925	003	Apr 29, 2019
+		50MG/PACKET	N207925	001	Mar 17, 2015
+!		75MG/PACKET	N207925	002	Mar 17, 2015

TABLET; ORAL

KALYDECO

+!	VERTEX PHARMS	150MG	N203188	001	Jan 31, 2012
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IVACAFTOR; IVACAFTOR, TEZACAFTOR

TABLET; ORAL

SYMDEKO (COPACKAGED)

+	VERTEX PHARMS INC	75MG;75MG, 50MG	N210491	002	Jun 21, 2019
+!		150MG;150MG, 100MG	N210491	001	Feb 12, 2018

IVACAFTOR; LUMACAFTOR

GRANULE; ORAL

ORKAMBI

+	VERTEX PHARMS INC	94MG/PACKET;75MG/PACKET	N211358	003	Sep 02, 2022
+		125MG/PACKET;100MG/PACKET	N211358	001	Aug 07, 2018
+!		188MG/PACKET;150MG/PACKET	N211358	002	Aug 07, 2018

TABLET; ORAL

ORKAMBI

+	VERTEX PHARMS INC	125MG;100MG	N206038	002	Sep 28, 2016
+!		125MG;200MG	N206038	001	Jul 02, 2015

IVERMECTIN

CREAM; TOPICAL

IVERMECTIN

AB		PADAGIS ISRAEL	1%	A210225	001	Apr 13, 2020
AB		TEVA PHARMS USA	1%	A210019	001	Sep 13, 2019
AB		ZYDUS	1%	A215210	001	Aug 02, 2022

SOOLANTRA

AB	+!	GALDERMA LABS LP	1%	N206255	001	Dec 19, 2014
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TABLET; ORAL

IVERMECTIN

AB		EDENBRIDGE PHARMS	3MG	A204154	001	Oct 24, 2014
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STROMECTOL

AB	+!	MERCK SHARP DOHME	3MG	N050742	002	Oct 08, 1998
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PRESCRIPTION DRUG PRODUCT LIST

IVOSIDENIB

TABLET; ORAL

TIBSOVO

+! SERVIER 250MG N211192 001 Jul 20, 2018

IXABEPILONE

INJECTABLE; INTRAVENOUS

IXEMPRA KIT

+! R-PHARM US LLC 15MG/VIAL N022065 001 Oct 16, 2007

+! 45MG/VIAL N022065 002 Oct 16, 2007

IXAZOMIB CITRATE

CAPSULE; ORAL

NINLARO

+ TAKEDA PHARMS USA EQ 2.3MG BASE N208462 001 Nov 20, 2015

+ EQ 3MG BASE N208462 002 Nov 20, 2015

+! EQ 4MG BASE N208462 003 Nov 20, 2015

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETALARAP +! PAR STERILE EQ 10MG BASE/ML N016812 001

PRODUCTS

AP +! EQ 50MG BASE/ML N016812 002AP +! EQ 100MG BASE/ML N016812 003KETAMINE HYDROCHLORIDEAP EUGIA PHARMA EQ 10MG BASE/ML A076092 001 Sep 30, 2008AP EQ 50MG BASE/ML A076092 002 Dec 28, 2001AP EQ 100MG BASE/ML A076092 003 Oct 25, 2002AP GLAND PHARMA LTD EQ 10MG BASE/ML A216809 001 Jan 24, 2023AP EQ 50MG BASE/ML A216809 002 Jan 24, 2023AP EQ 100MG BASE/ML A216809 003 Jan 24, 2023AP HIKMA EQ 50MG BASE/ML A074524 001 Mar 22, 1996AP EQ 100MG BASE/ML A074524 002 Mar 22, 1996AP HOSPIRA EQ 50MG BASE/ML A074549 001 Jun 27, 1996AP EQ 100MG BASE/ML A074549 002 Jun 27, 1996KETOCONAZOLE

AEROSOL, FOAM; TOPICAL

EXTINAAB +! MYLAN 2% N021738 001 Jun 12, 2007KETOCONAZOLEAB PADAGIS ISRAEL 2% A091550 001 Aug 25, 2011AB XIROMED 2% A213601 001 May 21, 2021

CREAM; TOPICAL

KETOCONAZOLEAB ENCUBE 2% A212443 001 May 20, 2021AB FOUGERA PHARMS 2% A076294 001 Apr 28, 2004AB TASMAN PHARMA 2% A215185 001 Nov 17, 2021AB ! TEVA 2% A075581 001 Apr 25, 2000KETOZOLEAB TARO 2% A075638 001 Dec 18, 2002

SHAMPOO; TOPICAL

KETOCONAZOLEAB ! PADAGIS ISRAEL 2% A076419 001 Jan 07, 2004AB TOLMAR 2% A076942 001 Apr 11, 2005

TABLET; ORAL

KETOCONAZOLEAB RISE PHARMA 200MG A075912 001 Jan 10, 2002AB STRIDES PHARMA 200MG A210457 001 Jun 18, 2018AB ! TARO 200MG A075319 001 Jun 15, 1999KETOPROFEN

CAPSULE; ORAL

KETOPROFENAB MISEMER 50MG A074014 002 Jan 29, 1993AB TEVA 50MG A073516 001 Dec 22, 1992

MISEMER 25MG A074014 001 Jan 29, 1993

! 75MG A074014 003 Jan 29, 1993

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

! MYLAN 200MG A075679 001 Feb 20, 2002

PRESCRIPTION DRUG PRODUCT LIST

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

<u>AP</u>	ALEMBIC	<u>15MG/ML</u>	<u>A214456 001</u>	Nov 02, 2022
<u>AP</u>		<u>30MG/ML</u>	<u>A214456 002</u>	Nov 02, 2022
<u>AP</u>	ASPIRO	<u>15MG/ML</u>	<u>A217166 001</u>	Jun 20, 2023
<u>AP</u>		<u>30MG/ML</u>	<u>A217166 002</u>	Jun 20, 2023
<u>AP</u>	BAXTER HLTHCARE CORP	<u>15MG/ML</u>	<u>A209900 002</u>	Jul 25, 2018
<u>AP</u>		<u>30MG/ML</u>	<u>A209900 001</u>	Sep 15, 2017
<u>AP</u>	CAPLIN	<u>15MG/ML</u>	<u>A217789 001</u>	May 10, 2023
<u>AP</u>		<u>30MG/ML</u>	<u>A217789 002</u>	May 10, 2023
<u>AP</u>	FRESENIUS KABI USA	<u>15MG/ML</u>	<u>A075784 001</u>	Jan 11, 2002
<u>AP</u>		<u>15MG/ML</u>	<u>A203242 001</u>	Oct 07, 2015
<u>AP</u>		<u>30MG/ML</u>	<u>A075784 002</u>	Jan 11, 2002
<u>AP</u>		<u>30MG/ML</u>	<u>A203242 002</u>	Oct 07, 2015
<u>AP</u>	GLAND	<u>15MG/ML</u>	<u>A204216 001</u>	Nov 01, 2016
<u>AP</u>		<u>30MG/ML</u>	<u>A204216 002</u>	Nov 01, 2016
<u>AP</u>	HIKMA	<u>15MG/ML</u>	<u>A075772 001</u>	Jul 21, 2004
<u>AP</u>		<u>30MG/ML</u>	<u>A075772 002</u>	Jul 21, 2004
<u>AP</u>	! HOSPIRA	<u>15MG/ML</u>	<u>A074802 001</u>	Jun 05, 1997
<u>AP</u>	!	<u>30MG/ML</u>	<u>A074802 002</u>	Jun 05, 1997
<u>AP</u>		<u>30MG/ML</u>	<u>A074993 002</u>	Jan 27, 1999
<u>AP</u>	NEPHRON	<u>30MG/ML</u>	<u>A211445 001</u>	Aug 20, 2020
<u>AP</u>	SAGENT PHARMS INC	<u>15MG/ML</u>	<u>A091065 001</u>	Nov 27, 2013
<u>AP</u>		<u>30MG/ML</u>	<u>A091065 002</u>	Nov 27, 2013
<u>AP</u>	SANDOZ	<u>30MG/ML</u>	<u>A076271 002</u>	Oct 06, 2004
<u>AP</u>	SUN PHARM	<u>15MG/ML</u>	<u>A078737 001</u>	Oct 06, 2008
<u>AP</u>		<u>30MG/ML</u>	<u>A078737 002</u>	Oct 06, 2008
<u>AP</u>	WOCKHARDT	<u>15MG/ML</u>	<u>A077942 001</u>	Mar 27, 2007
<u>AP</u>		<u>30MG/ML</u>	<u>A077942 002</u>	Mar 27, 2007

SOLUTION/DROPS;OPHTHALMIC

ACULAR

<u>AT</u>	+! ABBVIE	<u>0.5%</u>	<u>N019700 001</u>	Nov 09, 1992
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ACULAR LS

<u>AT</u>	+! ABBVIE	<u>0.4%</u>	<u>N021528 001</u>	May 30, 2003
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KETOROLAC TROMETHAMINE

<u>AT</u>	APOTEX INC	<u>0.4%</u>	<u>A077308 001</u>	Nov 05, 2009
<u>AT</u>		<u>0.5%</u>	<u>A076109 001</u>	Nov 05, 2009
<u>AT</u>	MICRO LABS LTD INDIA	<u>0.5%</u>	<u>A203410 001</u>	Apr 05, 2019
<u>AT</u>	SANDOZ	<u>0.5%</u>	<u>A076583 001</u>	Nov 05, 2009
<u>AT</u>	SUN PHARM	<u>0.5%</u>	<u>A090017 001</u>	Nov 05, 2009

ACUVAIL

+!	ABBVIE	0.45%	N022427 001	Jul 22, 2009
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SPRAY, METERED;NASAL

SPRIX

+!	ZYLA	15.75MG/SPRAY	N022382 001	May 14, 2010
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TABLET;ORAL

KETOROLAC TROMETHAMINE

<u>AB</u>	ATNAHS PHARMA US	<u>10MG</u>	<u>A216407 001</u>	Oct 21, 2022
<u>AB</u>	BIONPHARMA	<u>10MG</u>	<u>A216759 001</u>	Feb 23, 2023
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A210616 001</u>	Aug 16, 2018
<u>AB</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A216651 001</u>	Aug 25, 2022
<u>AB</u>	LEADING	<u>10MG</u>	<u>A215745 001</u>	Feb 23, 2023
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A074761 001</u>	May 16, 1997
<u>AB</u>	SENORES PHARMS	<u>10MG</u>	<u>A215788 001</u>	Mar 23, 2022
<u>AB</u>	! TEVA	<u>10MG</u>	<u>A074754 001</u>	May 16, 1997
<u>AB</u>	ZYDUS	<u>10MG</u>	<u>A217038 001</u>	Oct 21, 2022

KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION;IRRIGATION

KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE

<u>AT</u>	LUPIN LTD	<u>EQ 0.3% BASE;EQ 1% BASE</u>	<u>A210183 001</u>	Jul 01, 2019
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OMIDRIA

<u>AT</u>	+! RAYNER SURGICAL	<u>EQ 0.3% BASE;EQ 1% BASE</u>	<u>N205388 001</u>	May 30, 2014
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L-GLUTAMINE

FOR SOLUTION;ORAL

ENDARI

+	EMMAUS MEDCL	5GM/PACKET	N208587 001	Jul 07, 2017
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PRESCRIPTION DRUG PRODUCT LIST

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

<u>AP</u>	BAXTER HLTHCARE CORP	<u>5MG/ML</u>	<u>A076051 001</u>	Jul 05, 2002
<u>AP</u>	CAPLIN	<u>5MG/ML</u>	<u>A214533 001</u>	Sep 07, 2021
<u>AP</u>	GLAND PHARMA LTD	<u>5MG/ML</u>	<u>A090699 001</u>	Apr 03, 2012
<u>AP</u>	HIKMA	<u>5MG/ML</u>	<u>A075303 001</u>	May 28, 1999
<u>AP</u>	! HOSPIRA	<u>5MG/ML</u>	<u>A075239 001</u>	Nov 29, 1999
<u>AP</u>	! RISING	<u>5MG/ML</u>	<u>A075240 001</u>	Nov 29, 1999
<u>AP</u>	RISING	<u>5MG/ML</u>	<u>A075431 001</u>	Nov 29, 1999

SOLUTION; INTRAVENOUS

LABETALOL HYDROCHLORIDE

+!	HIKMA	10MG/2ML (5MG/ML)	N213330 005	Mar 18, 2022
	LABETALOL HYDROCHLORIDE IN DEXTROSE			
+!	HIKMA	200MG/200ML (1MG/ML)	N213330 001	Nov 09, 2020
	LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE			
+!	HIKMA	100MG/100ML (1MG/ML)	N213330 002	Nov 09, 2020
+!		200MG/200ML (1MG/ML)	N213330 003	Nov 09, 2020
+!		300MG/300ML (1MG/ML)	N213330 004	Nov 09, 2020

TABLET; ORAL

LABETALOL HYDROCHLORIDE

<u>AB</u>	APPCO	<u>100MG</u>	<u>A209603 001</u>	Jun 20, 2018
<u>AB</u>		<u>200MG</u>	<u>A209603 002</u>	Jun 20, 2018
<u>AB</u>		<u>300MG</u>	<u>A209603 003</u>	Jun 20, 2018
<u>AB</u>	CADILA PHARMS LTD	<u>100MG</u>	<u>A211325 001</u>	May 13, 2019
<u>AB</u>		<u>200MG</u>	<u>A211325 002</u>	May 13, 2019
<u>AB</u>		<u>300MG</u>	<u>A211325 003</u>	May 13, 2019
<u>AB</u>	EPIC PHARMA LLC	<u>100MG</u>	<u>A212990 001</u>	Sep 30, 2020
<u>AB</u>		<u>200MG</u>	<u>A212990 002</u>	Sep 30, 2020
<u>AB</u>		<u>300MG</u>	<u>A212990 003</u>	Sep 30, 2020
<u>AB</u>	EYWA	<u>100MG</u>	<u>A207863 001</u>	Feb 04, 2019
<u>AB</u>		<u>200MG</u>	<u>A207863 002</u>	Feb 04, 2019
<u>AB</u>		<u>300MG</u>	<u>A207863 003</u>	Feb 04, 2019
<u>AB</u>	HERITAGE PHARMA	<u>100MG</u>	<u>A074787 001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>A074787 002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>A074787 003</u>	Aug 03, 1998
<u>AB</u>	INNOGENIX	<u>100MG</u>	<u>A075215 001</u>	Jul 29, 1999
<u>AB</u>		<u>200MG</u>	<u>A075215 002</u>	Jul 29, 1999
<u>AB</u>		<u>300MG</u>	<u>A075215 003</u>	Jul 29, 1999
<u>AB</u>	PAR FORM	<u>100MG</u>	<u>A200908 001</u>	Jul 10, 2012
<u>AB</u>	!	<u>200MG</u>	<u>A200908 002</u>	Jul 10, 2012
<u>AB</u>		<u>300MG</u>	<u>A200908 003</u>	Jul 10, 2012
<u>AB</u>	RUBICON	<u>100MG</u>	<u>A211953 001</u>	Aug 18, 2021
<u>AB</u>		<u>200MG</u>	<u>A211953 002</u>	Aug 18, 2021
<u>AB</u>		<u>300MG</u>	<u>A211953 003</u>	Aug 18, 2021
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A075113 001</u>	Aug 04, 1998
<u>AB</u>	!	<u>200MG</u>	<u>A075113 002</u>	Aug 04, 1998
<u>AB</u>		<u>300MG</u>	<u>A075113 003</u>	Aug 04, 1998
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>A075133 001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>A075133 002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>A075133 003</u>	Aug 03, 1998
<u>AB</u>	ZYDUS PHARMS	<u>100MG</u>	<u>A207743 001</u>	Sep 19, 2017
<u>AB</u>		<u>200MG</u>	<u>A207743 002</u>	Sep 19, 2017
<u>AB</u>		<u>300MG</u>	<u>A207743 003</u>	Sep 19, 2017

LACOSAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

MOTPOLY XR

+	AUCTA	100MG	N216185 001	May 04, 2023
+		150MG	N216185 002	May 04, 2023
+!		200MG	N216185 003	May 04, 2023

SOLUTION; INTRAVENOUS

LACOSAMIDE

<u>AP</u>	APOTEX	<u>200MG/20ML (10MG/ML)</u>	<u>A216670 001</u>	Nov 13, 2023
<u>AP</u>	ASPIRO	<u>200MG/20ML (10MG/ML)</u>	<u>A216335 001</u>	Sep 14, 2022
<u>AP</u>	FRESENIUS KABI USA	<u>200MG/20ML (10MG/ML)</u>	<u>A214677 001</u>	May 03, 2022
<u>AP</u>	GLAND PHARMA LTD	<u>200MG/20ML (10MG/ML)</u>	<u>A215628 001</u>	Sep 19, 2022
<u>AP</u>	INDOCO	<u>200MG/20ML (10MG/ML)</u>	<u>A214301 001</u>	Apr 07, 2022
<u>AP</u>	MSN	<u>200MG/20ML (10MG/ML)</u>	<u>A215979 001</u>	Feb 09, 2023
<u>AP</u>	ZYDUS PHARMS	<u>200MG/20ML (10MG/ML)</u>	<u>A209465 001</u>	Jun 29, 2022
	<u>VIMPAT</u>			
<u>AP</u>	+! UCB INC	<u>200MG/20ML (10MG/ML)</u>	<u>N022254 001</u>	Oct 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

LACOSAMIDE

SOLUTION;ORAL

LACOSAMIDE

<u>AA</u>	ALKEM LABS LTD	<u>10MG/ML</u>	<u>A214672</u>	<u>001</u>	May 19, 2022
<u>AA</u>	APOTEX	<u>10MG/ML</u>	<u>A206355</u>	<u>001</u>	Sep 26, 2022
<u>AA</u>	HETERO LABS LTD III	<u>10MG/ML</u>	<u>A209301</u>	<u>001</u>	May 31, 2022
<u>AA</u>	LEADING PHARMA	<u>10MG/ML</u>	<u>A216461</u>	<u>001</u>	Feb 06, 2023
<u>AA</u>	MSN	<u>10MG/ML</u>	<u>A215379</u>	<u>001</u>	Dec 22, 2023
<u>AA</u>	NOVITIUM PHARMA	<u>10MG/ML</u>	<u>A216151</u>	<u>001</u>	Aug 26, 2022

VIMPAT

<u>AA</u>	+! UCB INC	<u>10MG/ML</u>	<u>N022255</u>	<u>001</u>	Apr 20, 2010
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TABLET;ORAL

LACOSAMIDE

<u>AB</u>	ALEMBIC	<u>50MG</u>	<u>A204974</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A204974</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A204974</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A204974</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	ALKEM LABS LTD	<u>50MG</u>	<u>A214695</u>	<u>001</u>	Mar 31, 2022
<u>AB</u>		<u>100MG</u>	<u>A214695</u>	<u>002</u>	Mar 31, 2022
<u>AB</u>		<u>150MG</u>	<u>A214695</u>	<u>003</u>	Mar 31, 2022
<u>AB</u>		<u>200MG</u>	<u>A214695</u>	<u>004</u>	Mar 31, 2022
<u>AB</u>	AMNEAL PHARMS	<u>50MG</u>	<u>A204857</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A204857</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A204857</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A204857</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A204994</u>	<u>001</u>	Jan 05, 2023
<u>AB</u>		<u>100MG</u>	<u>A204994</u>	<u>002</u>	Jan 05, 2023
<u>AB</u>		<u>150MG</u>	<u>A204994</u>	<u>003</u>	Jan 05, 2023
<u>AB</u>		<u>200MG</u>	<u>A204994</u>	<u>004</u>	Jan 05, 2023
<u>AB</u>	GLENMARK PHARMS LTD	<u>50MG</u>	<u>A205006</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A205006</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A205006</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A205006</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	HETERO LABS LTD V	<u>50MG</u>	<u>A204787</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A204787</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A204787</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A204787</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	INDOCO	<u>50MG</u>	<u>A208308</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A208308</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A208308</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A208308</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	MSN LABS PVT LTD	<u>50MG</u>	<u>A204921</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A204921</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A204921</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A204921</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	SCIEGEN PHARMS INC	<u>50MG</u>	<u>A205237</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A205237</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A205237</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A205237</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	SUN PHARM	<u>50MG</u>	<u>A205031</u>	<u>001</u>	Mar 17, 2022
<u>AB</u>		<u>100MG</u>	<u>A205031</u>	<u>002</u>	Mar 17, 2022
<u>AB</u>		<u>150MG</u>	<u>A205031</u>	<u>003</u>	Mar 17, 2022
<u>AB</u>		<u>200MG</u>	<u>A205031</u>	<u>004</u>	Mar 17, 2022
<u>AB</u>	ZYDUS PHARMS	<u>50MG</u>	<u>A204947</u>	<u>001</u>	Dec 15, 2023
<u>AB</u>		<u>100MG</u>	<u>A204947</u>	<u>002</u>	Dec 15, 2023
<u>AB</u>		<u>150MG</u>	<u>A204947</u>	<u>003</u>	Dec 15, 2023
<u>AB</u>		<u>200MG</u>	<u>A204947</u>	<u>004</u>	Dec 15, 2023
	<u>VIMPAT</u>				
<u>AB</u>	+ UCB INC	<u>50MG</u>	<u>N022253</u>	<u>001</u>	Oct 28, 2008
<u>AB</u>	+	<u>100MG</u>	<u>N022253</u>	<u>002</u>	Oct 28, 2008
<u>AB</u>	+	<u>150MG</u>	<u>N022253</u>	<u>003</u>	Oct 28, 2008
<u>AB</u>	+!	<u>200MG</u>	<u>N022253</u>	<u>004</u>	Oct 28, 2008

LACTULOSE

FOR SOLUTION;ORAL

LACTULOSE

!	CUMBERLAND PHARMS	10GM/PACKET	A074712	001	Dec 10, 1997
!		20GM/PACKET	A074712	002	Dec 10, 1997

SOLUTION;ORAL

LACTULOSE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>10GM/15ML</u>	<u>A074602</u>	<u>001</u>	Nov 14, 1996
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PRESCRIPTION DRUG PRODUCT LIST

LACTULOSE

SOLUTION;ORAL

LACTULOSE

AA	CHARTWELL RX	<u>10GM/15ML</u>	<u>A209517</u>	<u>001</u>	Nov 23, 2018
AA	FRESENIUS KABI	<u>10GM/15ML</u>	<u>A090503</u>	<u>001</u>	Jan 25, 2012
AA	! HIKMA	<u>10GM/15ML</u>	<u>A074076</u>	<u>001</u>	Jul 03, 1995
AA	LANNETT CO INC	<u>10GM/15ML</u>	<u>A075993</u>	<u>001</u>	Jul 26, 2001
AA	PHARM ASSOC	<u>10GM/15ML</u>	<u>A074623</u>	<u>001</u>	Jul 30, 1996
AA	XTTRIUM LABS INC	<u>10GM/15ML</u>	<u>A075911</u>	<u>001</u>	Feb 21, 2002

SOLUTION;ORAL, RECTAL

GENERLAC

AA	WOCKHARDT BIO AG	<u>10GM/15ML</u>	<u>A074603</u>	<u>001</u>	Oct 31, 1996
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LACTULOSE

AA	BAJAJ	<u>10GM/15ML</u>	<u>A076645</u>	<u>001</u>	Jul 28, 2003
AA	! FRESENIUS KABI	<u>10GM/15ML</u>	<u>A090502</u>	<u>001</u>	Jan 25, 2012
AA	PAI HOLDINGS PHARM	<u>10GM/15ML</u>	<u>A074077</u>	<u>001</u>	Jul 03, 1995

LAMIVUDINE

SOLUTION;ORAL

EPIVIR

AA	+! VIIV HLTHCARE	<u>10MG/ML</u>	<u>N020596</u>	<u>001</u>	Nov 17, 1995
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LAMIVUDINE

AA	AUROBINDO PHARMA	<u>10MG/ML</u>	<u>A077695</u>	<u>001</u>	Nov 21, 2016
AA	CHARTWELL MOLECULAR	<u>10MG/ML</u>	<u>A203564</u>	<u>001</u>	Oct 31, 2014
AA	HETERO LABS LTD III	<u>10MG/ML</u>	<u>A091475</u>	<u>001</u>	Oct 06, 2023

EPIVIR-HBV

+!	GLAXOSMITHKLINE	5MG/ML	N021004	001	Dec 08, 1998
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TABLET;ORAL

EPIVIR

AB	+ VIIV HLTHCARE	<u>150MG</u>	<u>N020564</u>	<u>001</u>	Nov 17, 1995
AB	+!	<u>300MG</u>	<u>N020564</u>	<u>003</u>	Jun 24, 2002

EPIVIR-HBV

AB	+! GLAXOSMITHKLINE	<u>100MG</u>	<u>N021003</u>	<u>001</u>	Dec 08, 1998
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LAMIVUDINE

AB	ANNORA	<u>100MG</u>	<u>A211306</u>	<u>001</u>	Mar 21, 2019
AB	APOTEX	<u>100MG</u>	<u>A202941</u>	<u>001</u>	Jan 02, 2014
AB		<u>150MG</u>	<u>A091606</u>	<u>001</u>	Dec 02, 2011
AB		<u>300MG</u>	<u>A091606</u>	<u>002</u>	Dec 02, 2011
AB	AUROBINDO PHARMA	<u>150MG</u>	<u>A077464</u>	<u>001</u>	Nov 21, 2016
AB		<u>300MG</u>	<u>A077464</u>	<u>002</u>	Nov 21, 2016
AB	BRECKENRIDGE	<u>150MG</u>	<u>A203586</u>	<u>001</u>	Nov 21, 2016
AB	CIPLA	<u>150MG</u>	<u>A077221</u>	<u>001</u>	Mar 03, 2017
AB		<u>300MG</u>	<u>A077221</u>	<u>002</u>	Mar 03, 2017
AB	HETERO LABS LTD V	<u>100MG</u>	<u>A203260</u>	<u>001</u>	Jan 02, 2014
AB		<u>150MG</u>	<u>A203277</u>	<u>001</u>	Jan 06, 2014
AB		<u>300MG</u>	<u>A203277</u>	<u>002</u>	Jan 06, 2014
AB	LUPIN LTD	<u>150MG</u>	<u>A205217</u>	<u>001</u>	Dec 18, 2014
AB		<u>300MG</u>	<u>A205217</u>	<u>002</u>	Dec 18, 2014
AB	MACLEODS PHARMS LTD	<u>150MG</u>	<u>A090198</u>	<u>001</u>	May 01, 2019
AB		<u>300MG</u>	<u>A090198</u>	<u>002</u>	May 01, 2019
AB	MYLAN LABS LTD	<u>150MG</u>	<u>A078545</u>	<u>001</u>	Mar 05, 2019
AB		<u>300MG</u>	<u>A078545</u>	<u>002</u>	Mar 05, 2019
AB	STRIDES PHARMA	<u>150MG</u>	<u>A090457</u>	<u>001</u>	Apr 19, 2018
AB		<u>300MG</u>	<u>A090457</u>	<u>002</u>	Apr 19, 2018
AB	UPSHER SMITH LABS	<u>150MG</u>	<u>A206974</u>	<u>001</u>	Nov 21, 2016
AB		<u>300MG</u>	<u>A206974</u>	<u>002</u>	Nov 21, 2016

LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

CIMDUO

+!	MYLAN LABS LTD	300MG;300MG	N022141	001	Feb 28, 2018
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LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

COMBIVIR

AB	+! VIIV HLTHCARE	<u>150MG;300MG</u>	<u>N020857</u>	<u>001</u>	Sep 26, 1997
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LAMIVUDINE AND ZIDOVUDINE

AB	ANDA REPOSITORY	<u>150MG;300MG</u>	<u>A206375</u>	<u>001</u>	Apr 10, 2018
AB	AUROBINDO PHARMA	<u>150MG;300MG</u>	<u>A077558</u>	<u>001</u>	May 05, 2017
AB	CIPLA	<u>150MG;300MG</u>	<u>A077411</u>	<u>001</u>	Sep 07, 2018
AB	HETERO LABS LTD III	<u>150MG;300MG</u>	<u>A079124</u>	<u>001</u>	Sep 17, 2015
AB	HETERO LABS LTD V	<u>150MG;300MG</u>	<u>A203259</u>	<u>001</u>	Feb 03, 2014
AB	LUPIN LTD	<u>150MG;300MG</u>	<u>A090246</u>	<u>001</u>	May 15, 2012
AB	MACLEODS PHARMS LTD	<u>150MG;300MG</u>	<u>A090679</u>	<u>001</u>	Aug 29, 2018

PRESCRIPTION DRUG PRODUCT LISTLAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

LAMIVUDINE AND ZIDOVUDINE

AB	STRIDES PHARMA	150MG;300MG	A079128 001	May 13, 2015
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LAMOTRIGINE

TABLET;ORAL

LAMICTAL

AB	+	GLAXOSMITHKLINE LLC	25MG	N020241 005	Dec 27, 1994
AB	+		100MG	N020241 001	Dec 27, 1994
AB	+		150MG	N020241 002	Dec 27, 1994
AB	+		200MG	N020241 003	Dec 27, 1994

LAMOTRIGINE

AB	ALEMBIC PHARMS LTD	25MG	A090607 001	Jan 13, 2011
AB		100MG	A090607 002	Jan 13, 2011
AB		150MG	A090607 003	Jan 13, 2011
AB		200MG	A090607 004	Jan 13, 2011
AB	ALKEM LABS LTD	25MG	A200694 001	Jun 14, 2013
AB		100MG	A200694 002	Jun 14, 2013
AB		150MG	A200694 003	Jun 14, 2013
AB		200MG	A200694 004	Jun 14, 2013
AB	AUROBINDO PHARMA	25MG	A078956 001	Jan 27, 2009
AB		100MG	A078956 002	Jan 27, 2009
AB		150MG	A078956 003	Jan 27, 2009
AB		200MG	A078956 004	Jan 27, 2009
AB	DR REDDYS LABS LTD	25MG	A076708 001	Jan 27, 2009
AB		100MG	A076708 002	Jan 27, 2009
AB		150MG	A076708 003	Jan 27, 2009
AB		200MG	A076708 004	Jan 27, 2009
AB	GLENMARK GENERICS	25MG	A090169 001	May 12, 2012
AB		100MG	A090169 002	May 12, 2012
AB		150MG	A090169 003	May 12, 2012
AB		200MG	A090169 004	May 12, 2012
AB	JUBILANT CADISTA	25MG	A079132 001	Jan 27, 2009
AB		100MG	A079132 002	Jan 27, 2009
AB		150MG	A079132 003	Jan 27, 2009
AB		200MG	A079132 004	Jan 27, 2009
AB	LUPIN LTD	25MG	A078691 001	Jun 01, 2010
AB		100MG	A078691 002	Jun 01, 2010
AB		150MG	A078691 003	Jun 01, 2010
AB		200MG	A078691 004	Jun 01, 2010
AB	RUBICON	25MG	A078625 001	Jan 27, 2009
AB		100MG	A078625 002	Jan 27, 2009
AB		150MG	A078625 003	Jan 27, 2009
AB		200MG	A078625 004	Jan 27, 2009
AB	TARO PHARM INDS	25MG	A078525 001	Jan 27, 2009
AB		100MG	A078525 002	Jan 27, 2009
AB		150MG	A078525 003	Jan 27, 2009
AB		200MG	A078525 004	Jan 27, 2009
AB	TORRENT PHARMS	25MG	A078947 001	Jan 27, 2009
AB		100MG	A078947 002	Jan 27, 2009
AB		150MG	A078947 003	Jan 27, 2009
AB		200MG	A078947 004	Jan 27, 2009
AB	UNICHEM LABS LTD	25MG	A090170 001	Oct 06, 2011
AB		100MG	A090170 002	Oct 06, 2011
AB		150MG	A090170 003	Oct 06, 2011
AB		200MG	A090170 004	Oct 06, 2011
AB	ZYDUS PHARMS USA	25MG	A077633 001	Jan 27, 2009
AB		100MG	A077633 003	Jan 27, 2009
AB		150MG	A077633 004	Jan 27, 2009
AB		200MG	A077633 005	Jan 27, 2009
		50MG	A077633 002	Jan 27, 2009
		250MG	A077633 006	Jan 27, 2009

TABLET, EXTENDED RELEASE;ORAL

LAMICTAL XR

AB	+	GLAXOSMITHKLINE LLC	25MG	N022115 001	May 29, 2009
AB	+		50MG	N022115 002	May 29, 2009
AB	+		100MG	N022115 003	May 29, 2009
AB	+		200MG	N022115 004	May 29, 2009
AB	+		250MG	N022115 006	Jun 21, 2011
AB	+		300MG	N022115 005	Apr 14, 2010

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A200672 003</u>	Oct 17, 2013
<u>AB</u>		<u>200MG</u>	<u>A200672 004</u>	Oct 17, 2013
<u>AB</u>		<u>25MG</u>	<u>A200672 001</u>	Oct 17, 2013
<u>AB</u>		<u>50MG</u>	<u>A200672 002</u>	Oct 17, 2013
<u>AB</u>		<u>250MG</u>	<u>A200672 006</u>	Nov 13, 2013
<u>AB</u>		<u>300MG</u>	<u>A200672 005</u>	Oct 17, 2013
<u>AB</u>	AMNEAL PHARMS	<u>25MG</u>	<u>A207497 001</u>	Nov 30, 2018
<u>AB</u>		<u>50MG</u>	<u>A207497 002</u>	Nov 30, 2018
<u>AB</u>		<u>100MG</u>	<u>A207497 003</u>	Nov 30, 2018
<u>AB</u>		<u>200MG</u>	<u>A207497 004</u>	Nov 30, 2018
<u>AB</u>		<u>250MG</u>	<u>A207497 005</u>	Nov 30, 2018
<u>AB</u>		<u>300MG</u>	<u>A207497 006</u>	Nov 30, 2018
<u>AB</u>	ANCHEN PHARMS	<u>25MG</u>	<u>A201374 001</u>	Dec 26, 2012
<u>AB</u>		<u>50MG</u>	<u>A201374 002</u>	Dec 26, 2012
<u>AB</u>		<u>100MG</u>	<u>A201374 003</u>	Dec 26, 2012
<u>AB</u>		<u>200MG</u>	<u>A201374 004</u>	Dec 26, 2012
<u>AB</u>		<u>250MG</u>	<u>A201374 005</u>	Dec 26, 2012
<u>AB</u>		<u>300MG</u>	<u>A201374 006</u>	Dec 26, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>25MG</u>	<u>A202383 001</u>	Jun 19, 2013
<u>AB</u>		<u>50MG</u>	<u>A202383 002</u>	Jun 19, 2013
<u>AB</u>		<u>100MG</u>	<u>A202383 003</u>	Jun 19, 2013
<u>AB</u>		<u>200MG</u>	<u>A202383 004</u>	Jun 19, 2013
<u>AB</u>		<u>250MG</u>	<u>A202383 006</u>	Sep 06, 2018
<u>AB</u>		<u>300MG</u>	<u>A202383 005</u>	Jun 19, 2013
<u>AB</u>	PAR PHARM	<u>25MG</u>	<u>A201791 001</u>	Jan 18, 2013
<u>AB</u>		<u>50MG</u>	<u>A201791 002</u>	Jan 18, 2013
<u>AB</u>		<u>100MG</u>	<u>A201791 003</u>	Jan 18, 2013
<u>AB</u>		<u>200MG</u>	<u>A201791 004</u>	Jan 18, 2013
<u>AB</u>		<u>250MG</u>	<u>A201791 005</u>	Jan 18, 2013
<u>AB</u>		<u>300MG</u>	<u>A201791 006</u>	Jan 18, 2013
<u>AB</u>	RUBICON	<u>100MG</u>	<u>A202887 003</u>	Jun 17, 2013
<u>AB</u>		<u>200MG</u>	<u>A202887 004</u>	Jun 17, 2013
<u>AB</u>	TORRENT	<u>25MG</u>	<u>A203370 001</u>	Dec 23, 2013
<u>AB</u>		<u>50MG</u>	<u>A203370 002</u>	Dec 23, 2013
<u>AB</u>		<u>100MG</u>	<u>A203370 003</u>	Dec 23, 2013
<u>AB</u>	WOCKHARDT BIO AG	<u>25MG</u>	<u>A202498 001</u>	Jan 04, 2013
<u>AB</u>		<u>50MG</u>	<u>A202498 002</u>	Jan 04, 2013
<u>AB</u>		<u>100MG</u>	<u>A202498 003</u>	Jan 04, 2013
<u>AB</u>		<u>200MG</u>	<u>A202498 004</u>	Jan 04, 2013
<u>AB</u>		<u>300MG</u>	<u>A202498 005</u>	Jan 04, 2013
<u>AB</u>	YILING	<u>25MG</u>	<u>A213949 001</u>	Dec 08, 2021
<u>AB</u>		<u>50MG</u>	<u>A213949 002</u>	Dec 08, 2021
<u>AB</u>		<u>100MG</u>	<u>A213949 003</u>	Dec 08, 2021
<u>AB</u>		<u>200MG</u>	<u>A213949 004</u>	Dec 08, 2021
<u>AB</u>		<u>250MG</u>	<u>A213949 005</u>	Dec 08, 2021
<u>AB</u>		<u>300MG</u>	<u>A213949 006</u>	Dec 08, 2021
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A207763 001</u>	Apr 01, 2020
<u>AB</u>		<u>50MG</u>	<u>A207763 002</u>	Apr 01, 2020
<u>AB</u>		<u>100MG</u>	<u>A207763 003</u>	Apr 01, 2020
<u>AB</u>		<u>200MG</u>	<u>A207763 004</u>	Apr 01, 2020
<u>AB</u>		<u>250MG</u>	<u>A207763 005</u>	Apr 01, 2020
<u>AB</u>		<u>300MG</u>	<u>A207763 006</u>	Apr 01, 2020

TABLET, FOR SUSPENSION;ORAL

LAMICTAL CD

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>2MG</u>	<u>N020764 004</u>	Sep 08, 2000
<u>AB</u>	+		<u>5MG</u>	<u>N020764 001</u>	Aug 24, 1998
<u>AB</u>	+		<u>25MG</u>	<u>N020764 002</u>	Aug 24, 1998

LAMOTRIGINE

<u>AB</u>	ALEMBIC	<u>5MG</u>	<u>A201168 001</u>	Jun 12, 2014
<u>AB</u>		<u>25MG</u>	<u>A201168 002</u>	Jun 12, 2014
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A090401 002</u>	Nov 04, 2009
<u>AB</u>		<u>25MG</u>	<u>A090401 003</u>	Nov 04, 2009
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076701 001</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076701 002</u>	Jan 22, 2009
<u>AB</u>	GLENMARK PHARMS LTD	<u>5MG</u>	<u>A079099 001</u>	Feb 19, 2009
<u>AB</u>		<u>25MG</u>	<u>A079099 002</u>	Feb 19, 2009
<u>AB</u>	TARO	<u>5MG</u>	<u>A079204 001</u>	Feb 04, 2009
<u>AB</u>		<u>25MG</u>	<u>A079204 002</u>	Feb 04, 2009
<u>AB</u>	WATSON LABS	<u>2MG</u>	<u>A076928 001</u>	Jan 22, 2009

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, FOR SUSPENSION;ORAL

LAMOTRIGINE

<u>AB</u>		<u>5MG</u>	<u>A076928 002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076928 003</u>	Jan 22, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A078009 002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A078009 003</u>	Jan 22, 2009

TABLET, ORALLY DISINTEGRATING;ORAL

LAMICTAL ODT

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N022251 001</u>	May 08, 2009
<u>AB</u>	+	!	<u>50MG</u>	<u>N022251 002</u>	May 08, 2009
<u>AB</u>	+		<u>100MG</u>	<u>N022251 003</u>	May 08, 2009
<u>AB</u>	+		<u>200MG</u>	<u>N022251 004</u>	May 08, 2009

LAMOTRIGINE

<u>AB</u>		AJANTA PHARMA LTD	<u>25MG</u>	<u>A213271 001</u>	Jan 19, 2021
<u>AB</u>			<u>50MG</u>	<u>A213271 002</u>	Jan 19, 2021
<u>AB</u>			<u>100MG</u>	<u>A213271 003</u>	Jan 19, 2021
<u>AB</u>			<u>200MG</u>	<u>A213271 004</u>	Jan 19, 2021
<u>AB</u>		AMRING PHARMS	<u>25MG</u>	<u>A214124 001</u>	Feb 03, 2022
<u>AB</u>			<u>50MG</u>	<u>A214124 002</u>	Feb 03, 2022
<u>AB</u>			<u>100MG</u>	<u>A214124 003</u>	Feb 03, 2022
<u>AB</u>			<u>200MG</u>	<u>A214124 004</u>	Feb 03, 2022
<u>AB</u>		PAR PHARM	<u>25MG</u>	<u>A204158 001</u>	Oct 27, 2015
<u>AB</u>			<u>50MG</u>	<u>A204158 002</u>	Oct 27, 2015
<u>AB</u>			<u>100MG</u>	<u>A204158 003</u>	Oct 27, 2015
<u>AB</u>			<u>200MG</u>	<u>A204158 004</u>	Oct 27, 2015
<u>AB</u>		SCIEGEN PHARMS INC	<u>25MG</u>	<u>A206382 001</u>	Jun 17, 2016
<u>AB</u>			<u>50MG</u>	<u>A206382 002</u>	Jun 17, 2016
<u>AB</u>			<u>100MG</u>	<u>A206382 003</u>	Jun 17, 2016
<u>AB</u>			<u>200MG</u>	<u>A206382 004</u>	Jun 17, 2016

LANREOTIDE ACETATE

SOLUTION;SUBCUTANEOUS

LANREOTIDE ACETATE

+	!	INVAGEN PHARMS	EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)	N215395 001	Dec 17, 2021
+	!		EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)	N215395 002	Dec 17, 2021
+	!		EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML)	N215395 003	Dec 17, 2021

SOMATULINE DEPOT

+	!	IPSEN PHARMA	EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)	N022074 001	Aug 30, 2007
+	!		EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)	N022074 002	Aug 30, 2007
+	!		EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML)	N022074 003	Aug 30, 2007

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

<u>AB</u>		ALKEM LABS LTD	<u>15MG</u>	<u>A207394 001</u>	Jan 18, 2019
<u>AB</u>			<u>30MG</u>	<u>A207394 002</u>	Jan 18, 2019
<u>AB</u>		CHARTWELL MOLECULAR	<u>15MG</u>	<u>A207156 001</u>	Sep 28, 2017
<u>AB</u>			<u>30MG</u>	<u>A207156 002</u>	Sep 28, 2017
<u>AB</u>		DR REDDYS LABS LTD	<u>15MG</u>	<u>A091269 001</u>	Oct 15, 2010
<u>AB</u>			<u>30MG</u>	<u>A091269 002</u>	Oct 15, 2010
<u>AB</u>		HETERO LABS LTD III	<u>15MG</u>	<u>A203083 001</u>	May 18, 2020
<u>AB</u>			<u>30MG</u>	<u>A203083 002</u>	May 18, 2020
<u>AB</u>		INVENTIA	<u>15MG</u>	<u>A205868 001</u>	Aug 30, 2017
<u>AB</u>			<u>30MG</u>	<u>A205868 002</u>	Aug 30, 2017
<u>AB</u>		MYLAN PHARMS INC	<u>15MG</u>	<u>A090763 001</u>	Nov 10, 2009
<u>AB</u>			<u>30MG</u>	<u>A090763 002</u>	Nov 10, 2009
<u>AB</u>		NATCO PHARMA LTD	<u>15MG</u>	<u>A201921 001</u>	Dec 18, 2012
<u>AB</u>			<u>30MG</u>	<u>A201921 002</u>	Dec 18, 2012
<u>AB</u>		SANDOZ	<u>15MG</u>	<u>A090331 001</u>	Apr 23, 2010
<u>AB</u>			<u>30MG</u>	<u>A090331 002</u>	Apr 23, 2010
<u>AB</u>		SUN PHARM	<u>15MG</u>	<u>A202637 001</u>	Sep 13, 2013
<u>AB</u>			<u>30MG</u>	<u>A091509 001</u>	Sep 13, 2013
<u>AB</u>		TEVA PHARMS	<u>15MG</u>	<u>A077255 001</u>	Nov 10, 2009
<u>AB</u>			<u>30MG</u>	<u>A077255 002</u>	Nov 10, 2009
<u>AB</u>		WOCKHARDT	<u>15MG</u>	<u>A202176 001</u>	Sep 14, 2012
<u>AB</u>			<u>30MG</u>	<u>A202176 002</u>	Sep 14, 2012
<u>AB</u>		XIROMED	<u>15MG</u>	<u>A203203 001</u>	Jul 25, 2016
<u>AB</u>			<u>30MG</u>	<u>A203203 002</u>	Jul 25, 2016
<u>AB</u>		ZYDUS HLTHCARE	<u>15MG</u>	<u>A202366 001</u>	Aug 19, 2013
<u>AB</u>			<u>30MG</u>	<u>A202366 002</u>	Aug 19, 2013

PRESCRIPTION DRUG PRODUCT LIST

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

PREVACID

AB	+	TAKEDA PHARMS USA	30MG	N020406	002	May 10, 1995
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TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

AB		AUROBINDO PHARMA LTD	15MG	A207167	001	Mar 28, 2023
AB			30MG	A207167	002	Mar 28, 2023
AB		DR REDDYS	15MG	A210465	001	Feb 01, 2021
AB			30MG	A210465	002	Feb 01, 2021
AB		MYLAN	15MG	A202396	001	Nov 28, 2018
AB			30MG	A202396	002	Nov 28, 2018
AB		TEVA PHARMS USA	15MG	A208784	001	Sep 21, 2017
AB			30MG	A208784	002	Sep 21, 2017
AB		ZYDUS PHARMS	15MG	A200816	001	Nov 27, 2018
AB			30MG	A200816	002	Nov 27, 2018

PREVACID

AB	+	TAKEDA PHARMS USA	15MG	N021428	001	Aug 30, 2002
AB	+		30MG	N021428	002	Aug 30, 2002

LANTHANUM CARBONATE

POWDER;ORAL

FOSRENOL

+	TAKEDA PHARMS USA	EQ 750MG BASE	N204734	001	Sep 24, 2014
+		EQ 1GM BASE	N204734	002	Sep 24, 2014

TABLET, CHEWABLE;ORAL

FOSRENOL

AB	+	TAKEDA PHARMS USA	EQ 500MG BASE	N021468	002	Oct 26, 2004
AB	+		EQ 750MG BASE	N021468	003	Nov 23, 2005
AB	+		EQ 1GM BASE	N021468	004	Nov 23, 2005

LANTHANUM CARBONATE

AB		BARR	EQ 500MG BASE	A090977	001	Jan 27, 2022
AB			EQ 750MG BASE	A090977	002	Jan 27, 2022
AB			EQ 1GM BASE	A090977	003	Jan 27, 2022
AB		INVAGEN PHARMS	EQ 500MG BASE	A206868	001	Jan 24, 2022
AB			EQ 750MG BASE	A206868	002	Jan 24, 2022
AB			EQ 1GM BASE	A206868	003	Jan 24, 2022
AB		NATCO PHARMA LTD	EQ 500MG BASE	A090978	001	Aug 11, 2017
AB			EQ 750MG BASE	A090978	002	Aug 11, 2017
AB			EQ 1GM BASE	A090978	003	Aug 11, 2017

LAPATINIB DITOSYLATE

TABLET;ORAL

LAPATINIB DITOSYLATE

AB		NATCO PHARMA LTD	EQ 250MG BASE	A203007	001	Sep 29, 2020
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TYKERB

AB	+	NOVARTIS	EQ 250MG BASE	N022059	001	Mar 13, 2007
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LAROTRECTINIB SULFATE

CAPSULE;ORAL

VITRAKVI

+	BAYER HLTHCARE	EQ 25MG BASE	N210861	001	Nov 26, 2018
+		EQ 100MG BASE	N210861	002	Nov 26, 2018

SOLUTION;ORAL

VITRAKVI

+	BAYER HEALTHCARE	EQ 20MG BASE/ML	N211710	001	Nov 26, 2018
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LASMITIDAN SUCCINATE

TABLET;ORAL

REYVOW

+	ELI LILLY AND CO	EQ 50MG BASE	N211280	001	Jan 31, 2020
+		EQ 100MG BASE	N211280	002	Jan 31, 2020

LATANOPROST

EMULSION;OPHTHALMIC

XELPROS

+	SUN PHARM	0.005%	N206185	001	Sep 12, 2018
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SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

AT		AMRING PHARMS	0.005%	A200925	001	Mar 22, 2011
AT		BAUSCH AND LOMB	0.005%	A201006	001	Mar 22, 2011
AT		CARNEGIE	0.005%	A202077	001	Feb 11, 2013
AT		FDC LTD	0.005%	A202442	001	Apr 22, 2016
AT		SANDOZ	0.005%	A091449	001	Mar 22, 2011

PRESCRIPTION DRUG PRODUCT LIST

LATANOPROST

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

AT	+	SOMERSET	0.005%	A201786 001	Mar 22, 2011
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XALATAN

AT	+	UPJOHN	0.005%	N020597 001	Jun 05, 1996
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IYUZEH

+	!	THEA PHARMA	0.005%	N216472 001	Dec 13, 2022
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LATANOPROST; NETARSUDIL DIMESYLATE

SOLUTION/DROPS;OPHTHALMIC

ROCKLATAN

+	!	ALCON LABS INC	0.005%;EQ 0.02% BASE	N208259 001	Mar 12, 2019
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LATANOPROSTENE BUNOD

SOLUTION/DROPS;OPHTHALMIC

VYZULTA

+	!	BAUSCH AND LOMB	0.024%	N207795 001	Nov 02, 2017
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LEDIPASVIR; SOFOSBUVIR

PELLETS;ORAL

HARVONI

+		GILEAD SCIENCES INC	33.75MG;150MG/PACKET	N212477 001	Aug 28, 2019
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+	!		45MG;200MG/PACKET	N212477 002	Aug 28, 2019
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TABLET;ORAL

HARVONI

+		GILEAD SCIENCES INC	45MG;200MG	N205834 002	Aug 28, 2019
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+	!		90MG;400MG	N205834 001	Oct 10, 2014
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LEFAMULIN ACETATE

TABLET;ORAL

XENLETA

+	!	NABRIVA	EQ 600MG BASE	N211672 001	Aug 19, 2019
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LEFLUNOMIDE

TABLET;ORAL

ARAVA

AB	+	SANOFI AVENTIS US	10MG	N020905 001	Sep 10, 1998
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AB	+	!	20MG	N020905 002	Sep 10, 1998
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LEFLUNOMIDE

AB		ABHAI LLC	10MG	A212453 001	Jun 03, 2019
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AB			20MG	A212453 002	Jun 03, 2019
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AB		AET PHARMA	10MG	A213497 001	May 10, 2021
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AB			20MG	A213497 002	May 10, 2021
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AB		ALEMBIC PHARMS LTD	10MG	A091369 001	Nov 21, 2011
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AB			20MG	A091369 002	Nov 21, 2011
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AB		APOTEX INC	10MG	A077090 001	Sep 13, 2005
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AB			20MG	A077090 002	Sep 13, 2005
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AB		AUROBINDO PHARMA	10MG	A213652 001	Mar 29, 2021
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AB			20MG	A213652 002	Mar 29, 2021
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AB		HERITAGE PHARMS	10MG	A077086 001	Sep 13, 2005
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AB			20MG	A077086 002	Sep 13, 2005
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AB		LUPIN LTD	10MG	A211863 001	Feb 04, 2020
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AB			20MG	A211863 002	Feb 04, 2020
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AB		WANBANG BIOPHARMS	10MG	A077087 001	Sep 13, 2005
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AB			20MG	A077087 002	Sep 13, 2005
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AB		ZYDUS	10MG	A212308 001	Apr 24, 2019
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AB			20MG	A212308 002	Apr 24, 2019
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ARAVA

+	!	SANOFI AVENTIS US	100MG	N020905 003	Sep 10, 1998
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LEMBOREXANT

TABLET;ORAL

DAYVIGO

+		EISAI INC	5MG	N212028 001	Apr 07, 2020
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+	!		10MG	N212028 002	Apr 07, 2020
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LENACAPAVIR SODIUM

SOLUTION;SUBCUTANEOUS

SUNLENCA

+	!	GILEAD SCIENCES INC	EQ 463.5MG BASE/1.5ML (EQ 309MG	N215973 001	Dec 22, 2022
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BASE/ML)

TABLET;ORAL

SUNLENCA

+	!	GILEAD SCIENCES INC	EQ 300MG BASE	N215974 001	Dec 22, 2022
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PRESCRIPTION DRUG PRODUCT LIST

LENALIDOMIDE

CAPSULE; ORAL

LENALIDOMIDE

<u>AB</u>	APOTEX	<u>2.5MG</u>	<u>A211022</u>	<u>005</u>	Mar 07, 2023
<u>AB</u>		<u>5MG</u>	<u>A211022</u>	<u>001</u>	Aug 30, 2022
<u>AB</u>		<u>10MG</u>	<u>A211022</u>	<u>002</u>	Aug 30, 2022
<u>AB</u>		<u>15MG</u>	<u>A211022</u>	<u>003</u>	Aug 30, 2022
<u>AB</u>		<u>20MG</u>	<u>A211022</u>	<u>006</u>	Mar 07, 2023
<u>AB</u>		<u>25MG</u>	<u>A211022</u>	<u>004</u>	Aug 30, 2022
<u>AB</u>	ARROW INTL	<u>2.5MG</u>	<u>A201452</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A201452</u>	<u>001</u>	May 21, 2021
<u>AB</u>		<u>10MG</u>	<u>A201452</u>	<u>002</u>	May 21, 2021
<u>AB</u>		<u>15MG</u>	<u>A201452</u>	<u>003</u>	May 21, 2021
<u>AB</u>		<u>20MG</u>	<u>A201452</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A201452</u>	<u>004</u>	May 21, 2021
<u>AB</u>	CIPLA	<u>2.5MG</u>	<u>A214618</u>	<u>001</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A210435</u>	<u>001</u>	Sep 06, 2022
<u>AB</u>		<u>10MG</u>	<u>A210435</u>	<u>002</u>	Sep 06, 2022
<u>AB</u>		<u>15MG</u>	<u>A210435</u>	<u>003</u>	Sep 06, 2022
<u>AB</u>		<u>20MG</u>	<u>A210435</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A210435</u>	<u>004</u>	Sep 06, 2022
<u>AB</u>	DR REDDYS	<u>2.5MG</u>	<u>A209348</u>	<u>001</u>	Oct 14, 2021
<u>AB</u>		<u>5MG</u>	<u>A209348</u>	<u>003</u>	Aug 30, 2022
<u>AB</u>		<u>10MG</u>	<u>A209348</u>	<u>004</u>	Aug 30, 2022
<u>AB</u>		<u>15MG</u>	<u>A209348</u>	<u>005</u>	Aug 30, 2022
<u>AB</u>		<u>20MG</u>	<u>A209348</u>	<u>002</u>	Oct 14, 2021
<u>AB</u>		<u>25MG</u>	<u>A209348</u>	<u>006</u>	Aug 30, 2022
<u>AB</u>	EUGIA PHARMA	<u>2.5MG</u>	<u>A213885</u>	<u>001</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A213885</u>	<u>002</u>	Mar 06, 2023
<u>AB</u>		<u>10MG</u>	<u>A213885</u>	<u>003</u>	Mar 06, 2023
<u>AB</u>		<u>15MG</u>	<u>A213885</u>	<u>004</u>	Mar 06, 2023
<u>AB</u>		<u>20MG</u>	<u>A213885</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A213885</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>	HETERO LABS LTD V	<u>2.5MG</u>	<u>A212414</u>	<u>001</u>	May 11, 2023
<u>AB</u>		<u>5MG</u>	<u>A212414</u>	<u>002</u>	May 11, 2023
<u>AB</u>		<u>10MG</u>	<u>A212414</u>	<u>003</u>	May 11, 2023
<u>AB</u>		<u>15MG</u>	<u>A212414</u>	<u>004</u>	May 11, 2023
<u>AB</u>		<u>20MG</u>	<u>A212414</u>	<u>005</u>	May 11, 2023
<u>AB</u>		<u>25MG</u>	<u>A212414</u>	<u>006</u>	May 11, 2023
<u>AB</u>	LOTUS PHARM CO LTD	<u>2.5MG</u>	<u>A210480</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A210480</u>	<u>001</u>	Aug 31, 2022
<u>AB</u>		<u>10MG</u>	<u>A210480</u>	<u>002</u>	Aug 31, 2022
<u>AB</u>		<u>15MG</u>	<u>A210480</u>	<u>003</u>	Aug 31, 2022
<u>AB</u>		<u>20MG</u>	<u>A210480</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A210480</u>	<u>004</u>	Aug 31, 2022
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A213912</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A213912</u>	<u>001</u>	Aug 30, 2022
<u>AB</u>		<u>10MG</u>	<u>A213912</u>	<u>002</u>	Aug 30, 2022
<u>AB</u>		<u>15MG</u>	<u>A213912</u>	<u>003</u>	Aug 30, 2022
<u>AB</u>		<u>20MG</u>	<u>A213912</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A213912</u>	<u>004</u>	Aug 30, 2022
<u>AB</u>	SUN PHARM	<u>2.5MG</u>	<u>A211846</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A211846</u>	<u>001</u>	Feb 08, 2023
<u>AB</u>		<u>10MG</u>	<u>A211846</u>	<u>002</u>	Feb 08, 2023
<u>AB</u>		<u>15MG</u>	<u>A211846</u>	<u>003</u>	Feb 08, 2023
<u>AB</u>		<u>20MG</u>	<u>A211846</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A211846</u>	<u>004</u>	Feb 08, 2023
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A210154</u>	<u>005</u>	Mar 06, 2023
<u>AB</u>		<u>5MG</u>	<u>A210154</u>	<u>001</u>	Sep 12, 2022
<u>AB</u>		<u>10MG</u>	<u>A210154</u>	<u>002</u>	Sep 12, 2022
<u>AB</u>		<u>15MG</u>	<u>A210154</u>	<u>003</u>	Sep 12, 2022
<u>AB</u>		<u>20MG</u>	<u>A210154</u>	<u>006</u>	Mar 06, 2023
<u>AB</u>		<u>25MG</u>	<u>A210154</u>	<u>004</u>	Sep 12, 2022
<u>REVLIMID</u>					
<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>2.5MG</u>	<u>N021880</u>	<u>005</u>	Dec 21, 2011
<u>AB</u>	+	<u>5MG</u>	<u>N021880</u>	<u>001</u>	Dec 27, 2005
<u>AB</u>	+	<u>10MG</u>	<u>N021880</u>	<u>002</u>	Dec 27, 2005
<u>AB</u>	+	<u>15MG</u>	<u>N021880</u>	<u>003</u>	Jun 29, 2006
<u>AB</u>	+	<u>20MG</u>	<u>N021880</u>	<u>006</u>	Jun 05, 2013
<u>AB</u>	+!	<u>25MG</u>	<u>N021880</u>	<u>004</u>	Jun 29, 2006

PRESCRIPTION DRUG PRODUCT LIST

LENIOLISIB PHOSPHATE

TABLET; ORAL

JOENJA

+! PHARMING EQ 70MG BASE N217759 001 Mar 24, 2023

LENVATINIB MESYLATE

CAPSULE; ORAL

LENVIMA

+ EISAI INC EQ 4MG BASE N206947 001 Feb 13, 2015

+! EQ 10MG BASE N206947 002 Feb 13, 2015

LETERMOVIR

SOLUTION; INTRAVENOUS

PREVMIS

+! MERCK SHARP DOHME 240MG/12ML (20MG/ML) N209940 001 Nov 08, 2017

+! 480MG/24ML (20MG/ML) N209940 002 Nov 08, 2017

TABLET; ORAL

PREVMIS

+ MERCK SHARP DOHME 240MG N209939 001 Nov 08, 2017

+! 480MG N209939 002 Nov 08, 2017

LETROZOLE

TABLET; ORAL

FEMARA**AB +! NOVARTIS PHARMS 2.5MG N020726 001 Jul 25, 1997****LETROZOLE****AB ACCORD HLTHCARE 2.5MG A090934 001 Jun 03, 2011****AB BEIJING YILING 2.5MG A205869 001 Nov 14, 2018****AB CARNEGIE 2.5MG A091191 001 Jun 03, 2011****AB EUGIA PHARMA 2.5MG A211717 001 Jan 11, 2019****AB NATCO PHARMA LTD 2.5MG A200161 001 Jun 03, 2011****AB TEVA PHARMS 2.5MG A090289 001 Jun 03, 2011**LETROZOLE; RIBOCICLIB SUCCINATE

TABLET; ORAL

KISQALI FEMARA CO-PACK (COPACKAGED)

+! NOVARTIS 2.5MG;EQ 200MG BASE N209935 001 May 04, 2017

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM**AP HAINAN POLY EQ 50MG BASE/VIAL A217021 001 Jul 10, 2023****AP EQ 100MG BASE/VIAL A217021 002 Jul 10, 2023****AP EQ 200MG BASE/VIAL A217021 003 Jul 10, 2023****AP EQ 500MG BASE/VIAL A217021 004 Jul 10, 2023****AP ! HIKMA EQ 50MG BASE/VIAL A089384 001 Sep 14, 1987****AP ! EQ 100MG BASE/VIAL A089717 001 Mar 28, 1988****AP KINDOS EQ 50MG BASE/VIAL A216590 001 Jul 19, 2023****AP EQ 100MG BASE/VIAL A216590 002 Jul 19, 2023****AP EQ 200MG BASE/VIAL A216590 003 Jul 19, 2023****AP EQ 350MG BASE/VIAL A216590 004 Jul 19, 2023****AP EQ 500MG BASE/VIAL A216590 005 Jul 19, 2023****LEUCOVORIN CALCIUM PRESERVATIVE FREE****AP FRESENIUS KABI USA EQ 200MG BASE/VIAL A040258 001 Feb 26, 1999****AP ! EQ 500MG BASE/VIAL A040286 001 Feb 26, 1999****AP ! HIKMA EQ 200MG BASE/VIAL A040056 001 May 23, 1995****AP ! EQ 350MG BASE/VIAL A040335 001 Apr 20, 2000****AP MYLAN LABS LTD EQ 100MG BASE/VIAL A203800 001 May 19, 2017****AP EQ 200MG BASE/VIAL A203800 002 May 19, 2017****AP EQ 350MG BASE/VIAL A203800 003 May 19, 2017****AP SAGENT PHARMS EQ 50MG BASE/VIAL A200753 001 Sep 06, 2012****AP EQ 100MG BASE/VIAL A200753 002 Sep 06, 2012****AP EQ 200MG BASE/VIAL A200753 003 Sep 06, 2012****AP EQ 350MG BASE/VIAL A200855 001 Sep 06, 2012****AP SAGENT PHARMS INC EQ 500MG BASE/VIAL A209110 001 Oct 26, 2017**

LEUCOVORIN CALCIUM

! FRESENIUS KABI USA EQ 10MG BASE/ML A207226 001 Jul 27, 2018

! EQ 10MG BASE/ML A207241 001 Mar 14, 2018

TABLET; ORAL

LEUCOVORIN CALCIUM**AB BARR EQ 5MG BASE A071198 001 Sep 24, 1987****AB EQ 25MG BASE A071199 001 Sep 24, 1987****AB EPIC PHARMA LLC EQ 5MG BASE A074544 001 Aug 28, 1997****AB EQ 10MG BASE A074544 003 May 19, 2021****AB EQ 15MG BASE A074544 004 May 19, 2021**

PRESCRIPTION DRUG PRODUCT LIST

LEUCOVORIN CALCIUM

TABLET; ORAL

LEUCOVORIN CALCIUM

<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A074544 002</u>	Aug 28, 1997
<u>AB</u>	HIKMA	<u>EQ 5MG BASE</u>	<u>A072733 001</u>	Feb 22, 1993
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A072734 001</u>	Feb 22, 1993
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A072735 001</u>	Feb 22, 1993
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A072736 001</u>	Feb 22, 1993
<u>AB</u>	LEADING	<u>EQ 5MG BASE</u>	<u>A213929 001</u>	Oct 22, 2020
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A213929 002</u>	Oct 22, 2020
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A213929 003</u>	Oct 22, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213929 004</u>	Oct 22, 2020
<u>AB</u>	NOVAST LABS	<u>EQ 5MG BASE</u>	<u>A211132 001</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A211132 002</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A211132 003</u>	Jul 30, 2020
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A211132 004</u>	Jul 30, 2020

LEUPROLIDE ACETATE

FOR SUSPENSION; INTRAMUSCULAR

LEUPROLIDE ACETATE FOR DEPOT SUSPENSION

+! INVAGEN PHARMS 22.5MG/VIAL

N205054 001 Aug 28, 2018

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

<u>AP</u>	AMNEAL	<u>1MG/0.2ML</u>	<u>A215336 001</u>	Oct 28, 2022
<u>AP</u>	EUGIA PHARMA	<u>1MG/0.2ML</u>	<u>A212963 001</u>	Jun 06, 2022
<u>AP</u>	MEITHEAL	<u>1MG/0.2ML</u>	<u>A075471 001</u>	Oct 25, 2000
<u>AP</u>	RK PHARMA	<u>1MG/0.2ML</u>	<u>A213829 001</u>	Aug 13, 2021
<u>AP</u>	!	<u>1MG/0.2ML</u>	<u>A074728 001</u>	Aug 04, 1998
<u>AP</u>	SUN PHARM	<u>1MG/0.2ML</u>	<u>A078885 001</u>	Mar 09, 2009

LUPRON DEPOT

+! ABBVIE ENDOCRINE
INC

3.75MG

N020011 002 Oct 26, 1995

+!

7.5MG

N019732 001 Jan 26, 1989

+!

11.25MG

N020708 001 Mar 07, 1997

+

22.5MG

N020517 001 Dec 22, 1995

+!

30MG

N020517 002 May 30, 1997

+!

45MG

N020517 003 Jun 17, 2011

POWDER; INTRAMUSCULAR

LUPRON DEPOT-PED KIT

+! ABBVIE ENDOCRINE
INC

7.5MG

N020263 002 Apr 16, 1993

+!

11.25MG

N020263 005 Jan 21, 1994

+!

11.25MG

N020263 007 Aug 15, 2011

+!

15MG

N020263 006 Jan 21, 1994

+!

30MG

N020263 008 Aug 15, 2011

+!

45MG

N020263 009 Apr 14, 2023

POWDER; SUBCUTANEOUS

ELIGARD KIT

+! TOLMAR

7.5MG

N021343 001 Jan 23, 2002

+!

22.5MG

N021379 001 Jul 24, 2002

+!

30MG

N021488 001 Feb 13, 2003

+!

45MG

N021731 001 Dec 14, 2004

FENSOLVI KIT

+! TOLMAR

45MG

N213150 001 May 01, 2020

LEUPROLIDE MESYLATE

EMULSION; SUBCUTANEOUS

CAMCEVI KIT

+! ACCORD

EQ 42MG BASE

N211488 001 May 25, 2021

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

<u>AN</u>	CIPLA	<u>EQ 0.021% BASE</u>	<u>A078171 002</u>	Dec 13, 2013
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A078171 003</u>	Dec 13, 2013
<u>AN</u>		<u>EQ 0.0103% BASE</u>	<u>A078171 001</u>	Dec 13, 2013
<u>AN</u>	IMPAX LABS INC	<u>EQ 0.0103% BASE</u>	<u>A077756 003</u>	Apr 09, 2008
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A077756 001</u>	Apr 09, 2008
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A077756 002</u>	Apr 09, 2008
<u>AN</u>	LUOXIN AUROVITAS	<u>EQ 0.25% BASE</u>	<u>A207628 001</u>	Jan 31, 2017
<u>AN</u>		<u>EQ 0.0103% BASE</u>	<u>A207625 001</u>	Dec 30, 2016
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A207625 002</u>	Dec 30, 2016
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A207625 003</u>	Dec 30, 2016
<u>AN</u>	MYLAN SPECIALITY LP	<u>EQ 0.25% BASE</u>	<u>A078309 001</u>	Mar 20, 2009

PRESCRIPTION DRUG PRODUCT LIST

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

<u>AN</u>	RITEDOSE CORP	<u>EQ 0.0103% BASE</u>	<u>A203653 001</u>	Mar 22, 2016
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A203653 002</u>	Mar 22, 2016
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A203653 003</u>	Mar 22, 2016
<u>AN</u>	SUN PHARM	<u>EQ 0.0103% BASE</u>	<u>A207820 001</u>	Nov 05, 2018
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A207820 002</u>	Nov 05, 2018
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A207820 003</u>	Nov 05, 2018
<u>AN</u>	TEVA PARENTERAL	<u>EQ 0.25% BASE</u>	<u>A200875 001</u>	Sep 11, 2014
<u>AN</u>	TEVA PHARMS USA	<u>EQ 0.0103% BASE</u>	<u>A090297 001</u>	Apr 26, 2013
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A090297 002</u>	Apr 26, 2013
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A090297 003</u>	Apr 26, 2013

XOPENEX

<u>AN</u>	+! HIKMA	<u>EQ 0.0103% BASE</u>	<u>N020837 003</u>	Jan 30, 2002
<u>AN</u>	+!	<u>EQ 0.021% BASE</u>	<u>N020837 001</u>	Mar 25, 1999
<u>AN</u>	+!	<u>EQ 0.042% BASE</u>	<u>N020837 002</u>	Mar 25, 1999
<u>AN</u>	+!	<u>EQ 0.25% BASE</u>	<u>N020837 004</u>	Jul 18, 2003

LEVALBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

+! LUPIN

EQ 0.045MG BASE/INH

N021730 001 Mar 11, 2005

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

KEPPRA

<u>AP</u>	+! UCB INC	<u>500MG/5ML (100MG/ML)</u>	<u>N021872 001</u>	Jul 31, 2006
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LEVETIRACETAM

<u>AP</u>	BAXTER HLTHCARE CORP	<u>500MG/100ML (5MG/ML)</u>	<u>A217059 001</u>	Oct 02, 2023
<u>AP</u>		<u>1GM/100ML (10MG/ML)</u>	<u>A217059 002</u>	Oct 02, 2023
<u>AP</u>		<u>1.5GM/100ML (15MG/ML)</u>	<u>A217059 003</u>	Oct 02, 2023
<u>AP</u>	BEOWULF ASSET	<u>500MG/5ML (100MG/ML)</u>	<u>A091485 001</u>	Aug 05, 2011
<u>AP</u>	EUGIA PHARMA	<u>500MG/5ML (100MG/ML)</u>	<u>A204312 001</u>	Feb 01, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>500MG/5ML (100MG/ML)</u>	<u>A090876 001</u>	Aug 13, 2015
<u>AP</u>	HAINAN POLY PHARM	<u>500MG/5ML (100MG/ML)</u>	<u>A209781 001</u>	Mar 20, 2018
<u>AP</u>	HIKMA FARMACEUTICA	<u>500MG/5ML (100MG/ML)</u>	<u>A090981 001</u>	Oct 13, 2011
<u>AP</u>	HOSPIRA INC	<u>500MG/5ML (100MG/ML)</u>	<u>A202869 001</u>	Apr 06, 2012
<u>AP</u>	MICRO LABS	<u>500MG/5ML (100MG/ML)</u>	<u>A211954 001</u>	Aug 09, 2019
<u>AP</u>	MSN	<u>500MG/5ML (100MG/ML)</u>	<u>A215980 001</u>	Nov 15, 2022
<u>AP</u>	MYLAN LABS LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A203308 001</u>	Sep 16, 2016
<u>AP</u>	PRINSTON INC	<u>500MG/5ML (100MG/ML)</u>	<u>A209474 001</u>	Mar 28, 2022
<u>AP</u>	SAGENT PHARMS	<u>500MG/5ML (100MG/ML)</u>	<u>A091627 001</u>	Jun 26, 2013
<u>AP</u>	SUN PHARM INDS LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A090754 001</u>	Jun 16, 2010

LEVETIRACETAM IN SODIUM CHLORIDE

<u>AP</u>	EUGIA PHARMA	<u>500MG/100ML (5MG/ML)</u>	<u>A207160 001</u>	Jan 04, 2017
<u>AP</u>		<u>1GM/100ML (10MG/ML)</u>	<u>A207160 002</u>	Jan 04, 2017
<u>AP</u>		<u>1.5GM/100ML (15MG/ML)</u>	<u>A207160 003</u>	Jan 04, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>500MG/100ML (5MG/ML)</u>	<u>A208619 001</u>	Jan 31, 2023
<u>AP</u>		<u>1GM/100ML (10MG/ML)</u>	<u>A208619 002</u>	Jan 31, 2023
<u>AP</u>		<u>1.5GM/100ML (15MG/ML)</u>	<u>A208619 003</u>	Jan 31, 2023
<u>AP</u>	GLAND PHARMA LTD	<u>500MG/100ML (5MG/ML)</u>	<u>A206880 001</u>	Oct 25, 2017
<u>AP</u>		<u>1GM/100ML (10MG/ML)</u>	<u>A206880 002</u>	Oct 25, 2017
<u>AP</u>		<u>1.5GM/100ML (15MG/ML)</u>	<u>A206880 003</u>	Oct 25, 2017
<u>AP</u>	+! HQ SPCLT PHARMA	<u>500MG/100ML (5MG/ML)</u>	<u>N202543 001</u>	Nov 09, 2011
<u>AP</u>	+!	<u>1GM/100ML (10MG/ML)</u>	<u>N202543 002</u>	Nov 09, 2011
<u>AP</u>	+!	<u>1.5GM/100ML (15MG/ML)</u>	<u>N202543 003</u>	Nov 09, 2011
<u>AP</u>	NEXUS	<u>500MG/100ML (5MG/ML)</u>	<u>A213532 001</u>	Jul 06, 2020
<u>AP</u>		<u>1GM/100ML (10MG/ML)</u>	<u>A213532 002</u>	Jul 06, 2020
<u>AP</u>		<u>1.5GM/100ML (15MG/ML)</u>	<u>A213532 003</u>	Jul 06, 2020
	+! HQ SPCLT PHARMA	<u>250MG/50ML (5MG/ML)</u>	<u>N202543 004</u>	Dec 14, 2020

SOLUTION; ORAL

KEPPRA

<u>AA</u>	+! UCB INC	<u>100MG/ML</u>	<u>N021505 001</u>	Jul 15, 2003
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LEVETIRACETAM

<u>AA</u>	ACI	<u>100MG/ML</u>	<u>A078582 001</u>	Jan 15, 2009
<u>AA</u>	ACTAVIS MID	<u>100MG/ML</u>	<u>A078976 001</u>	Jan 15, 2009
	ATLANTIC			
<u>AA</u>	ALEMBIC	<u>100MG/ML</u>	<u>A203067 001</u>	May 09, 2013
<u>AA</u>	AMNEAL PHARMS	<u>100MG/ML</u>	<u>A090992 001</u>	Oct 27, 2009
<u>AA</u>	AUROBINDO PHARMA	<u>100MG/ML</u>	<u>A079063 001</u>	Jan 15, 2009
<u>AA</u>	BELCHER	<u>100MG/ML</u>	<u>A090461 001</u>	Sep 30, 2010
<u>AA</u>	BIONPHARMA	<u>100MG/ML</u>	<u>A079120 001</u>	Jan 16, 2009

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

SOLUTION;ORAL

LEVETIRACETAM

<u>AA</u>	CHARTWELL MOLECULAR	<u>100MG/ML</u>	<u>A090263</u>	<u>001</u>	Apr 03, 2009
<u>AA</u>	HETERO LABS LTD III	<u>100MG/ML</u>	<u>A203052</u>	<u>001</u>	Feb 28, 2013
<u>AA</u>	HIKMA	<u>100MG/ML</u>	<u>A090601</u>	<u>001</u>	Feb 28, 2012
<u>AA</u>	LUPIN LTD	<u>100MG/ML</u>	<u>A090893</u>	<u>001</u>	Oct 17, 2011
<u>AA</u>	MSN	<u>100MG/ML</u>	<u>A214757</u>	<u>001</u>	Jul 15, 2022
<u>AA</u>	PHARM ASSOC	<u>100MG/ML</u>	<u>A201157</u>	<u>001</u>	Jun 04, 2015
<u>AA</u>	QUAGEN	<u>100MG/ML</u>	<u>A090079</u>	<u>001</u>	Apr 11, 2012
<u>AA</u>	STRIDES PHARMA	<u>100MG/ML</u>	<u>A214673</u>	<u>001</u>	Nov 20, 2023
<u>AA</u>	TARO	<u>100MG/ML</u>	<u>A078774</u>	<u>001</u>	Feb 10, 2009

TABLET;ORAL

KEPPRA

<u>AB</u>	<u>+</u>	UCB INC	<u>250MG</u>	<u>N021035</u>	<u>001</u>	Nov 30, 1999
<u>AB</u>	<u>+</u>		<u>500MG</u>	<u>N021035</u>	<u>002</u>	Nov 30, 1999
<u>AB</u>	<u>+</u>		<u>750MG</u>	<u>N021035</u>	<u>003</u>	Nov 30, 1999
<u>AB</u>	<u>+</u>		<u>1GM</u>	<u>N021035</u>	<u>004</u>	Jan 06, 2006

LEVETIRACETAM

<u>AB</u>		ACCORD HLTHCARE	<u>250MG</u>	<u>A090843</u>	<u>001</u>	Feb 14, 2011
<u>AB</u>			<u>500MG</u>	<u>A090843</u>	<u>002</u>	Feb 14, 2011
<u>AB</u>			<u>750MG</u>	<u>A090843</u>	<u>003</u>	Feb 14, 2011
<u>AB</u>			<u>1GM</u>	<u>A090843</u>	<u>004</u>	Feb 14, 2011
<u>AB</u>		ACI	<u>250MG</u>	<u>A078042</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078042</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078042</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A078042</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>		ALKEM LABS LTD	<u>250MG</u>	<u>A216375</u>	<u>001</u>	May 27, 2022
<u>AB</u>			<u>500MG</u>	<u>A216375</u>	<u>002</u>	May 27, 2022
<u>AB</u>			<u>750MG</u>	<u>A216375</u>	<u>003</u>	May 27, 2022
<u>AB</u>			<u>1GM</u>	<u>A216375</u>	<u>004</u>	May 27, 2022
<u>AB</u>		AUROBINDO PHARMA	<u>250MG</u>	<u>A078993</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078993</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078993</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A078993</u>	<u>004</u>	Jan 15, 2009
<u>AB</u>		CHARTWELL RX	<u>250MG</u>	<u>A201293</u>	<u>001</u>	Jun 14, 2011
<u>AB</u>			<u>500MG</u>	<u>A201293</u>	<u>002</u>	Jun 14, 2011
<u>AB</u>			<u>750MG</u>	<u>A201293</u>	<u>003</u>	Jun 14, 2011
<u>AB</u>			<u>1GM</u>	<u>A201293</u>	<u>004</u>	Jun 14, 2011
<u>AB</u>		DR REDDYS LABS LTD	<u>250MG</u>	<u>A076920</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A076920</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A076920</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A078904</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		GRANULES	<u>250MG</u>	<u>A217878</u>	<u>001</u>	Jun 13, 2023
<u>AB</u>			<u>500MG</u>	<u>A217878</u>	<u>002</u>	Jun 13, 2023
<u>AB</u>			<u>750MG</u>	<u>A217878</u>	<u>003</u>	Jun 13, 2023
<u>AB</u>			<u>1GM</u>	<u>A217878</u>	<u>004</u>	Jun 13, 2023
<u>AB</u>		HETERO LABS LTD III	<u>250MG</u>	<u>A090515</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>			<u>500MG</u>	<u>A090515</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>			<u>750MG</u>	<u>A090515</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>			<u>1GM</u>	<u>A090515</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		INVAGEN PHARMS	<u>250MG</u>	<u>A078234</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078234</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078234</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>		LUPIN	<u>250MG</u>	<u>A078154</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078154</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078154</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A090025</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>		MSN	<u>250MG</u>	<u>A214815</u>	<u>001</u>	Oct 07, 2022
<u>AB</u>			<u>500MG</u>	<u>A214815</u>	<u>002</u>	Oct 07, 2022
<u>AB</u>			<u>750MG</u>	<u>A214815</u>	<u>003</u>	Oct 07, 2022
<u>AB</u>			<u>1GM</u>	<u>A214815</u>	<u>004</u>	Oct 07, 2022
<u>AB</u>		MYLAN	<u>500MG</u>	<u>A076919</u>	<u>002</u>	Nov 04, 2008
<u>AB</u>			<u>750MG</u>	<u>A076919</u>	<u>003</u>	Nov 04, 2008
<u>AB</u>			<u>1GM</u>	<u>A090261</u>	<u>001</u>	Dec 08, 2009
<u>AB</u>		ORBION PHARMS	<u>250MG</u>	<u>A078526</u>	<u>001</u>	Jan 15, 2009
<u>AB</u>			<u>500MG</u>	<u>A078526</u>	<u>002</u>	Jan 15, 2009
<u>AB</u>			<u>750MG</u>	<u>A078526</u>	<u>003</u>	Jan 15, 2009
<u>AB</u>			<u>1GM</u>	<u>A090484</u>	<u>001</u>	Aug 05, 2010
<u>AB</u>		OXFORD PHARMS	<u>250MG</u>	<u>A077319</u>	<u>001</u>	Mar 20, 2009
<u>AB</u>			<u>500MG</u>	<u>A077319</u>	<u>002</u>	Mar 20, 2009
<u>AB</u>			<u>750MG</u>	<u>A077319</u>	<u>003</u>	Mar 20, 2009

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

<u>AB</u>	PRINSTON INC	<u>250MG</u>	<u>A078106 001</u>	Feb 10, 2009
<u>AB</u>		<u>500MG</u>	<u>A078106 002</u>	Feb 10, 2009
<u>AB</u>		<u>750MG</u>	<u>A078106 003</u>	Feb 10, 2009
<u>AB</u>		<u>1GM</u>	<u>A078106 004</u>	Feb 10, 2009
<u>AB</u>	RISING	<u>250MG</u>	<u>A090767 001</u>	Jul 28, 2010
<u>AB</u>		<u>500MG</u>	<u>A090767 002</u>	Jul 28, 2010
<u>AB</u>		<u>750MG</u>	<u>A090767 003</u>	Jul 28, 2010
<u>AB</u>		<u>1GM</u>	<u>A090767 004</u>	Jul 28, 2010
<u>AB</u>	SCIEGEN PHARMS INC	<u>250MG</u>	<u>A215069 003</u>	May 27, 2022
<u>AB</u>		<u>500MG</u>	<u>A215069 004</u>	May 27, 2022
<u>AB</u>		<u>750MG</u>	<u>A215069 001</u>	Jun 11, 2021
<u>AB</u>		<u>1GM</u>	<u>A215069 002</u>	Jun 11, 2021
<u>AB</u>	STALLION LABS	<u>250MG</u>	<u>A079042 001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A079042 002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A079042 003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A079042 004</u>	Jan 15, 2009
<u>AB</u>	TARO	<u>250MG</u>	<u>A078960 004</u>	Feb 01, 2010
<u>AB</u>		<u>500MG</u>	<u>A078960 003</u>	Feb 01, 2010
<u>AB</u>		<u>750MG</u>	<u>A078960 002</u>	Feb 01, 2010
<u>AB</u>		<u>1GM</u>	<u>A078960 001</u>	Feb 01, 2010
<u>AB</u>	TORRENT PHARMS	<u>250MG</u>	<u>A078858 001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078858 002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078858 003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078858 004</u>	Jan 15, 2009
<u>AB</u>	VIWIT PHARM	<u>250MG</u>	<u>A078869 001</u>	Mar 13, 2009
<u>AB</u>		<u>500MG</u>	<u>A078869 002</u>	Mar 13, 2009
<u>AB</u>		<u>750MG</u>	<u>A078869 003</u>	Mar 13, 2009
<u>AB</u>		<u>1GM</u>	<u>A078869 004</u>	Mar 13, 2009
<u>AB</u>	ZHEJIANG JINGXIN	<u>250MG</u>	<u>A091491 001</u>	Dec 14, 2010
<u>AB</u>		<u>500MG</u>	<u>A091491 002</u>	Dec 14, 2010
<u>AB</u>		<u>750MG</u>	<u>A091491 003</u>	Dec 14, 2010
<u>AB</u>		<u>1GM</u>	<u>A091491 004</u>	Dec 14, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>250MG</u>	<u>A078918 001</u>	Apr 29, 2009
<u>AB</u>		<u>1GM</u>	<u>A078918 002</u>	Apr 29, 2009

TABLET, EXTENDED RELEASE; ORAL

KEPPRA XR

<u>AB</u>	+ UCB INC	<u>500MG</u>	<u>N022285 001</u>	Sep 12, 2008
<u>AB</u>	+!	<u>750MG</u>	<u>N022285 002</u>	Feb 12, 2009

LEVETIRACETAM

<u>AB</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A091093 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091093 002</u>	Sep 12, 2011
<u>AB</u>	AIPING PHARM INC	<u>500MG</u>	<u>A204754 001</u>	Aug 26, 2016
<u>AB</u>		<u>750MG</u>	<u>A204754 002</u>	Aug 26, 2016
<u>AB</u>	ANCHEN PHARMS	<u>500MG</u>	<u>A091360 001</u>	Oct 04, 2011
<u>AB</u>		<u>750MG</u>	<u>A091360 002</u>	Oct 04, 2011
<u>AB</u>	ANDA REPOSITORY	<u>500MG</u>	<u>A204511 001</u>	Feb 23, 2016
<u>AB</u>		<u>750MG</u>	<u>A204511 002</u>	Feb 23, 2016
<u>AB</u>	APOTEX INC	<u>500MG</u>	<u>A091261 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091261 002</u>	Sep 12, 2011
<u>AB</u>	HISUN PHARM HANGZHOU	<u>500MG</u>	<u>A207175 001</u>	Sep 28, 2017
<u>AB</u>		<u>750MG</u>	<u>A207175 002</u>	Sep 28, 2017
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A091399 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091399 002</u>	Sep 12, 2011
<u>AB</u>	OVERSEAS	<u>500MG</u>	<u>A212688 002</u>	May 05, 2023
<u>AB</u>		<u>750MG</u>	<u>A212688 001</u>	Jun 11, 2020
<u>AB</u>	PHARMADAX INC	<u>500MG</u>	<u>A201464 001</u>	May 25, 2012
<u>AB</u>		<u>750MG</u>	<u>A201464 002</u>	May 25, 2012
<u>AB</u>	PRINSTON INC	<u>500MG</u>	<u>A202533 001</u>	Jul 20, 2012
<u>AB</u>		<u>500MG</u>	<u>A203468 001</u>	May 21, 2015
<u>AB</u>		<u>750MG</u>	<u>A202533 002</u>	Jul 20, 2012
<u>AB</u>		<u>750MG</u>	<u>A203468 002</u>	May 21, 2015
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A205130 001</u>	Nov 27, 2020
<u>AB</u>		<u>750MG</u>	<u>A205130 002</u>	Nov 27, 2020
<u>AB</u>	APOTEX	<u>1GM</u>	<u>A202958 001</u>	Feb 25, 2015

TABLET, FOR SUSPENSION; ORAL

SPRITAM

+	APRECIA PHARMS	250MG	<u>N207958 001</u>	Jul 31, 2015
+		500MG	<u>N207958 002</u>	Jul 31, 2015

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET, FOR SUSPENSION;ORAL

SPRITAM

+

+!

750MG

1GM

N207958 003 Jul 31, 2015

N207958 004 Jul 31, 2015

LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

BETAGAN**AT** +! ALLERGAN **0.5%****N019219 002** Dec 19, 1985LEVOBUNOLOL HYDROCHLORIDE**AT** BAUSCH AND LOMB **0.5%****A074326 001** Mar 04, 1994LEVOCARNITINE

INJECTABLE; INJECTION

CARNITOR**AP** +! LEADIANT BIOSCI INC **200MG/ML****N020182 001** Dec 16, 1992LEVOCARNITINE**AP** AM REGENT **200MG/ML****A075861 001** Jun 22, 2001**AP** HIKMA **200MG/ML****A075567 001** Mar 29, 2001

SOLUTION;ORAL

CARNITOR**AA** +! LEADIANT BIOSCI INC **1GM/10ML****N019257 001** Apr 10, 1986CARNITOR SF**AA** + LEADIANT BIOSCI INC **1GM/10ML****N019257 002** Mar 28, 2007LEVOCARNITINE**AA** HIKMA **1GM/10ML****A077399 001** Oct 25, 2007**AA** LYNE **1GM/10ML****A076851 001** Aug 10, 2004**AA** NOVITIUM PHARMA **1GM/10ML****A211676 001** Aug 14, 2019**AA** SAPTALIS PHARMS **1GM/10ML****A212533 001** Nov 10, 2021LEVOCARNITINE SF**AA** NOVITIUM PHARMA **1GM/10ML****A211676 002** Aug 14, 2019

TABLET;ORAL

CARNITOR**AB** +! LEADIANT BIOSCI INC **330MG****N018948 001** Dec 27, 1985LEVOCARNITINE**AB** NOVITIUM PHARMA **330MG****A216384 001** Dec 09, 2022**AB** RISING **330MG****A076858 001** Sep 20, 2004LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE**AA** CHARTWELL MOLECULAR **2.5MG/5ML****A204599 001** May 15, 2017**AA** ! PADAGIS US **2.5MG/5ML****A091263 001** Nov 07, 2011**AA** TARO **2.5MG/5ML****A202673 001** Jul 26, 2013LEVOCETIRIZINE HYDROCHLORIDE**AA** HETERO LABS LTD III **2.5MG/5ML****A210914 001** Apr 01, 2019

TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE**AB** DR REDDYS LABS LTD **5MG****A090392 001** Feb 24, 2011**AB** ! GLENMARK GENERICS **5MG****A090385 001** Feb 24, 2011**AB** HETERO LABS LTD III **5MG****A091264 001** Jun 29, 2012**AB** MACLEODS PHARMS LTD **5MG****A205564 001** Jan 11, 2016**AB** MICRO LABS LTD **5MG****A202046 001** Sep 17, 2013

INDIA

AB SCIEGEN PHARMS INC **5MG****A203646 001** Sep 09, 2014**AB** SUN PHARM **5MG****A090362 001** Jan 31, 2013**AB** SYNTHON PHARMS **5MG****A090229 001** Nov 26, 2010**AB** TEVA PHARMS **5MG****A090199 001** Aug 22, 2011LEVODOPA

POWDER; INHALATION

INBRIJA

+

+! ACORDA

42MG

N209184 001 Dec 21, 2018

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN**AP** BAXTER HLTHCARE **EQ 500MG/20ML (EQ 25MG/ML)****A091436 001** Jun 05, 2013

CORP

AP GLAND PHARMA LTD **EQ 500MG/20ML (EQ 25MG/ML)****A205540 001** Apr 22, 2020**AP** **EQ 750MG/30ML (EQ 25MG/ML)****A205540 002** Apr 22, 2020**AP** ! RISING **EQ 500MG/20ML (EQ 25MG/ML)****A091644 001** Jun 20, 2011**AP** ! **EQ 750MG/30ML (EQ 25MG/ML)****A091644 002** Jun 20, 2011

PRESCRIPTION DRUG PRODUCT LIST

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091397 001</u>	Aug 08, 2013
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091397 002</u>	Aug 08, 2013
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091397 003</u>	Aug 08, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A200674 001</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A200674 002</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A200674 003</u>	Jun 19, 2013
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A206908 001</u>	Dec 30, 2020
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A206908 002</u>	Dec 30, 2020
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A206908 003</u>	Dec 30, 2020
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091375 001</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091375 002</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091375 003</u>	Sep 16, 2011
<u>AP</u>	HOSPIRA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A078579 001</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A078579 002</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A078579 003</u>	Sep 03, 2015
<u>AP</u>	! INFORLIFE	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A090343 001</u>	Jul 07, 2011
<u>AP</u>	!	<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A090343 002</u>	Jul 07, 2011
<u>AP</u>	!	<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A090343 003</u>	Jul 07, 2011

SOLUTION; ORAL

LEVOFLOXACIN

<u>AA</u>	LANNETT CO INC	<u>250MG/10ML</u>	<u>A205222 001</u>	May 25, 2018
<u>AA</u>	! NOVITIUM PHARMA	<u>250MG/10ML</u>	<u>A091678 001</u>	Jun 20, 2011

SOLUTION/DROPS; OPHTHALMIC

LEVOFLOXACIN

	MICRO LABS LTD	1.5%	A205600 001	Feb 27, 2019
	INDIA			
	! RISING	0.5%	A077700 001	Dec 20, 2010

TABLET; ORAL

LEVOFLOXACIN

<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A090787 001</u>	Sep 29, 2011
<u>AB</u>		<u>500MG</u>	<u>A090787 002</u>	Sep 29, 2011
<u>AB</u>		<u>750MG</u>	<u>A090787 003</u>	Sep 29, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A201043 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A201043 002</u>	Jun 20, 2011
<u>AB</u>	!	<u>750MG</u>	<u>A201043 003</u>	Jun 20, 2011
<u>AB</u>	CADILA	<u>250MG</u>	<u>A077652 001</u>	Sep 07, 2012
<u>AB</u>		<u>500MG</u>	<u>A077652 002</u>	Sep 07, 2012
<u>AB</u>		<u>750MG</u>	<u>A077652 003</u>	Sep 07, 2012
<u>AB</u>	CELLTRION	<u>250MG</u>	<u>A090367 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A090367 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A090367 003</u>	Jun 20, 2011
<u>AB</u>	CHARTWELL MOLECULAR	<u>250MG</u>	<u>A076890 001</u>	Mar 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A076890 002</u>	Mar 30, 2012
<u>AB</u>		<u>750MG</u>	<u>A076890 003</u>	Mar 30, 2012
<u>AB</u>	DR REDDYS LABS INC	<u>250MG</u>	<u>A076710 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076710 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A076710 003</u>	Jun 20, 2011
<u>AB</u>	GLENMARK GENERICS	<u>250MG</u>	<u>A200250 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A200250 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A200250 003</u>	Jun 20, 2011
<u>AB</u>	HEC PHARM	<u>250MG</u>	<u>A204968 001</u>	Feb 05, 2019
<u>AB</u>		<u>500MG</u>	<u>A204968 002</u>	Feb 05, 2019
<u>AB</u>		<u>750MG</u>	<u>A204968 003</u>	Feb 05, 2019
<u>AB</u>	HETERO LABS LTD V	<u>250MG</u>	<u>A202801 001</u>	Jan 08, 2015
<u>AB</u>		<u>500MG</u>	<u>A202801 002</u>	Jan 08, 2015
<u>AB</u>		<u>750MG</u>	<u>A202801 003</u>	Jan 08, 2015
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A078424 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A078424 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A078424 003</u>	Jun 20, 2011
<u>AB</u>	MACLEODS PHARMS LTD	<u>250MG</u>	<u>A200839 001</u>	Mar 22, 2012
<u>AB</u>		<u>500MG</u>	<u>A200839 002</u>	Mar 22, 2012
<u>AB</u>		<u>750MG</u>	<u>A200839 003</u>	Mar 22, 2012
<u>AB</u>	ORBION PHARMS	<u>250MG</u>	<u>A202200 001</u>	Jan 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A202200 002</u>	Jan 30, 2012
<u>AB</u>		<u>750MG</u>	<u>A202200 003</u>	Jan 30, 2012
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A077438 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A077438 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A077438 003</u>	Jun 20, 2011

PRESCRIPTION DRUG PRODUCT LIST

LEVOFLOXACIN

TABLET; ORAL

LEVOFLOXACIN

AB	TEVA	250MG	A076361 001	Jun 20, 2011
AB		500MG	A076361 002	Jun 20, 2011
AB		750MG	A076361 003	Jun 20, 2011

LEVOKETOCONAZOLE

TABLET; ORAL

RECORLEV

+	STRONGBRIDGE	150MG	N214133 001	Dec 30, 2021
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LEVOLEUCOVORIN

POWDER; INTRAVENOUS

KHAPZORY

+	ACROTECH BIOPHARMA	175MG/VIAL	N211226 001	Oct 19, 2018
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LEVOLEUCOVORIN CALCIUM

POWDER; INTRAVENOUS

FUSILEV

AP	+	ACROTECH BIOPHARMA	EQ 50MG BASE/VIAL	N020140 001	Mar 07, 2008
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LEVOLEUCOVORIN CALCIUM

AP		MEITHEAL	EQ 50MG BASE/VIAL	A211003 001	Aug 22, 2019
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SOLUTION; INTRAVENOUS

LEVOLEUCOVORIN CALCIUM

AP		AMNEAL	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A207548 001	Sep 08, 2017
AP		GLAND PHARMA LTD	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A210892 001	Sep 14, 2018
AP	!		EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A210892 002	Sep 14, 2018
AP		HAINAN POLY PHARM	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A217314 001	May 22, 2023
AP			EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A217314 002	May 22, 2023
AP	!	MEITHEAL	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A211002 001	Aug 16, 2019
AP		SANDOZ	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A203563 001	Mar 09, 2015

LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

LEVOMILNACIPRAN HYDROCHLORIDE

AB		PRINSTON INC	EQ 20MG BASE	A210771 001	Mar 20, 2023
AB			EQ 40MG BASE	A210771 002	Mar 20, 2023
AB			EQ 80MG BASE	A210771 003	Mar 20, 2023
AB			EQ 120MG BASE	A210771 004	Mar 20, 2023
		FETZIMA			
	+	ABBVIE	EQ 20MG BASE	N204168 001	Jul 25, 2013
	+		EQ 40MG BASE	N204168 002	Jul 25, 2013
	+		EQ 80MG BASE	N204168 003	Jul 25, 2013
	+		EQ 120MG BASE	N204168 004	Jul 25, 2013

LEVONORGESTREL

INTRAUTERINE DEVICE; INTRAUTERINE

KYLEENA

+	BAYER HLTHCARE	19.5MG	N208224 001	Sep 16, 2016
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LILETTA

+	MEDICINES360	52MG	N206229 001	Feb 26, 2015
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MIRENA

+	BAYER HLTHCARE	52MG	N021225 001	Dec 06, 2000
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SKYLA

+	BAYER HLTHCARE	13.5MG	N203159 001	Jan 09, 2013
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LEVORPHANOL TARTRATE

TABLET; ORAL

LEVORPHANOL TARTRATE

AB		HIKMA	2MG	A074278 001	Mar 31, 2000
AB		NOVITIUM PHARMA	2MG	A213479 001	Jul 01, 2020
AB	!		3MG	A213479 002	Jan 12, 2021
AB		SPECGX LLC	2MG	A212024 001	Dec 13, 2019
AB		SUN PHARM INDS INC	2MG	A213906 001	Jun 17, 2021
AB			3MG	A213906 002	Apr 25, 2023
AB		VIRTUS	2MG	A211484 001	Dec 13, 2018
		NOVITIUM PHARMA	1MG	A213479 003	Jul 20, 2021

LEVOTHYROXINE SODIUM

CAPSULE; ORAL

TIROSINT

+	IBSA	0.013MG	N021924 013	Aug 01, 2007
+		0.025MG	N021924 002	Oct 13, 2006
+		0.0375MG	N021924 014	Jun 22, 2022
+		0.044MG	N021924 015	Jun 22, 2022

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM

CAPSULE; ORAL

TIROSINT

+		0.05MG	N021924	003	Oct 13, 2006
+		0.0625MG	N021924	016	Jun 22, 2022
+		0.075MG	N021924	004	Oct 13, 2006
+		0.088MG	N021924	010	Oct 02, 2009
+		0.1MG	N021924	005	Oct 13, 2006
+		0.112MG	N021924	008	Oct 02, 2009
+		0.125MG	N021924	006	Oct 13, 2006
+		0.137MG	N021924	009	Oct 02, 2009
+		0.15MG	N021924	007	Oct 13, 2006
+		0.175MG	N021924	011	Apr 25, 2017
+	!	0.2MG	N021924	012	Apr 25, 2017

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

<u>AP</u>	+	!	FRESENIUS KABI USA	<u>100MCG/VIAL</u>	<u>N202231</u>	<u>001</u>	Jun 24, 2011
<u>AP</u>	+	!		<u>200MCG/VIAL</u>	<u>N202231</u>	<u>002</u>	Jun 24, 2011
<u>AP</u>	+	!		<u>500MCG/VIAL</u>	<u>N202231</u>	<u>003</u>	Jun 24, 2011
<u>AP</u>			MAIA PHARMS INC	<u>100MCG/VIAL</u>	<u>A208749</u>	<u>001</u>	Dec 21, 2018
<u>AP</u>				<u>200MCG/VIAL</u>	<u>A208749</u>	<u>002</u>	Dec 21, 2018
<u>AP</u>				<u>500MCG/VIAL</u>	<u>A208749</u>	<u>003</u>	Dec 21, 2018
<u>AP</u>			PIRAMAL CRITICAL	<u>100MCG/VIAL</u>	<u>A206163</u>	<u>001</u>	Jun 29, 2016
<u>AP</u>				<u>500MCG/VIAL</u>	<u>A206163</u>	<u>002</u>	Jun 29, 2016
<u>AP</u>			ZYDUS PHARMS	<u>100MCG/VIAL</u>	<u>A217066</u>	<u>001</u>	Mar 24, 2023
<u>AP</u>				<u>200MCG/VIAL</u>	<u>A217066</u>	<u>002</u>	Mar 24, 2023
<u>AP</u>				<u>500MCG/VIAL</u>	<u>A217066</u>	<u>003</u>	Mar 24, 2023

SOLUTION; INTRAVENOUS

LEVOTHYROXINE SODIUM

+	!	FRESENIUS KABI USA	100MCG/5ML (20MCG/ML)	N210632	001	Apr 11, 2019
+	!		200MCG/5ML (40MCG/ML)	N210632	002	Apr 11, 2019
+	!		500MCG/5ML (100MCG/ML)	N210632	003	Apr 11, 2019
+	!	HIKMA	100MCG/ML	N214253	001	May 17, 2021

SOLUTION; ORAL

ERMEZA

+	!	MYLAN	150MCG/5ML	N215809	001	Apr 29, 2022
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THYQUIDITY

+	!	AZURITY	100MCG/5ML	N214047	001	Nov 30, 2020
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TIROSINT-SOL

+		IBSA	13MCG/ML	N206977	001	Dec 15, 2016
+			25MCG/ML	N206977	002	Dec 15, 2016
+			37.5MCG/ML	N206977	013	Jan 13, 2021
+			44MCG/ML	N206977	014	Jan 13, 2021
+			50MCG/ML	N206977	003	Dec 15, 2016
+			62.5MCG/ML	N206977	015	Jan 13, 2021
+			75MCG/ML	N206977	004	Dec 15, 2016
+			88MCG/ML	N206977	005	Dec 15, 2016
+			100MCG/ML	N206977	006	Dec 15, 2016
+			112MCG/ML	N206977	007	Dec 15, 2016
+			125MCG/ML	N206977	008	Dec 15, 2016
+			137MCG/ML	N206977	009	Dec 15, 2016
+			150MCG/ML	N206977	010	Dec 15, 2016
+			175MCG/ML	N206977	011	Dec 15, 2016
+	!		200MCG/ML	N206977	012	Dec 15, 2016

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET; ORAL

SYNTHROID

-->	+	ABBEVIE	--> <u>AB1, AB2</u>	<u>0.025MG</u>	N021402	001	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.05MG</u>	N021402	002	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.075MG</u>	N021402	003	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.088MG</u>	N021402	004	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.1MG</u>	N021402	005	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.112MG</u>	N021402	006	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.125MG</u>	N021402	007	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.137MG</u>	N021402	008	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.15MG</u>	N021402	009	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.175MG</u>	N021402	010	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.2MG</u>	N021402	012	Jul 24, 2002
-->	+		--> <u>AB1, AB2</u>	<u>0.3MG</u>	N021402	011	Jul 24, 2002

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVO-T

-->	CEDIPROF INC	--> <u>AB1, AB2, AB3</u>	<u>0.025MG</u>	N021342 001	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.05MG</u>	N021342 002	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.075MG</u>	N021342 003	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.088MG</u>	N021342 004	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.1MG</u>	N021342 005	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.112MG</u>	N021342 006	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.125MG</u>	N021342 007	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.137MG</u>	N021342 012	Dec 08, 2003
-->		--> <u>AB1, AB2, AB3</u>	<u>0.15MG</u>	N021342 008	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.175MG</u>	N021342 009	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.2MG</u>	N021342 010	Mar 01, 2002
-->		--> <u>AB1, AB2, AB3</u>	<u>0.3MG</u>	N021342 011	Mar 01, 2002

LEVOTHYROXINE SODIUM

-->	LUPIN	--> <u>AB1, AB2, AB3</u>	<u>0.025MG</u>	A209713 001	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.05MG</u>	A209713 002	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.075MG</u>	A209713 003	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.088MG</u>	A209713 004	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.1MG</u>	A209713 005	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.112MG</u>	A209713 006	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.125MG</u>	A209713 007	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.137MG</u>	A209713 008	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.15MG</u>	A209713 009	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.175MG</u>	A209713 010	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.2MG</u>	A209713 011	Jan 18, 2019
-->		--> <u>AB1, AB2, AB3</u>	<u>0.3MG</u>	A209713 012	Jan 18, 2019

UNITHROID

-->	+ STEVENS J	--> <u>AB1, AB2, AB3</u>	<u>0.025MG</u>	N021210 001	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.05MG</u>	N021210 002	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.075MG</u>	N021210 003	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.088MG</u>	N021210 004	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.1MG</u>	N021210 005	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.112MG</u>	N021210 006	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.125MG</u>	N021210 007	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.137MG</u>	N021210 012	Feb 08, 2008
-->		--> <u>AB1, AB2, AB3</u>	<u>0.15MG</u>	N021210 008	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.175MG</u>	N021210 009	Aug 21, 2000
-->		--> <u>AB1, AB2, AB3</u>	<u>0.2MG</u>	N021210 010	Aug 21, 2000
-->	+	--> <u>AB1, AB2, AB3</u>	<u>0.3MG</u>	N021210 011	Aug 21, 2000

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOLET

-->	GENUS LIFESCIENCES	-->	<u>0.025MG</u>	N021137 001 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.05MG</u>	N021137 002 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.075MG</u>	N021137 003 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.088MG</u>	N021137 004 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.1MG</u>	N021137 005 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.112MG</u>	N021137 006 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.125MG</u>	N021137 007 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.137MG</u>	N021137 008 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.15MG</u>	N021137 009 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.175MG</u>	N021137 010 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.2MG</u>	N021137 011 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.3MG</u>	N021137 012 Jun 06, 2003
		<u>AB1, AB2, AB3, AB4</u>		

LEVOTHYROXINE SODIUM

-->	ACCORD HLTHCARE	-->	<u>0.025MG</u>	A212399 001 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.05MG</u>	A212399 002 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.075MG</u>	A212399 003 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.088MG</u>	A212399 004 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.1MG</u>	A212399 005 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.112MG</u>	A212399 006 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.125MG</u>	A212399 007 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.137MG</u>	A212399 008 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.15MG</u>	A212399 009 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.175MG</u>	A212399 010 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.2MG</u>	A212399 011 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.3MG</u>	A212399 012 Oct 19, 2020
		<u>AB1, AB2, AB3, AB4</u>		
-->	MYLAN	-->	<u>0.025MG</u>	A076187 001 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.05MG</u>	A076187 002 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.075MG</u>	A076187 003 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.088MG</u>	A076187 004 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.1MG</u>	A076187 005 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.112MG</u>	A076187 006 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.125MG</u>	A076187 007 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.137MG</u>	A076187 012 Dec 13, 2006
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.15MG</u>	A076187 008 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.175MG</u>	A076187 009 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->		-->	<u>0.2MG</u>	A076187 010 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		
-->	!	-->	<u>0.3MG</u>	A076187 011 Jun 05, 2002
		<u>AB1, AB2, AB3, AB4</u>		

THYRO-TABS

-->	+ ALVOGEN	-->	<u>0.025MG</u>	N021116 001 Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>		

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

THYRO-TABS

-->	+	-->	<u>0.05MG</u>	N021116	002	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.075MG</u>	N021116	003	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.088MG</u>	N021116	010	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.1MG</u>	N021116	004	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.112MG</u>	N021116	011	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.125MG</u>	N021116	005	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.137MG</u>	N021116	012	Dec 07, 2004
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.15MG</u>	N021116	006	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.175MG</u>	N021116	007	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.2MG</u>	N021116	008	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				
-->	+	-->	<u>0.3MG</u>	N021116	009	Oct 24, 2002
		<u>AB1, AB2, AB3, AB4</u>				

LEVOXYL

-->	+	KING PHARMS	-->	<u>AB1, AB3</u>	<u>0.025MG</u>	N021301	001	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.05MG</u>	N021301	002	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.075MG</u>	N021301	003	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.088MG</u>	N021301	004	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.1MG</u>	N021301	005	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.112MG</u>	N021301	006	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.125MG</u>	N021301	007	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.137MG</u>	N021301	008	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.15MG</u>	N021301	009	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.175MG</u>	N021301	010	May 25, 2001
-->	+		-->	<u>AB1, AB3</u>	<u>0.2MG</u>	N021301	011	May 25, 2001

EUTHYROX

<u>AB2</u>	PROVELL	<u>0.025MG</u>	<u>N021292</u>	<u>001</u>	May 31, 2002
<u>AB2</u>		<u>0.05MG</u>	<u>N021292</u>	<u>002</u>	May 31, 2002
<u>AB2</u>		<u>0.075MG</u>	<u>N021292</u>	<u>003</u>	May 31, 2002
<u>AB2</u>		<u>0.088MG</u>	<u>N021292</u>	<u>004</u>	May 31, 2002
<u>AB2</u>		<u>0.1MG</u>	<u>N021292</u>	<u>005</u>	May 31, 2002
<u>AB2</u>		<u>0.112MG</u>	<u>N021292</u>	<u>006</u>	May 31, 2002
<u>AB2</u>		<u>0.125MG</u>	<u>N021292</u>	<u>007</u>	May 31, 2002
<u>AB2</u>		<u>0.137MG</u>	<u>N021292</u>	<u>008</u>	May 31, 2002
<u>AB2</u>		<u>0.15MG</u>	<u>N021292</u>	<u>009</u>	May 31, 2002
<u>AB2</u>		<u>0.175MG</u>	<u>N021292</u>	<u>010</u>	May 31, 2002
<u>AB2</u>		<u>0.2MG</u>	<u>N021292</u>	<u>011</u>	May 31, 2002

LEVOTHYROXINE SODIUM

<u>AB2</u>	MACLEODS PHARMS LTD	<u>0.025MG</u>	<u>A211417</u>	<u>001</u>	Dec 21, 2022
<u>AB2</u>		<u>0.05MG</u>	<u>A211417</u>	<u>002</u>	Dec 21, 2022
<u>AB2</u>		<u>0.075MG</u>	<u>A211417</u>	<u>003</u>	Dec 21, 2022
<u>AB2</u>		<u>0.088MG</u>	<u>A211417</u>	<u>004</u>	Dec 21, 2022
<u>AB2</u>		<u>0.1MG</u>	<u>A211417</u>	<u>005</u>	Dec 21, 2022
<u>AB2</u>		<u>0.112MG</u>	<u>A211417</u>	<u>006</u>	Dec 21, 2022
<u>AB2</u>		<u>0.125MG</u>	<u>A211417</u>	<u>007</u>	Dec 21, 2022
<u>AB2</u>		<u>0.137MG</u>	<u>A211417</u>	<u>008</u>	Dec 21, 2022
<u>AB2</u>		<u>0.15MG</u>	<u>A211417</u>	<u>009</u>	Dec 21, 2022
<u>AB2</u>		<u>0.175MG</u>	<u>A211417</u>	<u>010</u>	Dec 21, 2022
<u>AB2</u>		<u>0.2MG</u>	<u>A211417</u>	<u>011</u>	Dec 21, 2022
<u>AB2</u>		<u>0.3MG</u>	<u>A211417</u>	<u>012</u>	Dec 21, 2022
<u>AB2</u>	WATSON LABS TEVA	<u>0.025MG</u>	<u>A207588</u>	<u>001</u>	May 10, 2022
<u>AB2</u>		<u>0.05MG</u>	<u>A207588</u>	<u>002</u>	May 10, 2022
<u>AB2</u>		<u>0.075MG</u>	<u>A207588</u>	<u>003</u>	May 10, 2022
<u>AB2</u>		<u>0.088MG</u>	<u>A207588</u>	<u>004</u>	May 10, 2022
<u>AB2</u>		<u>0.1MG</u>	<u>A207588</u>	<u>005</u>	May 10, 2022
<u>AB2</u>		<u>0.112MG</u>	<u>A207588</u>	<u>006</u>	May 10, 2022
<u>AB2</u>		<u>0.125MG</u>	<u>A207588</u>	<u>007</u>	May 10, 2022
<u>AB2</u>		<u>0.137MG</u>	<u>A207588</u>	<u>008</u>	May 10, 2022
<u>AB2</u>		<u>0.15MG</u>	<u>A207588</u>	<u>009</u>	May 10, 2022
<u>AB2</u>		<u>0.175MG</u>	<u>A207588</u>	<u>010</u>	May 10, 2022
<u>AB2</u>		<u>0.2MG</u>	<u>A207588</u>	<u>011</u>	May 10, 2022
<u>AB2</u>		<u>0.3MG</u>	<u>A207588</u>	<u>012</u>	May 10, 2022

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOTHYROXINE SODIUM

<u>AB4</u>	ASCENT PHARMS INC	<u>0.025MG</u>	<u>A215259 001</u>	Jan 18, 2023
<u>AB4</u>		<u>0.05MG</u>	<u>A215259 002</u>	Jan 18, 2023
<u>AB4</u>		<u>0.075MG</u>	<u>A215259 003</u>	Jan 18, 2023
<u>AB4</u>		<u>0.088MG</u>	<u>A215259 004</u>	Jan 18, 2023
<u>AB4</u>		<u>0.1MG</u>	<u>A215259 005</u>	Jan 18, 2023
<u>AB4</u>		<u>0.112MG</u>	<u>A215259 006</u>	Jan 18, 2023
<u>AB4</u>		<u>0.125MG</u>	<u>A215259 007</u>	Jan 18, 2023
<u>AB4</u>		<u>0.137MG</u>	<u>A215259 008</u>	Jan 18, 2023
<u>AB4</u>		<u>0.15MG</u>	<u>A215259 009</u>	Jan 18, 2023
<u>AB4</u>		<u>0.175MG</u>	<u>A215259 010</u>	Jan 18, 2023
<u>AB4</u>		<u>0.2MG</u>	<u>A215259 011</u>	Jan 18, 2023
<u>AB4</u>		<u>0.3MG</u>	<u>A215259 012</u>	Jan 18, 2023

LIDOCAINE

OINTMENT;TOPICAL

LIDOCAINE

<u>AT</u>	ALEMBIC	<u>5%</u>	<u>A211469 001</u>	Nov 23, 2018
<u>AT</u>	ALKEM LABS LTD	<u>5%</u>	<u>A207810 001</u>	Mar 10, 2017
<u>AT</u>	AMNEAL PHARMS	<u>5%</u>	<u>A206297 001</u>	Aug 07, 2015
<u>AT</u>	COSETTE	<u>5%</u>	<u>A211019 001</u>	Dec 12, 2018
<u>AT</u>	DR REDDYS	<u>5%</u>	<u>A208660 001</u>	Jan 05, 2021
<u>AT</u>	+! FOUGERA PHARMS INC	<u>5%</u>	<u>A080198 001</u>	
<u>AT</u>	GLENMARK PHARMS LTD	<u>5%</u>	<u>A206498 001</u>	Sep 09, 2016
<u>AT</u>	MACLEODS PHARMS LTD	<u>5%</u>	<u>A211697 001</u>	Mar 16, 2020
<u>AT</u>	QUAGEN	<u>5%</u>	<u>A212695 001</u>	Apr 20, 2021
<u>AT</u>	SEPTODONT INC	<u>5%</u>	<u>A040911 001</u>	May 23, 2011
<u>AT</u>	STRIDES PHARMA	<u>5%</u>	<u>A210958 001</u>	Dec 11, 2018
<u>AT</u>	TARO	<u>5%</u>	<u>A086724 001</u>	

PATCH;TOPICAL

LIDOCAINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>5%</u>	<u>A200675 001</u>	Aug 23, 2012
<u>AB</u>	AMNEAL	<u>5%</u>	<u>A206463 001</u>	Aug 24, 2020
<u>AB</u>	IBSA	<u>5%</u>	<u>A209190 001</u>	Apr 30, 2020
<u>AB</u>	MYLAN TECHNOLOGIES	<u>5%</u>	<u>A202346 001</u>	Aug 07, 2015
<u>AB</u>	NAL PHARM	<u>5%</u>	<u>A205882 001</u>	Apr 29, 2021

LIDODERM

<u>AB</u>	+! TEIKOKU PHARMA USA	<u>5%</u>	<u>N020612 001</u>	Mar 19, 1999
	ZTLIDO			
	+! SCILEX PHARMS	1.8%	N207962 001	Feb 28, 2018

LIDOCAINE HYDROCHLORIDE

GEL;OPHTHALMIC

AKTEN

	+! THEA PHARMA	3.5%	N022221 001	Oct 07, 2008
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INJECTABLE;INJECTION

LIDOCAINE HYDROCHLORIDE

<u>AP</u>	AFAXYS	<u>1%</u>	<u>A215132 001</u>	Jun 09, 2022
<u>AP</u>	ASPIRO	<u>1%</u>	<u>A214336 001</u>	Nov 08, 2021
<u>AP</u>		<u>1%</u>	<u>A214339 001</u>	Nov 08, 2021
<u>AP</u>		<u>2%</u>	<u>A214336 002</u>	Dec 13, 2023
<u>AP</u>		<u>2%</u>	<u>A214339 002</u>	Nov 08, 2021
<u>AP</u>	B BRAUN MEDICAL INC	<u>1%</u>	<u>A208474 001</u>	Aug 03, 2018
<u>AP</u>	EUGIA PHARMA	<u>1%</u>	<u>A207182 001</u>	Oct 30, 2017
<u>AP</u>		<u>2%</u>	<u>A207182 002</u>	Oct 30, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A080404 002</u>	
<u>AP</u>		<u>2%</u>	<u>A080404 003</u>	
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088328 001</u>	May 17, 1984
<u>AP</u>	+!	<u>1%</u>	<u>A083158 001</u>	
<u>AP</u>		<u>1%</u>	<u>A088329 001</u>	May 17, 1984
<u>AP</u>		<u>2%</u>	<u>A040078 001</u>	Jun 23, 1995
<u>AP</u>	+!	<u>2%</u>	<u>A083158 002</u>	
<u>AP</u>		<u>2%</u>	<u>A088294 001</u>	May 17, 1984
<u>AP</u>	HUONS	<u>1%</u>	<u>A212821 001</u>	May 07, 2020
<u>AP</u>		<u>2%</u>	<u>A212821 002</u>	Jun 15, 2023
<u>AP</u>	INTL MEDICATION	<u>1%</u>	<u>A083173 001</u>	
<u>AP</u>		<u>2%</u>	<u>A083173 002</u>	
<u>AP</u>	MANKIND PHARMA	<u>1%</u>	<u>A217692 001</u>	Jun 16, 2023
<u>AP</u>		<u>1%</u>	<u>A217692 003</u>	Nov 29, 2023
<u>AP</u>		<u>1%</u>	<u>A217693 001</u>	Jul 11, 2023

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

<u>AP</u>		<u>2%</u>	<u>A217692 002</u>	Jun 16, 2023
<u>AP</u>		<u>2%</u>	<u>A217693 002</u>	Jul 11, 2023
<u>AP</u>	SINETICA US	<u>1%</u>	<u>A214267 001</u>	Sep 19, 2022
<u>AP</u>		<u>2%</u>	<u>A214267 002</u>	Sep 19, 2022
<u>AP</u>		<u>4%</u>	<u>A214269 001</u>	May 05, 2022
<u>AP</u>	SPECTRA MDCL DEVICES	<u>1%</u>	<u>A208017 001</u>	Apr 18, 2018
<u>AP</u>	WEST-WARD PHARMS INT	<u>1%</u>	<u>A080407 001</u>	
<u>AP</u>		<u>2%</u>	<u>A080407 002</u>	
	<u>LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	B BRAUN	<u>200MG/100ML</u>	<u>N019830 002</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML</u>	<u>N018461 002</u>	
	<u>LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	B BRAUN	<u>400MG/100ML</u>	<u>N019830 003</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>400MG/100ML</u>	<u>N018461 003</u>	
	<u>LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	B BRAUN	<u>800MG/100ML</u>	<u>N019830 004</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>800MG/100ML</u>	<u>N018461 004</u>	Feb 22, 1982
	<u>LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER</u>			
<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A088586 001</u>	Jul 24, 1985
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088325 001</u>	Jul 31, 1984
<u>AP</u>		<u>1%</u>	<u>A088299 001</u>	Jul 31, 1984
<u>AP</u>		<u>2%</u>	<u>A088327 001</u>	Jul 31, 1984
	<u>LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE</u>			
<u>AP</u>	EUGIA PHARMA	<u>1%</u>	<u>A203040 001</u>	Mar 14, 2013
<u>AP</u>		<u>1%</u>	<u>A203082 001</u>	Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>A203040 002</u>	Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>A203082 002</u>	Mar 14, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>2%</u>	<u>N017584 001</u>	
<u>AP</u>		<u>4%</u>	<u>N017584 002</u>	
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A080408 001</u>	
<u>AP</u>		<u>1.5%</u>	<u>A080408 002</u>	
<u>AP</u>	!	<u>4%</u>	<u>A088295 001</u>	May 17, 1984
<u>AP</u>	WEST-WARD PHARMS INT	<u>1%</u>	<u>A084625 001</u>	
<u>AP</u>		<u>2%</u>	<u>A084625 002</u>	
	<u>LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER</u>			
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A040302 001</u>	Sep 28, 1998
<u>AP</u>		<u>2%</u>	<u>A040302 002</u>	Sep 28, 1998
	<u>XYLOCAINE</u>			
<u>AP</u>	+! FRESENIUS KABI USA	<u>0.5%</u>	<u>N006488 008</u>	
<u>AP</u>	+!	<u>1%</u>	<u>N006488 007</u>	
<u>AP</u>	+!	<u>1.5%</u>	<u>N006488 010</u>	
<u>AP</u>	+!	<u>2%</u>	<u>N006488 002</u>	
	INJECTABLE; SPINAL			
	LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%			
	+! HOSPIRA	5%	A083914	001
	JELLY; TOPICAL			
	<u>GLYDO</u>			
<u>AT</u>	! SAGENT PHARMS INC	<u>2%</u>	<u>A201094 001</u>	Apr 28, 2014
	<u>LIDOCAINE HYDROCHLORIDE</u>			
<u>AT</u>	INTL MEDICATION	<u>2%</u>	<u>A086283 001</u>	
<u>AT</u>	SENTISS	<u>2%</u>	<u>A040433 001</u>	Feb 12, 2003
	SOLUTION; ORAL			
	<u>LIDOCAINE HYDROCHLORIDE</u>			
<u>AT</u>	! RUBICON	<u>2%</u>	<u>A216780 001</u>	Mar 28, 2023
<u>AT</u>	WOCKHARDT BIO AG	<u>2%</u>	<u>A087872 001</u>	Nov 18, 1982
	<u>LIDOCAINE HYDROCHLORIDE VISCOUS</u>			
<u>AT</u>	CHARTWELL MOLECULAR	<u>2%</u>	<u>A040708 001</u>	Feb 27, 2007
	<u>LIDOCAINE VISCOUS</u>			
<u>AT</u>	HIKMA	<u>2%</u>	<u>A088802 001</u>	Apr 26, 1985
	SOLUTION; TOPICAL			
	<u>LARYNG-O-JET KIT</u>			
<u>AT</u>	INTL MEDICATION	<u>4%</u>	<u>A086364 001</u>	
	<u>LIDOCAINE HYDROCHLORIDE</u>			
<u>AT</u>	! HIKMA	<u>4%</u>	<u>A088803 001</u>	Apr 03, 1985
<u>AT</u>	LANNETT CO INC	<u>4%</u>	<u>A040710 001</u>	Feb 27, 2007
<u>AT</u>	NOVITIUM PHARMA	<u>4%</u>	<u>A216250 001</u>	Mar 23, 2022
<u>AT</u>	PAI HOLDINGS PHARM	<u>4%</u>	<u>A204494 001</u>	Mar 12, 2014

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE HYDROCHLORIDE

SOLUTION; TOPICAL

LIDOCAINE HYDROCHLORIDE

AT	TARO	4%	A218182 001	Dec 07, 2023
AT	WOCKHARDT BIO AG	4%	A087881 001	Nov 18, 1982
	SYSTEM; INTRADERMAL			
	ZINGO			
	POWDER PHARMS	0.5MG	N022114 001	Aug 16, 2007

LIDOCAINE; PRILOCAINE

CREAM; TOPICAL

LIDOCAINE AND PRILOCAINE

AB	ALEMBIC	2.5%;2.5%	A213923 001	Apr 08, 2022
AB	ENCUBE	2.5%;2.5%	A076320 001	Aug 27, 2003
AB	! FOUGERA PHARMS	2.5%;2.5%	A076453 001	Aug 18, 2003
AB	HIKMA	2.5%;2.5%	A076290 001	Sep 25, 2003
AB	PADAGIS US	2.5%;2.5%	A212482 001	Jul 27, 2021
	GEL; PERIODONTAL			
	ORAQIX			
	+! DENTSPLY PHARM	2.5%;2.5%	N021451 001	Dec 19, 2003

LIFITEGRAST

SOLUTION/DROPS; OPHTHALMIC

LIFITEGRAST

AB	EUGIA PHARMA	5%	A215063 001	Nov 07, 2023
AB	MICRO LABS	5%	A215081 001	Aug 04, 2023
	<u>XIIDRA</u>			
AB	+! NOVARTIS	5%	N208073 001	Jul 11, 2016

LINACLOTIDE

CAPSULE; ORAL

LINZESS

	+! ABBVIE	72MCG	N202811 003	Jan 25, 2017
	+!	145MCG	N202811 001	Aug 30, 2012
	+	290MCG	N202811 002	Aug 30, 2012

LINAGLIPTIN

TABLET; ORAL

LINAGLIPTIN

AB	SUNSHINE	5MG	A208335 001	Aug 31, 2021
	<u>TRADJENTA</u>			
AB	+! BOEHRINGER INGELHEIM	5MG	N201280 001	May 02, 2011

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

JENTADUETO

AB	+ BOEHRINGER INGELHEIM	2.5MG;500MG	N201281 001	Jan 30, 2012
AB	+	2.5MG;850MG	N201281 002	Jan 30, 2012
AB	+!	2.5MG;1GM	N201281 003	Jan 30, 2012

LINAGLIPTIN AND METFORMIN HYDROCHLORIDE

AB	SUNSHINE	2.5MG;500MG	A208336 001	Aug 30, 2021
AB		2.5MG;850MG	A208336 002	Aug 30, 2021
AB		2.5MG;1GM	A208336 003	Aug 30, 2021

TABLET, EXTENDED RELEASE; ORAL

JENTADUETO XR

	+ BOEHRINGER INGELHEIM	2.5MG;1GM	N208026 001	May 27, 2016
	+!	5MG;1GM	N208026 002	May 27, 2016

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOCIN

AP	+! PFIZER	EQ 300MG BASE/ML	N050317 001	
	<u>LINCOMYCIN HYDROCHLORIDE</u>			
AP	GLAND PHARMA LTD	EQ 300MG BASE/ML	A215657 001	Sep 26, 2022
AP	MICRO LABS	EQ 300MG BASE/ML	A215082 001	Nov 08, 2021
AP	XGEN PHARMS	EQ 300MG BASE/ML	A201746 001	Jun 04, 2015

PRESCRIPTION DRUG PRODUCT LIST

LINEZOLID

FOR SUSPENSION; ORAL

LINEZOLID

AB	HETERO LABS	<u>100MG/5ML</u>	<u>A211813</u>	<u>001</u>	Oct 31, 2022
AB	HIKMA	<u>100MG/5ML</u>	<u>A200068</u>	<u>001</u>	Jun 03, 2015

ZYVOX

AB	+ ! PFIZER	<u>100MG/5ML</u>	<u>N021132</u>	<u>001</u>	Apr 18, 2000
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SOLUTION; INTRAVENOUS

LINEZOLID

AP	EUGIA PHARMA	<u>600MG/300ML (2MG/ML)</u>	<u>A206917</u>	<u>001</u>	Aug 04, 2016
AP	FRESENIUS KABI USA	<u>600MG/300ML (2MG/ML)</u>	<u>A204764</u>	<u>001</u>	Mar 15, 2016
AP	HIKMA	<u>600MG/300ML (2MG/ML)</u>	<u>A206454</u>	<u>001</u>	Aug 22, 2022
AP	HQ SPCLT PHARMA	<u>200MG/100ML (2MG/ML)</u>	<u>A207001</u>	<u>001</u>	Jul 07, 2017
AP		<u>600MG/300ML (2MG/ML)</u>	<u>A207001</u>	<u>002</u>	Jul 07, 2017
AP	MYLAN LABS LTD	<u>200MG/100ML (2MG/ML)</u>	<u>A205154</u>	<u>001</u>	Dec 06, 2017
AP		<u>600MG/300ML (2MG/ML)</u>	<u>A205154</u>	<u>002</u>	Dec 06, 2017
AP	NANG KUANG PHARM CO	<u>200MG/100ML (2MG/ML)</u>	<u>A207354</u>	<u>001</u>	Dec 20, 2016
AP		<u>600MG/300ML (2MG/ML)</u>	<u>A207354</u>	<u>002</u>	Dec 20, 2016
AP	SAGENT PHARMS INC	<u>200MG/100ML (2MG/ML)</u>	<u>A204696</u>	<u>001</u>	Mar 02, 2017
AP		<u>600MG/300ML (2MG/ML)</u>	<u>A204696</u>	<u>002</u>	Mar 02, 2017
AP	SANDOZ	<u>200MG/100ML (2MG/ML)</u>	<u>A200904</u>	<u>001</u>	Jul 16, 2015
AP		<u>600MG/300ML (2MG/ML)</u>	<u>A200904</u>	<u>002</u>	Jul 16, 2015

ZYVOX

AP	+	PFIZER	<u>200MG/100ML (2MG/ML)</u>	<u>N021131</u>	<u>001</u>	Apr 18, 2000
AP	+ !		<u>600MG/300ML (2MG/ML)</u>	<u>N021131</u>	<u>003</u>	Apr 18, 2000

LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+ !	HOSPIRA	600MG/300ML (2MG/ML)	N206473	001	Jun 18, 2015
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TABLET; ORAL

LINEZOLID

AB	ALEMBIC	<u>600MG</u>	<u>A205233</u>	<u>001</u>	Dec 21, 2015
AB	ALKEM LABS LTD	<u>600MG</u>	<u>A205517</u>	<u>001</u>	Dec 21, 2015
AB	CHARTWELL RX	<u>600MG</u>	<u>A210702</u>	<u>001</u>	Apr 25, 2019
AB	GLENMARK PHARMS	<u>600MG</u>	<u>A078987</u>	<u>001</u>	Dec 21, 2015
AB	HETERO LABS LTD V	<u>600MG</u>	<u>A204239</u>	<u>001</u>	Dec 21, 2015
AB	NOVEL LABS INC	<u>600MG</u>	<u>A207526</u>	<u>001</u>	Aug 22, 2016
AB	RISING	<u>600MG</u>	<u>A078845</u>	<u>001</u>	Dec 21, 2015
AB	ZYDUS PHARMS	<u>600MG</u>	<u>A206097</u>	<u>001</u>	Feb 22, 2017

ZYVOX

AB	+ !	PFIZER	<u>600MG</u>	<u>N021130</u>	<u>002</u>	Apr 18, 2000
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LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

LIOTHYRONINE SODIUM

!	XGEN PHARMS	EQ 0.01MG BASE/ML	A076923	001	Aug 17, 2005
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TABLET; ORAL

CYTOMEL

AB	+	KING PHARMS	<u>EQ 0.005MG BASE</u>	<u>N010379</u>	<u>001</u>
AB	+		<u>EQ 0.025MG BASE</u>	<u>N010379</u>	<u>002</u>
AB	+		<u>EQ 0.05MG BASE</u>	<u>N010379</u>	<u>003</u>

LIOTHYRONINE SODIUM

AB	DR REDDYS LABS SA	<u>EQ 0.005MG BASE</u>	<u>A090097</u>	<u>001</u>	Mar 20, 2009
AB		<u>EQ 0.025MG BASE</u>	<u>A090097</u>	<u>002</u>	Mar 20, 2009
AB		<u>EQ 0.05MG BASE</u>	<u>A090097</u>	<u>003</u>	Mar 20, 2009
AB	SIGMAPHARM LABS LLC	<u>EQ 0.005MG BASE</u>	<u>A200295</u>	<u>001</u>	Nov 29, 2012
AB		<u>EQ 0.025MG BASE</u>	<u>A200295</u>	<u>002</u>	Nov 29, 2012
AB	!	<u>EQ 0.05MG BASE</u>	<u>A200295</u>	<u>003</u>	Nov 29, 2012
AB	SUN PHARM	<u>EQ 0.005MG BASE</u>	<u>A091382</u>	<u>001</u>	Apr 20, 2016
AB		<u>EQ 0.025MG BASE</u>	<u>A091382</u>	<u>002</u>	Apr 20, 2016
AB		<u>EQ 0.05MG BASE</u>	<u>A091382</u>	<u>003</u>	Apr 20, 2016
AB	TEVA PHARMS USA	<u>EQ 0.005MG BASE</u>	<u>A211510</u>	<u>001</u>	Oct 26, 2018
AB		<u>EQ 0.025MG BASE</u>	<u>A211510</u>	<u>002</u>	Oct 26, 2018
AB		<u>EQ 0.05MG BASE</u>	<u>A211510</u>	<u>003</u>	Oct 26, 2018
AB	ZYDUS	<u>EQ 0.005MG BASE</u>	<u>A214803</u>	<u>001</u>	Jan 22, 2021
AB		<u>EQ 0.025MG BASE</u>	<u>A214803</u>	<u>002</u>	Jan 22, 2021
AB		<u>EQ 0.05MG BASE</u>	<u>A214803</u>	<u>003</u>	Jan 22, 2021

LIRAGLUTIDE RECOMBINANT

SOLUTION; SUBCUTANEOUS

SAXENDA

+ !	NOVO	18MG/3ML (6MG/ML)	N206321	001	Dec 23, 2014
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VICTOZA

+ !	NOVO NORDISK INC	18MG/3ML (6MG/ML)	N022341	001	Jan 25, 2010
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PRESCRIPTION DRUG PRODUCT LIST

LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

LISDEXAMFETAMINE DIMESYLATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A202802 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A202802 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202802 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202802 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202802 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202802 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202802 007</u>	Aug 25, 2023
<u>AB</u>	ALKEM LABS LTD	<u>10MG</u>	<u>A217194 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A217194 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A217194 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A217194 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A217194 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A217194 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A217194 007</u>	Aug 25, 2023
<u>AB</u>	AMNEAL	<u>20MG</u>	<u>A202830 001</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202830 002</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202830 003</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202830 004</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202830 005</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202830 006</u>	Aug 25, 2023
<u>AB</u>	APOTEX	<u>10MG</u>	<u>A216944 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A216944 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A216944 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A216944 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A216944 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A216944 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A216944 007</u>	Aug 25, 2023
<u>AB</u>	ASCENT PHARMS INC	<u>10MG</u>	<u>A217442 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A217442 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A217442 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A217442 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A217442 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A217442 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A217442 007</u>	Aug 25, 2023
<u>AB</u>	HIKMA	<u>20MG</u>	<u>A202827 001</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202827 002</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202827 003</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202827 004</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202827 005</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202827 006</u>	Aug 25, 2023
<u>AB</u>	LANNETT CO INC	<u>10MG</u>	<u>A215802 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A215802 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A215802 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A215802 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A215802 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A215802 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A215802 007</u>	Aug 25, 2023
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A202835 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A202835 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A202835 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A202835 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A202835 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A202835 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A202835 007</u>	Aug 25, 2023
<u>AB</u>	NORWICH	<u>10MG</u>	<u>A214547 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A214547 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A214547 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A214547 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A214547 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A214547 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A214547 007</u>	Aug 25, 2023
<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A216266 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A216266 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A216266 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A216266 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A216266 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A216266 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A216266 007</u>	Aug 25, 2023
<u>AB</u>	RHODES PHARMS	<u>10MG</u>	<u>A215330 001</u>	Aug 25, 2023

PRESCRIPTION DRUG PRODUCT LIST

LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

LISDEXAMFETAMINE DIMESYLATE

<u>AB</u>		<u>20MG</u>	<u>A215330 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A215330 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A215330 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A215330 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A215330 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A215330 007</u>	Aug 25, 2023
<u>AB</u>	SPECGX LLC	<u>10MG</u>	<u>A211840 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A211840 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A211840 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A211840 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A211840 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A211840 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A211840 007</u>	Aug 25, 2023
<u>AB</u>	SUN PHARM INDS INC	<u>10MG</u>	<u>A214484 001</u>	Aug 25, 2023
<u>AB</u>		<u>20MG</u>	<u>A214484 002</u>	Aug 25, 2023
<u>AB</u>		<u>30MG</u>	<u>A214484 003</u>	Aug 25, 2023
<u>AB</u>		<u>40MG</u>	<u>A214484 004</u>	Aug 25, 2023
<u>AB</u>		<u>50MG</u>	<u>A214484 005</u>	Aug 25, 2023
<u>AB</u>		<u>60MG</u>	<u>A214484 006</u>	Aug 25, 2023
<u>AB</u>		<u>70MG</u>	<u>A214484 007</u>	Aug 25, 2023

VYVANSE

<u>AB</u>	+	TAKEDA PHARMS USA	<u>10MG</u>	<u>N021977 007</u>	Oct 30, 2014
<u>AB</u>	+		<u>20MG</u>	<u>N021977 004</u>	Dec 10, 2007
<u>AB</u>	+		<u>30MG</u>	<u>N021977 001</u>	Feb 23, 2007
<u>AB</u>	+		<u>40MG</u>	<u>N021977 005</u>	Dec 10, 2007
<u>AB</u>	+		<u>50MG</u>	<u>N021977 002</u>	Feb 23, 2007
<u>AB</u>	+		<u>60MG</u>	<u>N021977 006</u>	Dec 10, 2007
<u>AB</u>	+		<u>70MG</u>	<u>N021977 003</u>	Feb 23, 2007

TABLET, CHEWABLE; ORAL

LISDEXAMFETAMINE DIMESYLATE

<u>AB</u>		ASCENT PHARMS INC	<u>10MG</u>	<u>A217068 001</u>	Aug 25, 2023
<u>AB</u>			<u>20MG</u>	<u>A217068 002</u>	Aug 25, 2023
<u>AB</u>			<u>30MG</u>	<u>A217068 003</u>	Aug 25, 2023
<u>AB</u>			<u>40MG</u>	<u>A217068 004</u>	Aug 25, 2023
<u>AB</u>			<u>50MG</u>	<u>A217068 005</u>	Aug 25, 2023
<u>AB</u>			<u>60MG</u>	<u>A217068 006</u>	Aug 25, 2023
<u>AB</u>		SUN PHARM INDS INC	<u>10MG</u>	<u>A214134 001</u>	Aug 25, 2023
<u>AB</u>			<u>20MG</u>	<u>A214134 002</u>	Aug 25, 2023
<u>AB</u>			<u>30MG</u>	<u>A214134 003</u>	Aug 25, 2023
<u>AB</u>			<u>40MG</u>	<u>A214134 004</u>	Aug 25, 2023
<u>AB</u>			<u>50MG</u>	<u>A214134 005</u>	Aug 25, 2023
<u>AB</u>			<u>60MG</u>	<u>A214134 006</u>	Aug 25, 2023
<u>AB</u>		TEVA PHARMS	<u>10MG</u>	<u>A215415 001</u>	Aug 25, 2023
<u>AB</u>			<u>20MG</u>	<u>A215415 002</u>	Aug 25, 2023
<u>AB</u>			<u>30MG</u>	<u>A215415 003</u>	Aug 25, 2023
<u>AB</u>			<u>40MG</u>	<u>A215415 004</u>	Aug 25, 2023
<u>AB</u>			<u>50MG</u>	<u>A215415 005</u>	Aug 25, 2023
<u>AB</u>			<u>60MG</u>	<u>A215415 006</u>	Aug 25, 2023

VYVANSE

<u>AB</u>	+	TAKEDA PHARMS USA	<u>10MG</u>	<u>N208510 001</u>	Jan 28, 2017
<u>AB</u>	+		<u>20MG</u>	<u>N208510 002</u>	Jan 28, 2017
<u>AB</u>	+		<u>30MG</u>	<u>N208510 003</u>	Jan 28, 2017
<u>AB</u>	+		<u>40MG</u>	<u>N208510 004</u>	Jan 28, 2017
<u>AB</u>	+		<u>50MG</u>	<u>N208510 005</u>	Jan 28, 2017
<u>AB</u>	+		<u>60MG</u>	<u>N208510 006</u>	Jan 28, 2017

LISINAPRIL

SOLUTION; ORAL

QBRELIS

+	AZURITY	1MG/ML	N208401 001	Jul 29, 2016
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TABLET; ORAL

LISINAPRIL

<u>AB</u>		ACCORD HLTHCARE	<u>2.5MG</u>	<u>A202554 001</u>	Jul 30, 2013
<u>AB</u>			<u>5MG</u>	<u>A202554 002</u>	Jul 30, 2013
<u>AB</u>			<u>10MG</u>	<u>A202554 003</u>	Jul 30, 2013
<u>AB</u>			<u>20MG</u>	<u>A202554 004</u>	Jul 30, 2013
<u>AB</u>			<u>30MG</u>	<u>A202554 005</u>	Jul 30, 2013
<u>AB</u>			<u>40MG</u>	<u>A202554 006</u>	Jul 30, 2013
<u>AB</u>		ASCENT PHARMS INC	<u>2.5MG</u>	<u>A075903 001</u>	Jul 01, 2002
<u>AB</u>			<u>5MG</u>	<u>A075903 002</u>	Jul 01, 2002

PRESCRIPTION DRUG PRODUCT LIST

LISINOPRIL

TABLET; ORAL

LISINOPRIL

<u>AB</u>		<u>10MG</u>	<u>A075903 003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075903 004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075903 005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075903 006</u>	Jul 01, 2002
<u>AB</u>	AUROBINDO	<u>2.5MG</u>	<u>A077622 001</u>	Feb 22, 2006
<u>AB</u>		<u>5MG</u>	<u>A077622 002</u>	Feb 22, 2006
<u>AB</u>		<u>10MG</u>	<u>A077622 003</u>	Feb 22, 2006
<u>AB</u>		<u>20MG</u>	<u>A077622 004</u>	Feb 22, 2006
<u>AB</u>		<u>30MG</u>	<u>A077622 005</u>	Feb 22, 2006
<u>AB</u>		<u>40MG</u>	<u>A077622 006</u>	Feb 22, 2006
<u>AB</u>	CHARTWELL RX	<u>2.5MG</u>	<u>A075994 001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075994 002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075994 003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075994 004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075994 005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075994 006</u>	Jul 01, 2002
<u>AB</u>	COREPHARMA	<u>2.5MG</u>	<u>A076102 001</u>	Sep 30, 2002
<u>AB</u>		<u>5MG</u>	<u>A076102 002</u>	Sep 30, 2002
<u>AB</u>		<u>10MG</u>	<u>A076102 003</u>	Sep 30, 2002
<u>AB</u>		<u>20MG</u>	<u>A076102 004</u>	Sep 30, 2002
<u>AB</u>		<u>30MG</u>	<u>A076102 005</u>	Sep 30, 2002
<u>AB</u>		<u>40MG</u>	<u>A076102 006</u>	Sep 30, 2002
<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A203508 001</u>	Oct 29, 2013
<u>AB</u>		<u>5MG</u>	<u>A203508 002</u>	Oct 29, 2013
<u>AB</u>		<u>10MG</u>	<u>A203508 003</u>	Oct 29, 2013
<u>AB</u>		<u>20MG</u>	<u>A203508 004</u>	Oct 29, 2013
<u>AB</u>		<u>30MG</u>	<u>A203508 005</u>	Oct 29, 2013
<u>AB</u>		<u>40MG</u>	<u>A203508 006</u>	Oct 29, 2013
<u>AB</u>	LUPIN	<u>2.5MG</u>	<u>A077321 001</u>	Sep 09, 2005
<u>AB</u>		<u>5MG</u>	<u>A077321 002</u>	Sep 09, 2005
<u>AB</u>		<u>10MG</u>	<u>A077321 003</u>	Sep 09, 2005
<u>AB</u>		<u>20MG</u>	<u>A077321 004</u>	Sep 09, 2005
<u>AB</u>		<u>30MG</u>	<u>A077321 005</u>	Sep 09, 2005
<u>AB</u>		<u>40MG</u>	<u>A077321 006</u>	Sep 09, 2005
<u>AB</u>	PRINSTON INC	<u>2.5MG</u>	<u>A075743 001</u>	Jul 01, 2002
<u>AB</u>		<u>2.5MG</u>	<u>A076164 004</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075743 002</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076164 005</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075743 003</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076164 006</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075743 004</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A076164 001</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075743 005</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A076164 002</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075743 006</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A076164 003</u>	Jul 01, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>2.5MG</u>	<u>A212041 001</u>	Sep 15, 2020
<u>AB</u>		<u>5MG</u>	<u>A212041 002</u>	Sep 15, 2020
<u>AB</u>		<u>10MG</u>	<u>A212041 003</u>	Sep 15, 2020
<u>AB</u>		<u>20MG</u>	<u>A208920 001</u>	Mar 04, 2021
<u>AB</u>		<u>30MG</u>	<u>A208920 002</u>	Mar 04, 2021
<u>AB</u>		<u>40MG</u>	<u>A208920 003</u>	Mar 04, 2021
<u>AB</u>	SUN PHARM INDS LTD	<u>2.5MG</u>	<u>A075944 001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075944 002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075944 003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075944 004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075944 006</u>	Feb 11, 2003
<u>AB</u>		<u>40MG</u>	<u>A075944 005</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A076059 001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076059 002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076059 003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A076059 004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A076059 005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A076059 006</u>	Jul 01, 2002
<u>AB</u>	WOCKHARDT BIO AG	<u>2.5MG</u>	<u>A078402 001</u>	Apr 19, 2007
<u>AB</u>		<u>5MG</u>	<u>A078402 002</u>	Apr 19, 2007
<u>AB</u>		<u>10MG</u>	<u>A078402 003</u>	Apr 19, 2007
<u>AB</u>		<u>20MG</u>	<u>A078402 004</u>	Apr 19, 2007
<u>AB</u>		<u>30MG</u>	<u>A078402 005</u>	Apr 19, 2007

PRESCRIPTION DRUG PRODUCT LIST

LISINAPRIL

TABLET; ORAL

LISINAPRIL

<u>AB</u>		<u>40MG</u>	<u>A078402 006</u>	Apr 19, 2007
<u>ZESTRIL</u>				
<u>AB</u>	+	TWI PHARMS	<u>2.5MG</u>	<u>N019777 005</u> Apr 29, 1993
<u>AB</u>	+		<u>5MG</u>	<u>N019777 001</u> May 19, 1988
<u>AB</u>	+		<u>10MG</u>	<u>N019777 002</u> May 19, 1988
<u>AB</u>	+		<u>20MG</u>	<u>N019777 003</u> May 19, 1988
<u>AB</u>	+		<u>30MG</u>	<u>N019777 006</u> Jan 20, 1999
<u>AB</u>	+		<u>40MG</u>	<u>N019777 004</u> May 19, 1988

LITHIUM CARBONATE

CAPSULE; ORAL

LITHIUM CARBONATE

<u>AB</u>		ALEMBIC LTD	<u>150MG</u>	<u>A079159 001</u> Jan 12, 2009
<u>AB</u>			<u>300MG</u>	<u>A079159 002</u> Jan 12, 2009
<u>AB</u>			<u>600MG</u>	<u>A079159 003</u> Jan 12, 2009
<u>AB</u>		GLENMARK GENERICS	<u>150MG</u>	<u>A079139 001</u> Feb 03, 2009
<u>AB</u>			<u>300MG</u>	<u>A079139 002</u> Feb 03, 2009
<u>AB</u>			<u>600MG</u>	<u>A079139 003</u> Feb 03, 2009
<u>AB</u>		HETERO LABS LTD III	<u>150MG</u>	<u>A090702 001</u> Sep 25, 2009
<u>AB</u>			<u>300MG</u>	<u>A090702 002</u> Sep 25, 2009
<u>AB</u>			<u>600MG</u>	<u>A090702 003</u> Sep 25, 2009
<u>AB</u>	+	HIKMA	<u>150MG</u>	<u>N017812 002</u> Jan 28, 1987
<u>AB</u>	+		<u>300MG</u>	<u>N017812 001</u>
<u>AB</u>	+		<u>600MG</u>	<u>N017812 003</u> Jan 28, 1987

TABLET; ORAL

LITHIUM CARBONATE

<u>AB</u>	+	HIKMA	<u>300MG</u>	<u>N018558 001</u> Jan 29, 1982
<u>AB</u>		SUN PHARM INDS INC	<u>300MG</u>	<u>A091027 001</u> Jun 24, 2010

TABLET, EXTENDED RELEASE; ORAL

LITHIUM CARBONATE

<u>AB</u>		GLENMARK GENERICS	<u>450MG</u>	<u>A091616 001</u> Feb 14, 2011
<u>AB</u>		GLENMARK PHARMS INC	<u>300MG</u>	<u>A091544 001</u> Dec 27, 2010
<u>AB</u>		HERITAGE PHARMA	<u>300MG</u>	<u>A205532 001</u> Sep 29, 2016
<u>AB</u>		HIKMA	<u>300MG</u>	<u>A076832 001</u> Oct 28, 2004
<u>AB</u>	!		<u>450MG</u>	<u>A076691 001</u> Jan 05, 2004
<u>AB</u>		MYLAN PHARMS INC	<u>300MG</u>	<u>A202288 001</u> Jun 29, 2012
<u>AB</u>			<u>450MG</u>	<u>A202219 001</u> Aug 08, 2012
<u>AB</u>		UNIQUE	<u>300MG</u>	<u>A204779 001</u> Jul 27, 2015
<u>AB</u>			<u>450MG</u>	<u>A205663 001</u> Jun 05, 2017

LITHOBID

<u>AB</u>	+	ANI PHARMS	<u>300MG</u>	<u>N018027 001</u>
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LITHIUM CITRATE

SYRUP; ORAL

LITHIUM CITRATE

<u>AA</u>	!	PAI HOLDINGS PHARM	<u>EQ 300MG CARBONATE/5ML</u>	<u>A070755 001</u> May 21, 1986
<u>AA</u>		RUBICON	<u>EQ 300MG CARBONATE/5ML</u>	<u>A218036 001</u> Aug 14, 2023

LODOXAMIDE TROMETHAMINE

SOLUTION/DROPS; OPHTHALMIC

ALOMIDE

+	NOVARTIS	EQ 0.1% BASE	<u>N020191 001</u>	Sep 23, 1993
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LOFEXIDINE HYDROCHLORIDE

TABLET; ORAL

LUCEMYRA

+	USWM	EQ 0.18MG BASE	<u>N209229 001</u>	May 16, 2018
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LOMITAPIDE MESYLATE

CAPSULE; ORAL

JUXTAPID

+	AMRYT	EQ 5MG BASE	<u>N203858 001</u>	Dec 21, 2012
+		EQ 10MG BASE	<u>N203858 002</u>	Dec 21, 2012
+		EQ 20MG BASE	<u>N203858 003</u>	Dec 21, 2012
+		EQ 30MG BASE	<u>N203858 004</u>	Apr 23, 2015

PRESCRIPTION DRUG PRODUCT LIST

LOMUSTINE

CAPSULE; ORAL

GLEOSTINE

+	CORDEN PHARMA	10MG	N017588	001	
+		40MG	N017588	002	
+	!	100MG	N017588	003	

LONAFARNIB

CAPSULE; ORAL

ZOKINVY

+	EIGER BIOPHARMS	50MG	N213969	001	Nov 20, 2020
+	!	75MG	N213969	002	Nov 20, 2020

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

LOPERAMIDE HYDROCHLORIDE

AB	AUROBINDO PHARMA LTD	2MG	A218122	001	Sep 05, 2023
AB	BIONPHARMA	2MG	A215579	001	Oct 22, 2021
AB	EDENBRIDGE PHARMS	2MG	A215001	001	Oct 06, 2021
AB	JUBILANT CADISTA	2MG	A217840	001	Jul 05, 2023
AB	! MYLAN	2MG	A072741	001	Sep 18, 1991
AB	RUBICON	2MG	A216876	001	Jan 26, 2023
AB	TEVA	2MG	A073192	001	Apr 30, 1992
AB	ZYDUS LIFESCIENCES	2MG	A217471	001	Mar 23, 2023

LOPINAVIR; RITONAVIR

SOLUTION; ORAL

KALETRA

AA	+	ABEVIE	80MG/ML; 20MG/ML	N021251	001	Sep 15, 2000
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LOPINAVIR AND RITONAVIR

AA	LANNETT CO INC	80MG/ML; 20MG/ML	A207407	001	Dec 27, 2016
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TABLET; ORAL

KALETRA

AB	+	ABEVIE	100MG; 25MG	N021906	002	Nov 09, 2007
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AB	+	!	200MG; 50MG	N021906	001	Oct 28, 2005
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LOPINAVIR AND RITONAVIR

AB	HETERO LABS LTD III	100MG; 25MG	A091677	001	Jun 04, 2021
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AB		200MG; 50MG	A091677	002	Jun 04, 2021
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AB	LAURUS	100MG; 25MG	A213857	001	Mar 21, 2022
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AB		200MG; 50MG	A213857	002	Mar 21, 2022
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LORAZEPAM

CAPSULE, EXTENDED RELEASE; ORAL

LOREEV XR

+	ALMATICA	1MG	N214826	001	Aug 27, 2021
+		1.5MG	N214826	004	Feb 16, 2022
+		2MG	N214826	002	Aug 27, 2021
+	!	3MG	N214826	003	Aug 27, 2021

CONCENTRATE; ORAL

LORAZEPAM

AA	AMNEAL PHARMS	2MG/ML	A091383	001	Dec 23, 2009
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AA	HIKMA	2MG/ML	A200169	001	Jan 30, 2012
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AA	LUPIN LTD	2MG/ML	A091407	001	Feb 19, 2013
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AA	PHARM ASSOC	2MG/ML	A090260	001	Jun 15, 2010
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LORAZEPAM INTENSOL

AA	!	HIKMA	2MG/ML	A072755	001	Jun 28, 1991
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INJECTABLE; INJECTION

ATIVAN

AP	+	!	2MG/ML	N018140	001
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AP	+	!	4MG/ML	N018140	002
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LORAZEPAM

AP	HOSPIRA	2MG/ML	A074243	001	Apr 12, 1994
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AP		2MG/ML	A074282	001	May 27, 1994
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AP		4MG/ML	A074243	002	Apr 12, 1994
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AP		4MG/ML	A074282	002	May 27, 1994
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AP	INTL MEDICATION SYS	2MG/ML	A076150	001	Nov 15, 2004
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AP	RISING	2MG/ML	A075025	001	Jul 23, 1998
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TABLET; ORAL

ATIVAN

AB	+	BAUSCH	0.5MG	N017794	001
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AB	+		1MG	N017794	002
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AB	+	!	2MG	N017794	003
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PRESCRIPTION DRUG PRODUCT LIST

LORAZEPAM

TABLET; ORAL

LORAZEPAM

<u>AB</u>	AUROLIFE PHARMA LLC	<u>0.5MG</u>	<u>A203572 001</u>	Dec 22, 2017
<u>AB</u>		<u>1MG</u>	<u>A203572 002</u>	Dec 22, 2017
<u>AB</u>		<u>2MG</u>	<u>A203572 003</u>	Dec 22, 2017
<u>AB</u>	LEADING	<u>0.5MG</u>	<u>A078203 001</u>	Jul 30, 2007
<u>AB</u>		<u>1MG</u>	<u>A078203 002</u>	Jul 30, 2007
<u>AB</u>		<u>2MG</u>	<u>A078203 003</u>	Jul 30, 2007
<u>AB</u>	OXFORD PHARMS	<u>0.5MG</u>	<u>A077754 001</u>	May 10, 2006
<u>AB</u>		<u>1MG</u>	<u>A077754 002</u>	May 10, 2006
<u>AB</u>		<u>2MG</u>	<u>A077754 003</u>	May 10, 2006
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A071141 002</u>	Apr 21, 1987
<u>AB</u>		<u>1MG</u>	<u>A071141 003</u>	Apr 21, 1987
<u>AB</u>		<u>2MG</u>	<u>A071141 001</u>	Apr 21, 1987
<u>AB</u>	SUN PHARM INDS LTD	<u>0.5MG</u>	<u>A076045 001</u>	Aug 29, 2001
<u>AB</u>		<u>1MG</u>	<u>A076045 002</u>	Aug 29, 2001
<u>AB</u>		<u>2MG</u>	<u>A076045 003</u>	Aug 29, 2001
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>A072926 001</u>	Oct 31, 1991
<u>AB</u>		<u>1MG</u>	<u>A072927 001</u>	Oct 31, 1991
<u>AB</u>		<u>2MG</u>	<u>A072928 001</u>	Oct 31, 1991

LORLATINIB

TABLET; ORAL

LORBRENA

+	PFIZER	25MG	N210868 001	Nov 02, 2018
+	!	100MG	N210868 002	Nov 02, 2018

LOSARTAN POTASSIUM

TABLET; ORAL

COZAAR

<u>AB</u>	+	ORGANON	<u>25MG</u>	<u>N020386 001</u>	Apr 14, 1995
<u>AB</u>	+		<u>50MG</u>	<u>N020386 002</u>	Apr 14, 1995
<u>AB</u>	+	!	<u>100MG</u>	<u>N020386 003</u>	Oct 13, 1998

LOSARTAN POTASSIUM

<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A090428 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090428 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090428 003</u>	Oct 06, 2010
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A090083 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090083 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090083 003</u>	Oct 06, 2010
<u>AB</u>	GRANULES	<u>25MG</u>	<u>A215959 001</u>	Feb 23, 2023
<u>AB</u>		<u>50MG</u>	<u>A215959 002</u>	Feb 23, 2023
<u>AB</u>		<u>100MG</u>	<u>A215959 003</u>	Feb 23, 2023
<u>AB</u>	HETERO LABS LTD V	<u>25MG</u>	<u>A203835 001</u>	Aug 12, 2015
<u>AB</u>		<u>50MG</u>	<u>A203835 002</u>	Aug 12, 2015
<u>AB</u>		<u>100MG</u>	<u>A203835 003</u>	Aug 12, 2015
<u>AB</u>	IPCA LABS LTD	<u>25MG</u>	<u>A200290 001</u>	Aug 30, 2013
<u>AB</u>		<u>50MG</u>	<u>A200290 002</u>	Aug 30, 2013
<u>AB</u>		<u>100MG</u>	<u>A200290 003</u>	Aug 30, 2013
<u>AB</u>	JUBILANT CADISTA	<u>25MG</u>	<u>A201170 001</u>	Sep 18, 2012
<u>AB</u>		<u>50MG</u>	<u>A201170 002</u>	Sep 18, 2012
<u>AB</u>		<u>100MG</u>	<u>A201170 003</u>	Sep 18, 2012
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A078232 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078232 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078232 003</u>	Oct 06, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>25MG</u>	<u>A202230 001</u>	May 30, 2012
<u>AB</u>		<u>50MG</u>	<u>A202230 002</u>	May 30, 2012
<u>AB</u>		<u>100MG</u>	<u>A202230 003</u>	May 30, 2012
<u>AB</u>	MICRO LABS	<u>25MG</u>	<u>A091541 001</u>	Sep 24, 2012
<u>AB</u>		<u>50MG</u>	<u>A091541 002</u>	Sep 24, 2012
<u>AB</u>		<u>100MG</u>	<u>A091541 003</u>	Sep 24, 2012
<u>AB</u>	MSN	<u>25MG</u>	<u>A217396 001</u>	Aug 14, 2023
<u>AB</u>		<u>50MG</u>	<u>A217396 002</u>	Aug 14, 2023
<u>AB</u>		<u>100MG</u>	<u>A217396 003</u>	Aug 14, 2023
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A091497 001</u>	Jun 06, 2011
<u>AB</u>		<u>50MG</u>	<u>A091497 002</u>	Jun 06, 2011
<u>AB</u>		<u>100MG</u>	<u>A091497 003</u>	Jun 06, 2011
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A077424 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A077424 002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A077424 003</u>	Oct 06, 2010
<u>AB</u>	STRIDES PHARMA	<u>25MG</u>	<u>A090382 001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090382 002</u>	Oct 06, 2010

PRESCRIPTION DRUG PRODUCT LIST

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

<u>AB</u>		<u>100MG</u>	<u>A090382</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A203030</u>	<u>001</u>	Oct 14, 2015
<u>AB</u>		<u>50MG</u>	<u>A203030</u>	<u>002</u>	Oct 14, 2015
<u>AB</u>		<u>100MG</u>	<u>A203030</u>	<u>003</u>	Oct 14, 2015
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A091129</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A091129</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A091129</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078243</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078243</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078243</u>	<u>003</u>	Oct 06, 2010

LOTEPREDNOL ETABONATE

GEL; OPHTHALMIC

LOTEMAX

<u>AB</u>	<u>+</u> !	BAUSCH AND LOMB INC	<u>0.5%</u>	<u>N202872</u>	<u>001</u>	Sep 28, 2012
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LOTEPREDNOL ETABONATE

<u>AB</u>		SENTISS	<u>0.5%</u>	<u>A212080</u>	<u>001</u>	Feb 10, 2021
<u>AB</u>		SLAYBACK PHARMA LLC	<u>0.5%</u>	<u>A213956</u>	<u>001</u>	Nov 29, 2023
<u>AB</u>		SUN PHARM	<u>0.5%</u>	<u>A215384</u>	<u>001</u>	Aug 09, 2023

LOTEMAX SM

	<u>+</u> !	BAUSCH AND LOMB INC	0.38%	N208219	001	Feb 22, 2019
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OINTMENT; OPHTHALMIC

LOTEMAX

	<u>+</u> !	BAUSCH AND LOMB	0.5%	N200738	001	Apr 15, 2011
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SUSPENSION/DROPS; OPHTHALMIC

ALREX

<u>AB</u>	<u>+</u> !	BAUSCH AND LOMB	<u>0.2%</u>	<u>N020803</u>	<u>001</u>	Mar 09, 1998
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LOTEMAX

<u>AB</u>	<u>+</u> !	BAUSCH AND LOMB	<u>0.5%</u>	<u>N020583</u>	<u>001</u>	Mar 09, 1998
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LOTEPREDNOL ETABONATE

<u>AB</u>		LUPIN LTD	<u>0.2%</u>	<u>A215550</u>	<u>001</u>	Dec 26, 2023
<u>AB</u>		SENTISS	<u>0.2%</u>	<u>A215933</u>	<u>001</u>	Apr 12, 2023
<u>AB</u>			<u>0.5%</u>	<u>A207609</u>	<u>001</u>	Apr 17, 2019
<u>AB</u>		SUN PHARM	<u>0.5%</u>	<u>A212450</u>	<u>001</u>	Feb 26, 2021

EYSUVIS

	<u>+</u> !	ALCON LABS INC	0.25%	N210933	001	Oct 26, 2020
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INVELTYS

	<u>+</u> !	ALCON LABS INC	1%	N210565	001	Aug 22, 2018
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LOTEPREDNOL ETABONATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC

ZYLET

	<u>+</u> !	BAUSCH AND LOMB	0.5%; 0.3%	N050804	001	Dec 14, 2004
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LOTILANER

SOLUTION/DROPS; OPHTHALMIC

XDEMY

	<u>+</u> !	TARSUS	0.25%	N217603	001	Jul 24, 2023
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LOVASTATIN

TABLET; ORAL

LOVASTATIN

<u>AB</u>		ACTAVIS ELIZABETH	<u>10MG</u>	<u>A075828</u>	<u>001</u>	Dec 17, 2001
<u>AB</u>			<u>20MG</u>	<u>A075828</u>	<u>002</u>	Dec 17, 2001
<u>AB</u>			<u>40MG</u>	<u>A075828</u>	<u>003</u>	Dec 17, 2001
<u>AB</u>		CARLSBAD	<u>10MG</u>	<u>A075991</u>	<u>001</u>	Jun 05, 2002
<u>AB</u>			<u>20MG</u>	<u>A075991</u>	<u>002</u>	Jun 05, 2002
<u>AB</u>	<u>!</u>		<u>40MG</u>	<u>A075991</u>	<u>003</u>	Jun 05, 2002
<u>AB</u>		CHARTWELL RX	<u>10MG</u>	<u>A075300</u>	<u>001</u>	Dec 17, 2001
<u>AB</u>			<u>20MG</u>	<u>A075300</u>	<u>002</u>	Dec 17, 2001
<u>AB</u>			<u>40MG</u>	<u>A075300</u>	<u>003</u>	Dec 17, 2001
<u>AB</u>		COREPHARMA	<u>10MG</u>	<u>A077748</u>	<u>001</u>	Feb 28, 2007
<u>AB</u>			<u>20MG</u>	<u>A077748</u>	<u>002</u>	Feb 28, 2007
<u>AB</u>			<u>40MG</u>	<u>A077748</u>	<u>003</u>	Feb 28, 2007
<u>AB</u>		EPIC PHARMA LLC	<u>10MG</u>	<u>A075636</u>	<u>001</u>	Dec 17, 2001
<u>AB</u>			<u>20MG</u>	<u>A075636</u>	<u>002</u>	Dec 17, 2001
<u>AB</u>			<u>40MG</u>	<u>A075636</u>	<u>003</u>	Dec 17, 2001
<u>AB</u>		LUPIN	<u>10MG</u>	<u>A078296</u>	<u>001</u>	Mar 14, 2008
<u>AB</u>			<u>20MG</u>	<u>A078296</u>	<u>002</u>	Nov 01, 2007
<u>AB</u>			<u>40MG</u>	<u>A078296</u>	<u>003</u>	Nov 01, 2007
<u>AB</u>		TEVA	<u>10MG</u>	<u>A075551</u>	<u>003</u>	Dec 17, 2001

PRESCRIPTION DRUG PRODUCT LIST

LOVASTATIN

TABLET; ORAL

LOVASTATIN

<u>AB</u>		<u>20MG</u>	<u>A075551 002</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075551 001</u>	Dec 17, 2001

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

+	COVIS	20MG	N021316 002	Jun 26, 2002
+		40MG	N021316 003	Jun 26, 2002
+	!	60MG	N021316 004	Jun 26, 2002

LOXAPINE

POWDER; INHALATION

ADASUVE

+	!	ALEXZA PHARMS	10MG	N022549 001	Dec 21, 2012
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LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

<u>AB</u>	ELITE LABS INC	<u>EQ 5MG BASE</u>	<u>A076868 001</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076868 002</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A076868 003</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076868 004</u>	Aug 04, 2005
<u>AB</u>	LANNETT CO INC	<u>EQ 5MG BASE</u>	<u>A090695 001</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090695 002</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A090695 003</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090695 004</u>	Sep 26, 2011
<u>AB</u>	WATSON LABS	<u>EQ 5MG BASE</u>	<u>A072204 001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A072205 001</u>	Jun 15, 1988
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A072206 001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A072062 001</u>	Jun 15, 1988

LUBIPROSTONE

CAPSULE; ORAL

AMITIZA

<u>AB</u>	+	SUCAMPO PHARMA LLC	<u>8MCG</u>	<u>N021908 002</u>	Apr 29, 2008
<u>AB</u>	+	!	<u>24MCG</u>	<u>N021908 001</u>	Jan 31, 2006

LUBIPROSTONE

<u>AB</u>		AMNEAL	<u>8MCG</u>	<u>A209450 001</u>	Nov 30, 2021
<u>AB</u>			<u>24MCG</u>	<u>A209450 002</u>	Nov 30, 2021
<u>AB</u>		DR REDDYS	<u>8MCG</u>	<u>A206994 001</u>	Feb 08, 2022
<u>AB</u>			<u>24MCG</u>	<u>A206994 002</u>	Feb 08, 2022
<u>AB</u>		TEVA PHARMS USA INC	<u>8MCG</u>	<u>A209920 001</u>	Jan 18, 2022
<u>AB</u>			<u>24MCG</u>	<u>A209920 002</u>	Jan 18, 2022
<u>AB</u>		ZYDUS PHARMS	<u>8MCG</u>	<u>A214131 001</u>	Mar 23, 2023
<u>AB</u>			<u>24MCG</u>	<u>A214131 002</u>	Mar 23, 2023

LULICONAZOLE

CREAM; TOPICAL

LUZU

+	!	BAUSCH	1%	N204153 001	Nov 14, 2013
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LUMASIRAN SODIUM

SOLUTION; SUBCUTANEOUS

OXLUMO

+	!	ALNYLAM PHARMS INC	EQ 94.5MG BASE/0.5ML (EQ 94.5MG BASE/0.5ML)	N214103 001	Nov 23, 2020
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LUMATEPERONE TOSYLATE

CAPSULE; ORAL

CAPLYTA

+		INTRA-CELLULAR	EQ 10.5MG BASE	N209500 002	Apr 22, 2022
+			EQ 21MG BASE	N209500 003	Apr 22, 2022
+	!		EQ 42MG BASE	N209500 001	Dec 20, 2019

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LATUDA

<u>AB</u>	+	SUNOVION PHARMS INC	<u>20MG</u>	<u>N200603 003</u>	Dec 07, 2011
<u>AB</u>	+	!	<u>40MG</u>	<u>N200603 001</u>	Oct 28, 2010
<u>AB</u>	+		<u>60MG</u>	<u>N200603 005</u>	Jul 12, 2013
<u>AB</u>	+		<u>80MG</u>	<u>N200603 002</u>	Oct 28, 2010
<u>AB</u>	+		<u>120MG</u>	<u>N200603 004</u>	Apr 26, 2012

LURASIDONE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>20MG</u>	<u>A208049 001</u>	Jan 03, 2019
<u>AB</u>			<u>40MG</u>	<u>A208049 002</u>	Jan 03, 2019

PRESCRIPTION DRUG PRODUCT LIST

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

<u>AB</u>		<u>60MG</u>	<u>A208049 003</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208049 004</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208049 005</u>	Jan 03, 2019
<u>AB</u>	ALEMBIC	<u>20MG</u>	<u>A213248 001</u>	May 13, 2021
<u>AB</u>		<u>40MG</u>	<u>A213248 002</u>	May 13, 2021
<u>AB</u>		<u>60MG</u>	<u>A213248 003</u>	May 13, 2021
<u>AB</u>		<u>80MG</u>	<u>A213248 004</u>	May 13, 2021
<u>AB</u>		<u>120MG</u>	<u>A213248 005</u>	May 13, 2021
<u>AB</u>	ALKEM LABS LTD	<u>20MG</u>	<u>A212244 001</u>	Dec 13, 2022
<u>AB</u>		<u>40MG</u>	<u>A212244 002</u>	Dec 13, 2022
<u>AB</u>		<u>60MG</u>	<u>A212244 003</u>	Dec 13, 2022
<u>AB</u>		<u>80MG</u>	<u>A212244 004</u>	Dec 13, 2022
<u>AB</u>		<u>120MG</u>	<u>A212244 005</u>	Dec 13, 2022
<u>AB</u>	AMNEAL PHARMS CO	<u>20MG</u>	<u>A208002 001</u>	Jan 03, 2019
<u>AB</u>		<u>40MG</u>	<u>A208002 002</u>	Jan 03, 2019
<u>AB</u>		<u>60MG</u>	<u>A208002 003</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208002 004</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208002 005</u>	Jan 03, 2019
<u>AB</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A208045 001</u>	Mar 10, 2023
<u>AB</u>		<u>40MG</u>	<u>A208045 002</u>	Mar 10, 2023
<u>AB</u>		<u>60MG</u>	<u>A208045 003</u>	Mar 10, 2023
<u>AB</u>		<u>80MG</u>	<u>A208045 004</u>	Mar 10, 2023
<u>AB</u>		<u>120MG</u>	<u>A208045 005</u>	Mar 10, 2023
<u>AB</u>	AVET LIFESCIENCES	<u>20MG</u>	<u>A208058 001</u>	Sep 04, 2019
<u>AB</u>		<u>40MG</u>	<u>A208058 002</u>	Sep 04, 2019
<u>AB</u>		<u>60MG</u>	<u>A208058 003</u>	Sep 04, 2019
<u>AB</u>		<u>80MG</u>	<u>A208058 004</u>	Sep 04, 2019
<u>AB</u>		<u>120MG</u>	<u>A208058 005</u>	Feb 28, 2023
<u>AB</u>	DR REDDYS	<u>20MG</u>	<u>A208047 001</u>	Aug 24, 2021
<u>AB</u>		<u>40MG</u>	<u>A208047 002</u>	Aug 24, 2021
<u>AB</u>		<u>60MG</u>	<u>A208047 003</u>	Aug 24, 2021
<u>AB</u>		<u>80MG</u>	<u>A208047 004</u>	Aug 24, 2021
<u>AB</u>	INVAGEN PHARMS	<u>20MG</u>	<u>A208028 001</u>	Jan 03, 2019
<u>AB</u>		<u>40MG</u>	<u>A208028 002</u>	Jan 03, 2019
<u>AB</u>		<u>60MG</u>	<u>A208028 003</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208028 004</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208028 005</u>	Jan 03, 2019
<u>AB</u>	LUPIN LTD	<u>20MG</u>	<u>A208031 001</u>	Jan 03, 2019
<u>AB</u>		<u>40MG</u>	<u>A208031 002</u>	Jan 03, 2019
<u>AB</u>		<u>60MG</u>	<u>A208031 003</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208031 004</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208031 005</u>	Jan 03, 2019
<u>AB</u>	MACLEODS PHARMS LTD	<u>20MG</u>	<u>A212124 001</u>	Jun 09, 2023
<u>AB</u>		<u>40MG</u>	<u>A212124 002</u>	Jun 09, 2023
<u>AB</u>		<u>60MG</u>	<u>A212124 003</u>	Jun 09, 2023
<u>AB</u>		<u>80MG</u>	<u>A212124 004</u>	Jun 09, 2023
<u>AB</u>		<u>120MG</u>	<u>A212124 005</u>	Jun 09, 2023
<u>AB</u>	MSN	<u>20MG</u>	<u>A208037 001</u>	Sep 09, 2022
<u>AB</u>		<u>40MG</u>	<u>A208037 002</u>	Sep 09, 2022
<u>AB</u>		<u>60MG</u>	<u>A208037 003</u>	Sep 09, 2022
<u>AB</u>		<u>80MG</u>	<u>A208037 004</u>	Sep 09, 2022
<u>AB</u>		<u>120MG</u>	<u>A208037 005</u>	Sep 09, 2022
<u>AB</u>	PIRAMAL HLTHCARE UK	<u>20MG</u>	<u>A212091 001</u>	Dec 28, 2020
<u>AB</u>		<u>40MG</u>	<u>A212091 002</u>	Dec 28, 2020
<u>AB</u>		<u>60MG</u>	<u>A212091 003</u>	Dec 28, 2020
<u>AB</u>		<u>80MG</u>	<u>A212091 004</u>	Dec 28, 2020
<u>AB</u>		<u>120MG</u>	<u>A212091 005</u>	Dec 28, 2020
<u>AB</u>	SUN PHARM	<u>20MG</u>	<u>A208066 001</u>	Jan 04, 2019
<u>AB</u>		<u>40MG</u>	<u>A208066 002</u>	Jan 04, 2019
<u>AB</u>		<u>60MG</u>	<u>A208066 003</u>	Jan 04, 2019
<u>AB</u>		<u>80MG</u>	<u>A208066 004</u>	Jan 04, 2019
<u>AB</u>		<u>120MG</u>	<u>A208066 005</u>	Jan 04, 2019
<u>AB</u>	TORRENT	<u>20MG</u>	<u>A208055 001</u>	Jan 03, 2019
<u>AB</u>		<u>40MG</u>	<u>A208055 002</u>	Jan 03, 2019
<u>AB</u>		<u>80MG</u>	<u>A208055 003</u>	Jan 03, 2019
<u>AB</u>		<u>120MG</u>	<u>A208055 004</u>	Jan 03, 2019
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A208052 001</u>	Mar 19, 2019
<u>AB</u>		<u>40MG</u>	<u>A208052 002</u>	Mar 19, 2019
<u>AB</u>		<u>60MG</u>	<u>A208052 003</u>	Mar 19, 2019

PRESCRIPTION DRUG PRODUCT LISTLURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

AB		80MG	A208052 004	Mar 19, 2019
AB		120MG	A208052 005	Mar 19, 2019

LURBINECTEDIN

POWDER; INTRAVENOUS

ZEPZELCA

+	!	JAZZ	4MG/VIAL	N213702 001	Jun 15, 2020
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LUSUTROMBOPAG

TABLET; ORAL

MULPLETA

+	!	VANCOCIN ITALIA	3MG	N210923 001	Jul 31, 2018
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LUTETIUM LU 177 DOTATATE

SOLUTION; INTRAVENOUS

LUTATHERA

+	!	AAA USA INC	10mCi/ML	N208700 001	Jan 26, 2018
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LUTETIUM LU-177 VIPIVOTIDE TETRAJETAN

SOLUTION; INTRAVENOUS

PLUVICTO

+	!	NOVARTIS	27mCi/ML	N215833 001	Mar 23, 2022
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MACITENTAN

TABLET; ORAL

MACITENTAN

AB		APOTEX	10MG	A211195 001	Jan 09, 2024
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OPSUMIT

AB	+	!	ACTELION	10MG	N204410 001	Oct 18, 2013
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MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYLON

+	!	RISING	EQ 85MG BASE/GM	N016763 001	
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MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

+	!	B BRAUN	30MG/100ML; 37MG/100ML; 0.82MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML; 12MG/100ML	N019696 001	Sep 29, 1989
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MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 5.5

AP		FRESENIUS KABI USA	30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	A215370 001	Jun 29, 2022
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MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 7.4

AP		FRESENIUS KABI USA	30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	A215371 001	Jun 08, 2022
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PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER

AP	+	!	BAXTER HLTHCARE	30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	N017378 001
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PLASMA-LYTE A IN PLASTIC CONTAINER

AP	+	!	BAXTER HLTHCARE	30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	N017378 002	Nov 22, 1982
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ISOLYTE S IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML	N019711 001	Sep 29, 1989
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NORMOSOL-R IN PLASTIC CONTAINER

ICU MEDICAL INC

30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML	N017586 001	
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SOLUTION; IRRIGATION

PHYSIOLYTE IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML	N019024 001	Jun 08, 1984
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PHYSIOSOL IN PLASTIC CONTAINER

ICU MEDICAL INC

30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML	N017637 002	Jul 08, 1982
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PRESCRIPTION DRUG PRODUCT LIST

MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

SOLUTION; INJECTION

NORMOCARB HF 25

+! DIALYSIS SUPS 0.21GM/100ML; 2.8GM/100ML; 9.07GM/100ML N021910 001 Jul 26, 2006

NORMOCARB HF 35

+! DIALYSIS SUPS 0.21GM/100ML; 3.97GM/100ML; 8.3GM/100ML N021910 002 Jul 26, 2006

MAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATEAP B BRAUN MEDICAL INC 2GM/50ML (40MG/ML) A207967 001 Apr 26, 2021AP 4GM/100ML (40MG/ML) A207967 002 Apr 26, 2021AP 4GM/50ML (80MG/ML) A207967 003 Apr 26, 2021MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINERAP B BRAUN MEDICAL INC 1GM/100ML A207966 001 Oct 27, 2020AP BAXTER HLTHCARE 1GM/100ML A211965 001 Aug 11, 2020

CORP

AP FRESENIUS KABI USA 1GM/100ML A206486 001 Mar 07, 2016AP +! HOSPIRA 1GM/100ML N020488 001 Jul 11, 1995AP HQ SPCLT PHARMA 1GM/100ML A207349 001 Mar 02, 2016AP MYLAN LABS LTD 1GM/100ML A209932 001 Sep 10, 2018MAGNESIUM SULFATE IN PLASTIC CONTAINERAP AMNEAL 2GM/50ML (40MG/ML) A216597 001 Feb 15, 2023AP 4GM/100ML (40MG/ML) A216597 002 Feb 15, 2023AP BAXTER HLTHCARE 2GM/50ML (40MG/ML) A211966 001 Jun 01, 2020

CORP

AP 4GM/50ML (80MG/ML) A211966 002 Jun 01, 2020AP 4GM/100ML (40MG/ML) A211966 003 Jun 01, 2020AP FRESENIUS KABI USA 4GM/100ML (40MG/ML) A206485 001 Mar 15, 2016AP 4GM/50ML (80MG/ML) A206485 002 Mar 15, 2016AP 2GM/50ML (40MG/ML) A206485 003 Mar 15, 2016AP 20GM/500ML (40MG/ML) A206485 004 Mar 15, 2016AP 40GM/1000ML (40MG/ML) A206485 005 Mar 15, 2016AP GLAND PHARMA LTD 2GM/50ML (40MG/ML) A213917 001 Jul 10, 2020AP 4GM/100ML (40MG/ML) A213917 002 Jul 10, 2020AP + HOSPIRA 2GM/50ML (40MG/ML) N020309 003 Jan 26, 2007AP +! 4GM/100ML (40MG/ML) N020309 001 Jun 24, 1994AP +! 4GM/50ML (80MG/ML) N020309 002 Jun 24, 1994AP + 20GM/500ML (40MG/ML) N020309 004 Jan 18, 1995AP + 40GM/1000ML (40MG/ML) N020309 005 Jan 18, 1995AP HQ SPCLT PHARMA 2GM/50ML (40MG/ML) A207350 001 Dec 06, 2017AP 4GM/100ML (40MG/ML) A207350 002 Dec 06, 2017AP 4GM/50ML (80MG/ML) A207350 003 Dec 06, 2017AP 20GM/500ML (40MG/ML) A207350 004 Dec 06, 2017AP 40GM/1000ML (40MG/ML) A207350 005 Dec 06, 2017AP MILLA PHARMS 2GM/50ML (40MG/ML) A209642 001 Nov 08, 2021AP 4GM/100ML (40MG/ML) A209642 002 Nov 08, 2021AP 4GM/50ML (80MG/ML) A209642 003 Nov 08, 2021AP MYLAN LABS LTD 2GM/50ML (40MG/ML) A209911 001 Sep 14, 2018AP 4GM/100ML (40MG/ML) A209911 002 Sep 14, 2018MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+! HOSPIRA 2GM/100ML N020488 002 Jul 11, 1995

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATEAP EXELA PHARMA 5GM/10ML (500MG/ML) A206039 001 Dec 18, 2014AP +! FRESENIUS KABI USA 5GM/10ML (500MG/ML) N019316 001 Sep 08, 1986AP +! 10GM/20ML (500MG/ML) N019316 003 Jan 29, 2016AP ! HOSPIRA 5GM/10ML (500MG/ML) A075151 001 Apr 25, 2000AP 10GM/20ML (500MG/ML) A202411 001 May 14, 2015

+! FRESENIUS KABI USA 1GM/2ML (500MG/ML) N019316 002 Sep 08, 1986

+! 25GM/50ML (500MG/ML) N019316 004 Jan 29, 2016

MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

SUFLAVE

+! BRAINTREE LABS 0.9GM/BOT; 178.7GM/BOT; 1.12GM/BOT; 0.5GM/BOT; 7.3GM/BOT N215344 001 Jun 15, 2023

PRESCRIPTION DRUG PRODUCT LIST

MAGNESIUM SULFATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

TIS-U-SOL

AT	BAXTER HLTHCARE	20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML	N018508 001	Feb 19, 1982
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TIS-U-SOL IN PLASTIC CONTAINER

AT	BAXTER HLTHCARE	20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML	N018336 001	
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MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM SULFATE

TABLET; ORAL

SUTAB

+	BRAINTREE LABS	0.225GM; 0.188GM; 1.479GM	N213135 001	Nov 10, 2020
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MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

SOLUTION; ORAL

SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE

AA	ALKEM LABS LTD	1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT	A213924 001	Dec 29, 2023
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AA	NOVEL LABS INC	1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT	A202511 001	Feb 23, 2017
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AA	STRIDES PHARMA	1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT	A215469 001	Nov 22, 2023
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SUPREP BOWEL PREP KIT

AA	+	BRAINTREE LABS	1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT	N022372 001	Aug 05, 2010
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MALATHION

LOTION; TOPICAL

MALATHION

!	SUVEN PHARMS	0.5%	A091559 001	May 23, 2012
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MANGANESE CHLORIDE

INJECTABLE; INJECTION

MANGANESE CHLORIDE IN PLASTIC CONTAINER

+	HOSPIRA	EQ 0.1MG MANGANESE/ML	N018962 001	Jun 26, 1986
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MANNITOL

INJECTABLE; INJECTION

MANNITOL 10% IN PLASTIC CONTAINER

AP	B BRAUN	10GM/100ML	N020006 002	Jul 26, 1993
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MANNITOL 15% IN PLASTIC CONTAINER

AP	B BRAUN	15GM/100ML	N020006 003	Jul 26, 1993
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MANNITOL 20% IN PLASTIC CONTAINER

AP	B BRAUN	20GM/100ML	N020006 004	Jul 26, 1993
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AP	ICU MEDICAL INC	20GM/100ML	N019603 004	Jan 08, 1990
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MANNITOL 25%

AP	FRESENIUS KABI USA	12.5GM/50ML	A080677 001	
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AP	HOSPIRA	12.5GM/50ML	N016269 006	Aug 25, 1994
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MANNITOL 5% IN PLASTIC CONTAINER

AP	B BRAUN	5GM/100ML	N020006 001	Jul 26, 1993
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OSMITROL 10% IN WATER

AP	BAXTER HLTHCARE	10GM/100ML	N013684 002	
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OSMITROL 10% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	10GM/100ML	N013684 006	
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OSMITROL 15% IN WATER

AP	BAXTER HLTHCARE	15GM/100ML	N013684 004	
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OSMITROL 15% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	15GM/100ML	N013684 008	
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OSMITROL 20% IN WATER

AP	BAXTER HLTHCARE	20GM/100ML	N013684 003	
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OSMITROL 20% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	20GM/100ML	N013684 007	
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OSMITROL 5% IN WATER

AP	BAXTER HLTHCARE	5GM/100ML	N013684 001	
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OSMITROL 5% IN WATER IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML	N013684 005	
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POWDER; INHALATION

ARIDOL KIT

+	ARNA PHARMA	N/A, 5MG, 10MG, 20MG, 40MG	N022368 001	Oct 05, 2010
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BRONCHITOL

+	CHIESI	40MG	N202049 001	Oct 30, 2020
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PRESCRIPTION DRUG PRODUCT LISTMANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL IN PLASTIC CONTAINER

+! ICU MEDICAL INC 540MG/100ML; 2.7GM/100ML N018316 001

MARALIXIBAT CHLORIDE

SOLUTION; ORAL

LIVMARLI

+! MIRUM EQ 9.5MG BASE/ML N214662 001 Sep 29, 2021

MARAVIROC

SOLUTION; ORAL

SELZENTRY

+! VIIV HLTHCARE 20MG/ML N208984 001 Nov 04, 2016

TABLET; ORAL

MARAVIROC**AB** HETERO LABS LTD III **150MG** **A203347 001** Feb 07, 2022**AB** **300MG** **A203347 002** Feb 07, 2022**AB** I 3 PHARMS **150MG** **A217114 001** Aug 17, 2023**AB** **300MG** **A217114 002** Aug 17, 2023SELZENTRY**AB** + VIIV HLTHCARE **150MG** **N022128 001** Aug 06, 2007**AB** +! **300MG** **N022128 002** Aug 06, 2007

+ 25MG N022128 003 Nov 04, 2016

+ 75MG N022128 004 Nov 04, 2016

MARIBAVIR

TABLET; ORAL

LIVTENCITY

+! TAKEDA PHARMS USA 200MG N215596 001 Nov 23, 2021

MAVACAMTEN

CAPSULE; ORAL

CAMZYOS

+ BRISTOL 2.5MG N214998 001 Apr 28, 2022

+! 5MG N214998 002 Apr 28, 2022

+ 10MG N214998 003 Apr 28, 2022

+ 15MG N214998 004 Apr 28, 2022

MEBENDAZOLE

TABLET, CHEWABLE; ORAL

EMVERM

! IMPAX LABS INC 100MG A073580 001 Jan 04, 1995

MECAMYLAMINE HYDROCHLORIDE

TABLET; ORAL

MECAMYLAMINE HYDROCHLORIDE

! LGM PHARMA 2.5MG A204054 001 Mar 19, 2013

MECHLORETHAMINE HYDROCHLORIDE

GEL; TOPICAL

VALCHLOR

+! HELSINN EQ 0.016% BASE N202317 001 Aug 23, 2013

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

ANTIVERT**AA** + CASPER PHARMA LLC **12.5MG** **N010721 006****AA** + **25MG** **N010721 004****AA** + **50MG** **N010721 001** Jan 20, 1982MECLIZINE HYDROCHLORIDE**AA** ! AMNEAL PHARMS **12.5MG** **A201451 001** Feb 23, 2011**AA** ! **25MG** **A201451 002** Feb 23, 2011**AA** APNAR PHARMA LP **12.5MG** **A087128 002****AA** **25MG** **A087128 001****AA** AUROBINDO PHARMA USA **12.5MG** **A202640 001** Sep 17, 2012**AA** **25MG** **A202640 002** Sep 17, 2012**AA** **50MG** **A202640 003** Sep 17, 2012**AA** EPIC PHARMA LLC **12.5MG** **A200294 001** Apr 13, 2012**AA** **25MG** **A200294 002** Apr 13, 2012**AA** INVAGEN PHARMS **12.5MG** **A200432 001** Feb 17, 2022**AA** **25MG** **A200432 002** Feb 17, 2022**AA** **50MG** **A200432 003** Feb 17, 2022**AA** INVATECH **25MG** **A084092 003** May 22, 1989**AA** JUBILANT CADISTA **12.5MG** **A040659 001** Jun 04, 2010**AA** **25MG** **A040659 002** Jun 04, 2010

PRESCRIPTION DRUG PRODUCT LIST

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

MECLIZINE HYDROCHLORIDE

<u>AA</u>	LUPIN LTD	<u>12.5MG</u>	<u>A203003 001</u>	Aug 11, 2022
<u>AA</u>		<u>25MG</u>	<u>A203003 002</u>	Aug 11, 2022
<u>AA</u>		<u>50MG</u>	<u>A203003 003</u>	Aug 11, 2022
<u>AA</u>	SANDOZ	<u>12.5MG</u>	<u>A084843 002</u>	May 22, 1989
<u>AA</u>	WILSHIRE PHARMS INC	<u>12.5MG</u>	<u>A205136 001</u>	Feb 22, 2019
<u>AA</u>		<u>25MG</u>	<u>A205136 002</u>	Feb 22, 2019
<u>AA</u>		<u>50MG</u>	<u>A205136 003</u>	Feb 22, 2019
<u>AA</u>	ZYDUS	<u>12.5MG</u>	<u>A213957 001</u>	Jun 23, 2020
<u>AA</u>		<u>25MG</u>	<u>A213957 002</u>	Jun 23, 2020

TABLET, CHEWABLE; ORAL

ANTIVERT

<u>AA</u>	+ CASPER PHARMA LLC	<u>25MG</u>	<u>N010721 005</u>	
<u>AA</u>	<u>MECLIZINE HYDROCHLORIDE</u> INVAGEN PHARMS	<u>25MG</u>	<u>A200791 001</u>	Feb 17, 2022

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLOFENAMATE SODIUM

	MYLAN	EQ 50MG BASE	A071081 002	Sep 03, 1986
!		EQ 100MG BASE	A071081 001	Sep 03, 1986

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

<u>AB</u>	+! PFIZER	<u>150MG/ML</u>	<u>N020246 001</u>	Oct 29, 1992
<u>AB</u>	<u>MECLOFENAMATE SODIUM</u> AMNEAL	<u>150MG/ML</u>	<u>A215397 001</u>	Jun 07, 2023
<u>AB</u>	AMPHASTAR PHARMS INC	<u>150MG/ML</u>	<u>A077235 001</u>	Nov 28, 2017
<u>AB</u>		<u>150MG/ML</u>	<u>A077334 001</u>	Nov 28, 2017
<u>AB</u>	EUGIA PHARMA	<u>150MG/ML</u>	<u>A212824 001</u>	Aug 22, 2022
<u>AB</u>		<u>150MG/ML</u>	<u>A212844 001</u>	Aug 31, 2022
<u>AB</u>	MEITHEAL	<u>150MG/ML</u>	<u>A076553 001</u>	Jul 28, 2004
<u>AB</u>	MYLAN LABS LTD	<u>150MG/ML</u>	<u>A210227 001</u>	Oct 12, 2018
	INJECTABLE; SUBCUTANEOUS DEPO-SUBQ PROVERA 104 +! PFIZER	104MG/0.65ML	N021583 001	Dec 17, 2004

TABLET; ORAL

MECLOFENAMATE SODIUM

<u>AB</u>	BARR	<u>2.5MG</u>	<u>A040159 001</u>	Aug 09, 1996
<u>AB</u>		<u>5MG</u>	<u>A040159 002</u>	Aug 09, 1996
<u>AB</u>		<u>10MG</u>	<u>A040159 003</u>	Aug 09, 1996
<u>AB</u>	<u>PROVERA</u> + PFIZER	<u>2.5MG</u>	<u>N011839 001</u>	
<u>AB</u>	+	<u>5MG</u>	<u>N011839 003</u>	
<u>AB</u>	+!	<u>10MG</u>	<u>N011839 004</u>	

MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

<u>AB</u>	BELCHER	<u>250MG</u>	<u>A091608 001</u>	Jun 02, 2014
<u>AB</u>	LUPIN LTD	<u>250MG</u>	<u>A091322 001</u>	Jul 22, 2011
<u>AB</u>	MICRO LABS	<u>250MG</u>	<u>A090562 001</u>	Nov 19, 2010
<u>AB</u>	<u>PONSTEL</u> +! AVION PHARMS	<u>250MG</u>	<u>N015034 003</u>	

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

MEFLOQUINE HYDROCHLORIDE

<u>AB</u>	! BARR	<u>250MG</u>	<u>A076392 001</u>	Dec 29, 2003
<u>AB</u>	HIKMA	<u>250MG</u>	<u>A076523 001</u>	Oct 01, 2004

MEGESTROL ACETATE

SUSPENSION; ORAL

MEGESTROL ACETATE

<u>AB</u>	+! ENDO PHARMS INC	<u>125MG/ML</u>	<u>N021778 001</u>	Jul 05, 2005
<u>AB</u>	<u>MEGESTROL ACETATE</u> HIKMA	<u>40MG/ML</u>	<u>A203960 001</u>	Jun 09, 2017
<u>AB</u>	! STRIDES PHARMA	<u>40MG/ML</u>	<u>A075671 001</u>	Jul 25, 2001
<u>AB</u>	TWI PHARMS	<u>125MG/ML</u>	<u>A203139 001</u>	Aug 27, 2014

PRESCRIPTION DRUG PRODUCT LIST

MEGESTROL ACETATE

TABLET; ORAL

MEGESTROL ACETATE

<u>AB</u>	BARR	<u>20MG</u>	<u>A074621</u>	<u>002</u>	Aug 16, 1996
<u>AB</u>		<u>40MG</u>	<u>A074621</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>	STRIDES PHARMA	<u>20MG</u>	<u>A072422</u>	<u>001</u>	Aug 08, 1988
<u>AB</u>	!	<u>40MG</u>	<u>A072423</u>	<u>001</u>	Aug 08, 1988

MELOXICAM

CAPSULE; ORAL

MELOXICAM

<u>AB</u>	LUPIN LTD	<u>5MG</u>	<u>A209487</u>	<u>001</u>	Jun 01, 2020
<u>AB</u>	!	<u>10MG</u>	<u>A209487</u>	<u>002</u>	Jun 01, 2020
<u>AB</u>	NOVITIUM PHARMA	<u>5MG</u>	<u>A211398</u>	<u>001</u>	Mar 09, 2021
<u>AB</u>		<u>10MG</u>	<u>A211398</u>	<u>002</u>	Mar 09, 2021

SUSPENSION; ORAL

MELOXICAM

+! AVONDALE PHARMS 7.5MG/5ML N021530 001 Jun 01, 2004

TABLET; ORAL

MELOXICAM

<u>AB</u>	ASCENT PHARMS INC	<u>7.5MG</u>	<u>A217579</u>	<u>001</u>	Jun 07, 2023
<u>AB</u>		<u>15MG</u>	<u>A217579</u>	<u>002</u>	Jun 07, 2023
<u>AB</u>	AUROBINDO PHARMA	<u>7.5MG</u>	<u>A078008</u>	<u>001</u>	Oct 02, 2006
<u>AB</u>		<u>15MG</u>	<u>A078008</u>	<u>002</u>	Oct 02, 2006
<u>AB</u>	CIPLA	<u>7.5MG</u>	<u>A077929</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077929</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	COREPHARMA	<u>7.5MG</u>	<u>A077882</u>	<u>001</u>	Jul 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077882</u>	<u>002</u>	Jul 20, 2006
<u>AB</u>	DR REDDYS LABS INC	<u>7.5MG</u>	<u>A077931</u>	<u>001</u>	Jul 25, 2006
<u>AB</u>		<u>15MG</u>	<u>A077931</u>	<u>002</u>	Jul 25, 2006
<u>AB</u>	GLENMARK GENERICS	<u>7.5MG</u>	<u>A077932</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077932</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	LUPIN PHARMS	<u>7.5MG</u>	<u>A077944</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077944</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	PURACAP PHARM	<u>7.5MG</u>	<u>A077938</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077938</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	STRIDES PHARMA	<u>7.5MG</u>	<u>A077928</u>	<u>001</u>	May 13, 2009
<u>AB</u>		<u>15MG</u>	<u>A077928</u>	<u>002</u>	May 13, 2009
<u>AB</u>	TARO	<u>7.5MG</u>	<u>A078102</u>	<u>001</u>	Nov 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A078102</u>	<u>002</u>	Nov 07, 2006
<u>AB</u>	UNICHEM	<u>7.5MG</u>	<u>A077927</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077927</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>	YUNG SHIN PHARM	<u>7.5MG</u>	<u>A077918</u>	<u>001</u>	Dec 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A077918</u>	<u>002</u>	Dec 07, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>7.5MG</u>	<u>A077921</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077921</u>	<u>002</u>	Jul 19, 2006

MOBIC

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>7.5MG</u>	<u>N020938</u>	<u>001</u>	Apr 13, 2000
<u>AB</u>	+!	<u>15MG</u>	<u>N020938</u>	<u>002</u>	Aug 23, 2000

MELPHALAN

TABLET; ORAL

MELPHALAN

! ALVOGEN 2MG A207809 001 Mar 22, 2017

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

MELPHALAN HYDROCHLORIDE

<u>AP</u>	ACTAVIS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A206018</u>	<u>001</u>	Dec 19, 2016
<u>AP</u>	ALMAJECT	<u>EQ 50MG BASE/VIAL</u>	<u>A204817</u>	<u>001</u>	May 17, 2019
<u>AP</u>	ARTHUR GRP	<u>EQ 50MG BASE/VIAL</u>	<u>A211463</u>	<u>001</u>	Sep 13, 2019
<u>AP</u>	BPI LABS	<u>EQ 50MG BASE/VIAL</u>	<u>A209197</u>	<u>001</u>	May 08, 2020
<u>AP</u>	DR REDDYS	<u>EQ 50MG BASE/VIAL</u>	<u>A203655</u>	<u>001</u>	Dec 08, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 50MG BASE/VIAL</u>	<u>A203393</u>	<u>001</u>	Dec 22, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 50MG BASE/VIAL</u>	<u>A209826</u>	<u>001</u>	May 28, 2019
<u>AP</u>	HIKMA	<u>EQ 50MG BASE/VIAL</u>	<u>A090303</u>	<u>001</u>	Oct 28, 2010
<u>AP</u>	INGENUS PHARMS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A210947</u>	<u>001</u>	Feb 18, 2020
<u>AP</u>	MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A212960</u>	<u>001</u>	May 28, 2021
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>EQ 50MG BASE/VIAL</u>	<u>A090270</u>	<u>001</u>	Jun 09, 2009
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 50MG BASE/VIAL</u>	<u>A201379</u>	<u>001</u>	Feb 28, 2017

POWDER; INTRA-ARTERIAL

HEPZATO

+! DELCATH SYSTEMS INC EQ 50MG BASE/VIAL N201848 001 Aug 14, 2023

PRESCRIPTION DRUG PRODUCT LIST

MELPHALAN HYDROCHLORIDE

POWDER; INTRAVENOUS

EVOMELA

+! ACROTECH BIOPHARMA EQ 50MG BASE/VIAL N207155 001 Mar 10, 2016

SOLUTION; INTRAVENOUS

MELPHALAN HYDROCHLORIDE

+! APOTEX EQ 90MG BASE/ML (EQ 90MG BASE/ML) N217110 001 Aug 18, 2023

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>7MG</u>	<u>A205825</u>	<u>001</u>	Oct 12, 2016
<u>AB</u>		<u>14MG</u>	<u>A205825</u>	<u>002</u>	Oct 12, 2016
<u>AB</u>		<u>21MG</u>	<u>A205825</u>	<u>003</u>	Oct 12, 2016
<u>AB</u>		<u>28MG</u>	<u>A205825</u>	<u>004</u>	Oct 12, 2016
<u>AB</u>	APOTEX	<u>7MG</u>	<u>A206135</u>	<u>001</u>	Nov 22, 2016
<u>AB</u>		<u>14MG</u>	<u>A206135</u>	<u>002</u>	Nov 22, 2016
<u>AB</u>		<u>21MG</u>	<u>A206135</u>	<u>003</u>	Nov 22, 2016
<u>AB</u>		<u>28MG</u>	<u>A206135</u>	<u>004</u>	Nov 22, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>7MG</u>	<u>A214651</u>	<u>001</u>	Aug 09, 2021
<u>AB</u>		<u>14MG</u>	<u>A214651</u>	<u>002</u>	Aug 09, 2021
<u>AB</u>		<u>21MG</u>	<u>A214651</u>	<u>003</u>	Aug 09, 2021
<u>AB</u>		<u>28MG</u>	<u>A214651</u>	<u>004</u>	Aug 09, 2021
<u>AB</u>	LUPIN LTD	<u>7MG</u>	<u>A206028</u>	<u>001</u>	Sep 28, 2016
<u>AB</u>		<u>14MG</u>	<u>A206028</u>	<u>002</u>	Sep 28, 2016
<u>AB</u>		<u>21MG</u>	<u>A206028</u>	<u>003</u>	Sep 28, 2016
<u>AB</u>		<u>28MG</u>	<u>A206028</u>	<u>004</u>	Sep 28, 2016
<u>AB</u>	XIAMEN LP PHARM CO	<u>7MG</u>	<u>A213985</u>	<u>001</u>	Oct 11, 2022
<u>AB</u>		<u>14MG</u>	<u>A213985</u>	<u>002</u>	Oct 11, 2022
<u>AB</u>		<u>28MG</u>	<u>A213985</u>	<u>003</u>	Oct 11, 2022
<u>AB</u>	YICHANG HUMANWELL	<u>7MG</u>	<u>A211100</u>	<u>001</u>	Apr 02, 2021
<u>AB</u>		<u>14MG</u>	<u>A211100</u>	<u>002</u>	Apr 02, 2021
<u>AB</u>		<u>21MG</u>	<u>A211100</u>	<u>003</u>	Apr 02, 2021
<u>AB</u>		<u>28MG</u>	<u>A211100</u>	<u>004</u>	Apr 02, 2021
<u>AB</u>	ZYDUS PHARMS	<u>7MG</u>	<u>A203293</u>	<u>001</u>	Aug 03, 2017
<u>AB</u>		<u>14MG</u>	<u>A203293</u>	<u>002</u>	Aug 03, 2017
<u>AB</u>		<u>21MG</u>	<u>A203293</u>	<u>003</u>	Aug 03, 2017
<u>AB</u>		<u>28MG</u>	<u>A203293</u>	<u>004</u>	Aug 03, 2017
<u>NAMENDA XR</u>					
<u>AB</u>	+ ABBVIE	<u>14MG</u>	<u>N022525</u>	<u>002</u>	Jun 21, 2010
<u>AB</u>	+	<u>21MG</u>	<u>N022525</u>	<u>003</u>	Jun 21, 2010
<u>AB</u>	+!	<u>28MG</u>	<u>N022525</u>	<u>004</u>	Jun 21, 2010

SOLUTION; ORAL

MEMANTINE HYDROCHLORIDE

<u>AA</u>	APOTEX	<u>2MG/ML</u>	<u>A209955</u>	<u>001</u>	Feb 09, 2018
<u>AA</u>	! CHARTWELL MOLECULAR	<u>2MG/ML</u>	<u>A204033</u>	<u>001</u>	Oct 13, 2015
<u>AA</u>	MACLEODS PHARMS LTD	<u>2MG/ML</u>	<u>A202790</u>	<u>001</u>	Oct 13, 2015
<u>AA</u>	SETON PHARMS	<u>2MG/ML</u>	<u>A210319</u>	<u>001</u>	Aug 31, 2020

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>5MG</u>	<u>A206528</u>	<u>001</u>	Nov 30, 2015
<u>AB</u>		<u>10MG</u>	<u>A206528</u>	<u>002</u>	Nov 30, 2015
<u>AB</u>	ALEMBIC	<u>5MG</u>	<u>A200891</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A200891</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A090041</u>	<u>001</u>	Apr 10, 2015
<u>AB</u>		<u>10MG</u>	<u>A090041</u>	<u>002</u>	Apr 10, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A203175</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A203175</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	CADILA	<u>5MG</u>	<u>A090961</u>	<u>001</u>	Jul 10, 2017
<u>AB</u>		<u>10MG</u>	<u>A090961</u>	<u>002</u>	Jul 10, 2017
<u>AB</u>	CELLTRION	<u>5MG</u>	<u>A090073</u>	<u>001</u>	Sep 04, 2015
<u>AB</u>		<u>10MG</u>	<u>A090073</u>	<u>002</u>	Sep 04, 2015
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A090048</u>	<u>001</u>	Apr 14, 2010
<u>AB</u>		<u>10MG</u>	<u>A090048</u>	<u>002</u>	Apr 14, 2010
<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A204389</u>	<u>001</u>	Sep 26, 2022
<u>AB</u>		<u>10MG</u>	<u>A204389</u>	<u>002</u>	Sep 26, 2022
<u>AB</u>	LUPIN LTD	<u>5MG</u>	<u>A090051</u>	<u>001</u>	Apr 10, 2015
<u>AB</u>		<u>10MG</u>	<u>A090051</u>	<u>002</u>	Apr 10, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A202840</u>	<u>001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A202840</u>	<u>002</u>	Oct 13, 2015
<u>AB</u>	POLYGEN PHARMS	<u>5MG</u>	<u>A210587</u>	<u>001</u>	Dec 11, 2020

PRESCRIPTION DRUG PRODUCT LIST

MEMANTINE HYDROCHLORIDE

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>		<u>10MG</u>	<u>A210587 002</u>	Dec 11, 2020
<u>AB</u>	PURACAP PHARM LLC	<u>5MG</u>	<u>A206855 001</u>	Nov 17, 2015
<u>AB</u>		<u>10MG</u>	<u>A206855 002</u>	Nov 17, 2015
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A202350 001</u>	May 23, 2017
<u>AB</u>		<u>10MG</u>	<u>A202350 002</u>	May 23, 2017
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A090058 001</u>	May 05, 2010
<u>AB</u>		<u>10MG</u>	<u>A090058 002</u>	May 05, 2010
<u>AB</u>	UNICHEM	<u>5MG</u>	<u>A200022 001</u>	Oct 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A200022 002</u>	Oct 13, 2015
<u>AB</u>	UPSHER SMITH LABS	<u>5MG</u>	<u>A090043 001</u>	Jul 31, 2015
<u>AB</u>		<u>10MG</u>	<u>A090043 002</u>	Jul 31, 2015
<u>AB</u>	WESTMINSTER PHARMS	<u>5MG</u>	<u>A209527 001</u>	May 07, 2018
<u>AB</u>		<u>10MG</u>	<u>A209527 002</u>	May 07, 2018

NAMENDA

<u>AB</u>	+ ABBEVIE	<u>5MG</u>	<u>N021487 001</u>	Oct 16, 2003
<u>AB</u>	+!	<u>10MG</u>	<u>N021487 002</u>	Oct 16, 2003

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

<u>AP</u>	+! HOSPIRA	<u>25MG/ML</u>	<u>N021171 001</u>	
<u>AP</u>	+!	<u>50MG/ML</u>	<u>N021171 002</u>	
<u>AP</u>	+!	<u>75MG/ML</u>	<u>N021171 003</u>	
<u>AP</u>	+!	<u>100MG/ML</u>	<u>N021171 004</u>	

MEPERIDINE HYDROCHLORIDE

<u>AP</u>	WEST-WARD PHARMS INT	<u>25MG/ML</u>	<u>A080445 001</u>	
<u>AP</u>		<u>25MG/ML</u>	<u>A080455 007</u>	
<u>AP</u>		<u>50MG/ML</u>	<u>A080445 002</u>	
<u>AP</u>		<u>50MG/ML</u>	<u>A080455 008</u>	
<u>AP</u>		<u>75MG/ML</u>	<u>A080445 003</u>	
<u>AP</u>		<u>75MG/ML</u>	<u>A080455 009</u>	
<u>AP</u>		<u>100MG/ML</u>	<u>A080445 004</u>	
<u>AP</u>		<u>100MG/ML</u>	<u>A080455 010</u>	

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

!	WEST-WARD PHARMS INT	10MG/ML	A081002 001	Jul 30, 1993
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SYRUP; ORAL

MEPERIDINE HYDROCHLORIDE

!	HIKMA	50MG/5ML	A088744 001	Jan 30, 1985
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TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

	EPIC PHARMA LLC	50MG	A040331 001	May 28, 1999
		100MG	A040331 002	May 28, 1999

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE

<u>AP</u>	+! HOSPIRA	<u>1%</u>	<u>N012250 001</u>	
<u>AP</u>	+!	<u>1.5%</u>	<u>N012250 005</u>	
<u>AP</u>	+!	<u>2%</u>	<u>N012250 002</u>	

POLOCAINE

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A089407 001</u>	Dec 01, 1986
<u>AP</u>		<u>2%</u>	<u>A089410 001</u>	Dec 01, 1986

POLOCAINE-MPF

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A089406 001</u>	Dec 01, 1986
<u>AP</u>		<u>1.5%</u>	<u>A089408 001</u>	Dec 01, 1986
<u>AP</u>		<u>2%</u>	<u>A089409 001</u>	Dec 01, 1986

SCANDONEST PLAIN

!	DEPROCO	3%	A088387 001	Oct 10, 1984
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MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

<u>AA</u>	! ALEMBIC PHARMS LTD	<u>200MG</u>	<u>A090122 001</u>	Feb 18, 2009
<u>AA</u>	!	<u>400MG</u>	<u>A090122 002</u>	Feb 18, 2009
<u>AA</u>	INVAGEN PHARMS	<u>200MG</u>	<u>A040797 001</u>	Feb 27, 2008
<u>AA</u>		<u>400MG</u>	<u>A040797 002</u>	Feb 27, 2008

PRESCRIPTION DRUG PRODUCT LIST

MERCAPTOPURINE

SUSPENSION; ORAL

PURIXAN

+! NOVA LABS LTD 20MG/ML

N205919 001 Apr 28, 2014

TABLET; ORAL

MERCAPTOPURINE**AB** DR REDDYS LABS SA **50MG****A040461 001** Feb 11, 2004**AB** ! HIKMA **50MG****A040528 001** Feb 13, 2004**AB** MYLAN **50MG****A040594 001** Jul 01, 2005PURINETHOL**AB** + STASON PHARMS **50MG****N009053 002**MEROPENEM

INJECTABLE; INJECTION

MEROPENEM**AP** ACS DOBFAR **500MG/VIAL****A091404 001** Oct 26, 2011**AP** **1GM/VIAL****A091404 002** Oct 26, 2011**AP** ACS DOBFAR SPA **500MG/VIAL****A204139 001** Jun 09, 2016**AP** **1GM/VIAL****A204139 002** Jun 09, 2016**AP** AMNEAL PHARMS **500MG/VIAL****A205883 001** Apr 12, 2016**AP** **1GM/VIAL****A205883 002** Apr 12, 2016**AP** BROOKS STERISCIENCE **500MG/VIAL****A216154 001** Aug 18, 2022**AP** **1GM/VIAL****A216154 002** Aug 18, 2022**AP** DAEWOONG PHARM CO **500MG/VIAL****A204854 001** Dec 18, 2015**AP** **1GM/VIAL****A204854 002** Dec 18, 2015**AP** EUGIA PHARMA **500MG/VIAL****A205835 001** Mar 27, 2017**AP** **1GM/VIAL****A205835 002** Mar 27, 2017**AP** GLAND **500MG/VIAL****A206141 001** Jun 08, 2016**AP** **1GM/VIAL****A206141 002** Jun 08, 2016**AP** HQ SPCLT PHARMA **500MG/VIAL****A210773 001** Aug 16, 2019**AP** **1GM/VIAL****A210773 002** Aug 16, 2019**AP** SAVIOR LIFETEC CORP **500MG/VIAL****A206086 001** Apr 19, 2016**AP** **1GM/VIAL****A206086 002** Apr 19, 2016MERREM**AP** +! PFIZER **500MG/VIAL****N050706 003** Jun 21, 1996**AP** +! **1GM/VIAL****N050706 001** Jun 21, 1996

POWDER; INTRAVENOUS

MEROPENEM

+! HQ SPCLT PHARMA 2GM/VIAL

N215212 001 Jul 26, 2023

MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER

B BRAUN MEDICAL INC 500MG/VIAL

N202106 001 Apr 30, 2015

1GM/VIAL

N202106 002 Apr 30, 2015

MEROPENEM; VABORBACTAM

POWDER; INTRAVENOUS

VABOMERE

+! REMPEX 1GM/VIAL; 1GM/VIAL

N209776 001 Aug 29, 2017

MESALAMINE

CAPSULE, DELAYED RELEASE; ORAL

DELZICOL**AB** +! ABBVIE **400MG****N204412 001** Feb 01, 2013MESALAMINE**AB** TEVA PHARMS USA **400MG****A207873 001** May 09, 2019

CAPSULE, EXTENDED RELEASE; ORAL

APRISO**AB** +! SALIX **375MG****N022301 001** Oct 31, 2008MESALAMINE**AB** ALEMBIC **375MG****A216967 001** Oct 28, 2022**AB** ALKEM LABS LTD **375MG****A214242 001** Jul 15, 2021**AB** AMTA **375MG****A217533 001** Jun 06, 2023**AB** AUROBINDO PHARMA**A214477 001** Mar 31, 2023

LTD

AB MYLAN **375MG****A207271 001** Nov 20, 2019**AB** SUN PHARM **500MG****A214585 001** May 11, 2022**AB** ZYDUS PHARMS **375MG****A208954 001** Aug 12, 2021PENTASA**AB** +! TAKEDA PHARMS USA **500MG****N020049 002** Jul 08, 2004

+ 250MG

N020049 001 May 10, 1993

ENEMA; RECTAL

MESALAMINE**AB** ENCUBE **4GM/60ML****A216941 001** May 30, 2023**AB** PADAGIS ISRAEL **4GM/60ML****A076751 001** Sep 17, 2004

PRESCRIPTION DRUG PRODUCT LIST

MESALAMINE

ENEMA; RECTAL

ROWASA

AB	+!	MYLAN SPECIALITY LP	4GM/60ML	N019618	001	Dec 24, 1987
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SFROWASA

AB	+	MYLAN SPECIALITY LP	4GM/60ML	N019618	002	Jun 20, 2008
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SUPPOSITORY; RECTAL

CANASA

AB	+!	ABEVIE	1GM	N021252	002	Nov 05, 2004
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MESALAMINE

AB		ACTAVIS MID	1GM	A205654	001	Aug 14, 2020
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ATLANTIC

AB		AMRING PHARMS	1GM	A208362	001	Jun 21, 2019
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AB		ANNORA PHARMA	1GM	A213377	001	Mar 19, 2020
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AB		MYLAN	1GM	A204354	001	Nov 24, 2015
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AB		PHARM SOURCING	1GM	A207448	001	Apr 19, 2019
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AB		SANDOZ	1GM	A202065	001	Jun 12, 2019
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AB		ZYDUS PHARMS	1GM	A208953	001	Feb 12, 2020
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TABLET, DELAYED RELEASE; ORAL

LIALDA

AB	+!	TAKEDA PHARMS USA	1.2GM	N022000	001	Jan 16, 2007
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MESALAMINE

AB		ACTAVIS LABS FL	1.2GM	A203817	001	Mar 23, 2018
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AB		SINOTHERAPEUTICS	1.2GM	A217337	001	May 12, 2023
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INC

AB		SUN PHARM	1.2GM	A211858	001	Jan 25, 2019
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AB		ZYDUS PHARMS	1.2GM	A091640	001	Jun 05, 2017
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!

			800MG	A203286	001	Jul 21, 2017
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MESNA

INJECTABLE; INTRAVENOUS

MESNA

AP		FRESENIUS KABI USA	100MG/ML	A075811	001	Apr 26, 2001
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AP		GLAND	100MG/ML	A206992	001	Dec 18, 2017
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AP		HIKMA	100MG/ML	A075739	001	Jan 09, 2004
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AP		SAGENT PHARMS INC	100MG/ML	A090913	001	Apr 13, 2010
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MESNEX

AP	+!	BAXTER HLTHCARE	100MG/ML	N019884	001	Dec 30, 1988
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TABLET; ORAL

MESNEX

+!		BAXTER HLTHCARE	400MG	N020855	001	Mar 21, 2002
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METAPROTERENOL SULFATE

SYRUP; ORAL

METAPROTERENOL SULFATE

!		ANDA REPOSITORY	10MG/5ML	A073632	001	Jul 22, 1992
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METARAMINOL BITARTRATE

INJECTABLE; INJECTION

METARAMINOL BITARTRATE

		SLAYBACK PHARMA LLC	EQ 10MG BASE/ML	A211304	001	Aug 24, 2021
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METAXALONE

TABLET; ORAL

METAXALONE

AB		ACTAVIS LABS FL INC	800MG	A203695	001	Jun 15, 2017
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AB	!	AMNEAL PHARMS	800MG	A203399	001	Jun 21, 2013
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AB		LANNETT CO INC	800MG	A204770	001	Nov 22, 2016
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AB		MOUNTAIN	400MG	A040486	001	Feb 27, 2015
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AB		SANDOZ	800MG	A040445	001	Mar 31, 2010
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AB		SCIEGEN PHARMS INC	400MG	A207466	002	Mar 13, 2020
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AB			800MG	A207466	001	Aug 31, 2017
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BX		APPCO	800MG	A208774	001	Sep 24, 2018
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METFORMIN HYDROCHLORIDE

SOLUTION; ORAL

METFORMIN HYDROCHLORIDE

AB	!	SAPTALIS PHARMS	500MG/5ML	A211309	001	Mar 03, 2020
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AB		VISTAPHARM	500MG/5ML	A212677	001	Aug 19, 2022
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TABLET; ORAL

METFORMIN HYDROCHLORIDE

AB		ALKEM	500MG	A091184	001	Nov 01, 2010
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AB			850MG	A091184	002	Nov 01, 2010
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AB			1GM	A091184	003	Nov 01, 2010
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AB		AMNEAL PHARMS NY	500MG	A077880	001	Jun 05, 2006
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PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>		<u>850MG</u>	<u>A077880</u>	<u>002</u>	Jun 05, 2006
<u>AB</u>		<u>1GM</u>	<u>A077880</u>	<u>003</u>	Jun 05, 2006
<u>AB</u>	APOTEX	<u>500MG</u>	<u>A075984</u>	<u>001</u>	Apr 23, 2002
<u>AB</u>		<u>500MG</u>	<u>A090666</u>	<u>001</u>	Dec 07, 2011
<u>AB</u>		<u>850MG</u>	<u>A075984</u>	<u>002</u>	Apr 23, 2002
<u>AB</u>		<u>850MG</u>	<u>A090666</u>	<u>002</u>	Dec 07, 2011
<u>AB</u>		<u>1GM</u>	<u>A075984</u>	<u>003</u>	Apr 23, 2002
<u>AB</u>		<u>1GM</u>	<u>A090666</u>	<u>003</u>	Dec 07, 2011
<u>AB</u>	ATLAS PHARMS LLC	<u>500MG</u>	<u>A076033</u>	<u>001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A076033</u>	<u>002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A076033</u>	<u>003</u>	Jan 24, 2002
<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>A077095</u>	<u>001</u>	Jan 14, 2005
<u>AB</u>		<u>850MG</u>	<u>A077095</u>	<u>002</u>	Jan 14, 2005
<u>AB</u>		<u>1GM</u>	<u>A077095</u>	<u>003</u>	Jan 14, 2005
<u>AB</u>	CHARTWELL	<u>500MG</u>	<u>A075972</u>	<u>001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A075972</u>	<u>002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A075972</u>	<u>003</u>	Jan 24, 2002
<u>AB</u>	CSPC OUYI	<u>500MG</u>	<u>A205096</u>	<u>001</u>	Jul 11, 2016
<u>AB</u>		<u>850MG</u>	<u>A205096</u>	<u>002</u>	Jul 11, 2016
<u>AB</u>		<u>1GM</u>	<u>A205096</u>	<u>003</u>	Jul 11, 2016
<u>AB</u>	DR REDDYS LABS INC	<u>500MG</u>	<u>A077787</u>	<u>001</u>	Aug 23, 2006
<u>AB</u>		<u>850MG</u>	<u>A077787</u>	<u>002</u>	Aug 23, 2006
<u>AB</u>		<u>1GM</u>	<u>A077787</u>	<u>003</u>	Aug 23, 2006
<u>AB</u>	EPIC PHARMA LLC	<u>500MG</u>	<u>A075965</u>	<u>001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075965</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075965</u>	<u>003</u>	Jan 25, 2002
<u>AB</u>	GLENMARK GENERICS	<u>500MG</u>	<u>A078170</u>	<u>001</u>	May 23, 2008
<u>AB</u>		<u>850MG</u>	<u>A078170</u>	<u>002</u>	May 23, 2008
<u>AB</u>		<u>1GM</u>	<u>A078170</u>	<u>003</u>	May 23, 2008
<u>AB</u>	GRANULES INDIA	<u>500MG</u>	<u>A090564</u>	<u>001</u>	Apr 22, 2010
<u>AB</u>		<u>850MG</u>	<u>A090564</u>	<u>002</u>	Apr 22, 2010
<u>AB</u>	!	<u>1GM</u>	<u>A090564</u>	<u>003</u>	Apr 22, 2010
<u>AB</u>	HARMAN FINOCHEM	<u>500MG</u>	<u>A213320</u>	<u>001</u>	Dec 03, 2021
<u>AB</u>		<u>850MG</u>	<u>A213320</u>	<u>002</u>	Dec 03, 2021
<u>AB</u>		<u>1GM</u>	<u>A213320</u>	<u>003</u>	Dec 03, 2021
<u>AB</u>	LAURUS	<u>500MG</u>	<u>A209882</u>	<u>001</u>	Aug 27, 2018
<u>AB</u>		<u>850MG</u>	<u>A209882</u>	<u>002</u>	Aug 27, 2018
<u>AB</u>		<u>1GM</u>	<u>A209882</u>	<u>003</u>	Aug 27, 2018
<u>AB</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090888</u>	<u>001</u>	Mar 12, 2012
<u>AB</u>		<u>850MG</u>	<u>A090888</u>	<u>002</u>	Mar 12, 2012
<u>AB</u>		<u>1GM</u>	<u>A090888</u>	<u>003</u>	Mar 12, 2012
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A075973</u>	<u>001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075973</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075973</u>	<u>003</u>	Jan 25, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A203769</u>	<u>001</u>	Sep 11, 2013
<u>AB</u>		<u>850MG</u>	<u>A203769</u>	<u>002</u>	Sep 11, 2013
<u>AB</u>		<u>1GM</u>	<u>A203769</u>	<u>003</u>	Sep 11, 2013
<u>AB</u>	ZYDUS HLTHCARE	<u>500MG</u>	<u>A203686</u>	<u>001</u>	Aug 28, 2014
<u>AB</u>		<u>850MG</u>	<u>A203686</u>	<u>002</u>	Aug 28, 2014
<u>AB</u>		<u>1GM</u>	<u>A203686</u>	<u>003</u>	Aug 28, 2014
<u>AB</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>A077064</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>850MG</u>	<u>A077064</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>		<u>1GM</u>	<u>A077064</u>	<u>003</u>	Apr 18, 2005
	CHARTWELL	625MG	A075972	005	Jan 24, 2002
		750MG	A075972	004	Jan 24, 2002

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>750MG</u>	<u>A076869</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	!	<u>750MG</u>	<u>A206145</u>	<u>002</u>	Oct 22, 2018
<u>AB</u>	AMNEAL PHARMS NY	<u>750MG</u>	<u>A078596</u>	<u>002</u>	Jan 03, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>750MG</u>	<u>A079118</u>	<u>002</u>	Jul 20, 2012
<u>AB</u>	BEXIMCO PHARMS USA	<u>750MG</u>	<u>A207427</u>	<u>002</u>	Dec 13, 2016
<u>AB</u>	CADILA	<u>750MG</u>	<u>A077078</u>	<u>001</u>	Apr 21, 2005
<u>AB</u>	CSPC OUYI	<u>750MG</u>	<u>A078321</u>	<u>002</u>	Apr 17, 2008
<u>AB</u>	GRANULES	<u>750MG</u>	<u>A209313</u>	<u>002</u>	Mar 16, 2018
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>750MG</u>	<u>A202306</u>	<u>002</u>	Feb 23, 2017
<u>AB</u>	LAURUS	<u>750MG</u>	<u>A217631</u>	<u>002</u>	Oct 05, 2023
<u>AB</u>	MACLEODS PHARMS LTD	<u>750MG</u>	<u>A206955</u>	<u>002</u>	Dec 07, 2016

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	MARKSANS PHARMA	<u>750MG</u>	<u>A090295</u>	<u>002</u>	Apr 29, 2016
<u>AB</u>	NOSTRUM PHARMS LLC	<u>750MG</u>	<u>A076756</u>	<u>002</u>	Dec 12, 2011
<u>AB</u>	PRINSTON INC	<u>750MG</u>	<u>A208880</u>	<u>002</u>	Sep 10, 2018
<u>AB</u>	SUN PHARM INDS (IN)	<u>750MG</u>	<u>A077336</u>	<u>002</u>	Feb 09, 2006
<u>AB</u>	TEVA	<u>750MG</u>	<u>A076864</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	UNICHEM	<u>750MG</u>	<u>A213359</u>	<u>002</u>	Aug 11, 2021
<u>AB</u>	YICHANG HUMANWELL	<u>750MG</u>	<u>A211052</u>	<u>002</u>	Sep 24, 2018
<u>AB1</u>	ALIGNSCIENCE PHARMA	<u>500MG</u>	<u>A209303</u>	<u>001</u>	Mar 19, 2018
<u>AB1</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A206145</u>	<u>001</u>	Oct 22, 2018
<u>AB1</u>	AMNEAL PHARMS NY	<u>500MG</u>	<u>A078596</u>	<u>001</u>	Jan 03, 2008
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A079118</u>	<u>001</u>	Jul 20, 2012
<u>AB1</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A207427</u>	<u>001</u>	Dec 13, 2016
<u>AB1</u>	CADILA	<u>500MG</u>	<u>A077060</u>	<u>001</u>	Apr 20, 2005
<u>AB1</u>	CSPC OUYI	<u>500MG</u>	<u>A078321</u>	<u>001</u>	Apr 17, 2008
<u>AB1</u>	GRANULES	<u>500MG</u>	<u>A209313</u>	<u>001</u>	Mar 16, 2018
<u>AB1</u>	INTELLIPHARMACEUTIC S	<u>500MG</u>	<u>A202306</u>	<u>001</u>	Feb 23, 2017
<u>AB1</u>	INVENTIA	<u>500MG</u>	<u>A201991</u>	<u>001</u>	Jan 18, 2012
<u>AB1</u>	LAURUS	<u>500MG</u>	<u>A217631</u>	<u>001</u>	Oct 05, 2023
<u>AB1</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A206955</u>	<u>001</u>	Dec 07, 2016
<u>AB1</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090295</u>	<u>001</u>	Apr 29, 2016
<u>AB1</u>	NOSTRUM PHARMS LLC	<u>500MG</u>	<u>A076756</u>	<u>001</u>	Jul 26, 2006
<u>AB1</u>	PRINSTON INC	<u>500MG</u>	<u>A208880</u>	<u>001</u>	Sep 10, 2018
<u>AB1</u>	SANDOZ	<u>500MG</u>	<u>A076873</u>	<u>001</u>	Dec 14, 2004
<u>AB1</u>	SUN PHARM INDS (IN)	<u>500MG</u>	<u>A077336</u>	<u>001</u>	Feb 09, 2006
<u>AB1</u>	TEVA	<u>500MG</u>	<u>A076269</u>	<u>001</u>	Jun 18, 2004
<u>AB1</u>	UNICHEM	<u>500MG</u>	<u>A213359</u>	<u>001</u>	Aug 11, 2021
<u>AB1</u>	YICHANG HUMANWELL	<u>500MG</u>	<u>A211052</u>	<u>001</u>	Sep 24, 2018

FORTAMET

<u>AB2</u>	+ ANDRX LABS LLC	<u>1GM</u>	<u>N021574</u>	<u>002</u>	Apr 27, 2004
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METFORMIN HYDROCHLORIDE

<u>AB2</u>	AJANTA PHARMA LTD	<u>500MG</u>	<u>A213651</u>	<u>001</u>	Apr 09, 2020
<u>AB2</u>		<u>1GM</u>	<u>A213651</u>	<u>002</u>	Apr 09, 2020
<u>AB2</u>	LUPIN LTD	<u>500MG</u>	<u>A090692</u>	<u>001</u>	Jun 29, 2011
<u>AB2</u>	!	<u>1GM</u>	<u>A090692</u>	<u>002</u>	Jun 29, 2011
<u>AB2</u>	MYLAN PHARMS INC	<u>500MG</u>	<u>A200690</u>	<u>001</u>	Aug 01, 2012
<u>AB2</u>		<u>1GM</u>	<u>A200690</u>	<u>002</u>	Aug 01, 2012
<u>AB2</u>	NOSTRUM LABS INC	<u>500MG</u>	<u>A203832</u>	<u>001</u>	Dec 26, 2017
<u>AB2</u>		<u>1GM</u>	<u>A203832</u>	<u>002</u>	Dec 26, 2017
<u>AB2</u>	NOVAST LABS	<u>500MG</u>	<u>A209674</u>	<u>001</u>	Nov 02, 2018
<u>AB2</u>		<u>1GM</u>	<u>A209674</u>	<u>002</u>	Nov 02, 2018
<u>AB2</u>	QINGDAO BAHEAL PHARM	<u>500MG</u>	<u>A209993</u>	<u>001</u>	Dec 27, 2018
<u>AB2</u>		<u>1GM</u>	<u>A209993</u>	<u>002</u>	Dec 27, 2018
<u>AB2</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A214629</u>	<u>001</u>	Feb 22, 2022
<u>AB2</u>		<u>1GM</u>	<u>A214629</u>	<u>002</u>	Feb 22, 2022
<u>AB2</u>	TWI PHARMS	<u>500MG</u>	<u>A213247</u>	<u>001</u>	Sep 29, 2021
<u>AB2</u>		<u>1GM</u>	<u>A213247</u>	<u>002</u>	Sep 29, 2021

GLUMETZA

<u>AB3</u>	+ SANTARUS INC	<u>500MG</u>	<u>N021748</u>	<u>001</u>	Jun 03, 2005
<u>AB3</u>	+!	<u>1GM</u>	<u>N021748</u>	<u>002</u>	Jun 03, 2005

METFORMIN HYDROCHLORIDE

<u>AB3</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A203755</u>	<u>001</u>	Aug 01, 2016
<u>AB3</u>		<u>1GM</u>	<u>A203755</u>	<u>002</u>	Aug 01, 2016
<u>AB3</u>	AJANTA PHARMA LTD	<u>500MG</u>	<u>A213962</u>	<u>001</u>	Mar 09, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213962</u>	<u>002</u>	Mar 09, 2021
<u>AB3</u>	APOTEX	<u>500MG</u>	<u>A213356</u>	<u>001</u>	Dec 13, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213356</u>	<u>002</u>	Dec 13, 2021
<u>AB3</u>	GLENMARK PHARMS LTD	<u>500MG</u>	<u>A212969</u>	<u>001</u>	Nov 25, 2019
<u>AB3</u>		<u>1GM</u>	<u>A212969</u>	<u>002</u>	Nov 25, 2019
<u>AB3</u>	GRANULES	<u>500MG</u>	<u>A213344</u>	<u>001</u>	Jan 12, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213344</u>	<u>002</u>	Jan 12, 2021
<u>AB3</u>	LUPIN LTD	<u>500MG</u>	<u>A091664</u>	<u>001</u>	Jul 19, 2013
<u>AB3</u>		<u>1GM</u>	<u>A091664</u>	<u>002</u>	Jul 19, 2013
<u>AB3</u>	MICRO LABS	<u>500MG</u>	<u>A212448</u>	<u>001</u>	Jan 10, 2023
<u>AB3</u>		<u>1GM</u>	<u>A212448</u>	<u>002</u>	Jan 10, 2023
<u>AB3</u>	PRINSTON INC	<u>500MG</u>	<u>A212681</u>	<u>001</u>	Jun 09, 2022
<u>AB3</u>		<u>1GM</u>	<u>A212681</u>	<u>002</u>	Jun 09, 2022
<u>AB3</u>	RK PHARMA	<u>500MG</u>	<u>A215629</u>	<u>001</u>	Jun 20, 2023

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

<u>AB3</u>		<u>1GM</u>	<u>A215629</u>	<u>002</u>	Jun 20, 2023
<u>AB3</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A213334</u>	<u>001</u>	Apr 16, 2021
<u>AB3</u>		<u>1GM</u>	<u>A213334</u>	<u>002</u>	Apr 16, 2021
<u>AB3</u>	SUN PHARM	<u>500MG</u>	<u>A202917</u>	<u>001</u>	Aug 01, 2016
<u>AB3</u>		<u>1GM</u>	<u>A202917</u>	<u>002</u>	Aug 01, 2016

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOPLUS MET

<u>AB</u>	+!	TAKEDA PHARMS USA	<u>850MG;EQ 15MG BASE</u>	<u>N021842</u>	<u>002</u>	Aug 29, 2005
<u>PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE</u>						
<u>AB</u>		AUROBINDO PHARMA LTD	<u>500MG;EQ 15MG BASE</u>	<u>A200823</u>	<u>001</u>	Feb 13, 2013
<u>AB</u>			<u>850MG;EQ 15MG BASE</u>	<u>A200823</u>	<u>002</u>	Feb 13, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>500MG;EQ 15MG BASE</u>	<u>A204802</u>	<u>001</u>	Nov 05, 2015
<u>AB</u>			<u>850MG;EQ 15MG BASE</u>	<u>A204802</u>	<u>002</u>	Nov 05, 2015
<u>AB</u>		TEVA PHARMS USA	<u>500MG;EQ 15MG BASE</u>	<u>A091155</u>	<u>001</u>	Mar 10, 2014
<u>AB</u>			<u>850MG;EQ 15MG BASE</u>	<u>A091155</u>	<u>002</u>	Mar 10, 2014

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

<u>AB</u>		DR REDDYS LABS SA	<u>500MG;EQ 5MG BASE</u>	<u>A207678</u>	<u>001</u>	Aug 09, 2023
<u>AB</u>			<u>1GM;EQ 2.5MG BASE</u>	<u>A207678</u>	<u>002</u>	Aug 09, 2023
<u>AB</u>	!		<u>1GM;EQ 5MG BASE</u>	<u>A207678</u>	<u>003</u>	Aug 09, 2023
<u>SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE</u>						
<u>AB</u>		MYLAN	<u>500MG;EQ 5MG BASE</u>	<u>A205981</u>	<u>001</u>	Jul 31, 2023
<u>AB</u>			<u>1GM;EQ 2.5MG BASE</u>	<u>A205981</u>	<u>002</u>	Jul 31, 2023
<u>AB</u>			<u>1GM;EQ 5MG BASE</u>	<u>A205981</u>	<u>003</u>	Jul 31, 2023
<u>AB</u>		SUN PHARM	<u>500MG;EQ 5MG BASE</u>	<u>A206081</u>	<u>001</u>	Jul 31, 2023
<u>AB</u>			<u>1GM;EQ 2.5MG BASE</u>	<u>A206081</u>	<u>002</u>	Jul 31, 2023
<u>AB</u>			<u>1GM;EQ 5MG BASE</u>	<u>A206081</u>	<u>003</u>	Jul 31, 2023

METFORMIN HYDROCHLORIDE; SITAGLIPTIN

TABLET;ORAL

ZITUVIMET

	+	ZYDUS	500MG;50MG	N216743	001	Nov 03, 2023
	+	!	1GM;50MG	N216743	002	Nov 03, 2023

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET;ORAL

JANUMET

	+	MSD SUB MERCK	500MG;EQ 50MG BASE	N022044	001	Mar 30, 2007
	+	!	1GM;EQ 50MG BASE	N022044	002	Mar 30, 2007

TABLET, EXTENDED RELEASE;ORAL

JANUMET XR

	+	MSD SUB MERCK	500MG;EQ 50MG BASE	N202270	001	Feb 02, 2012
	+		1GM;EQ 50MG BASE	N202270	002	Feb 02, 2012
	+	!	1GM;EQ 100MG BASE	N202270	003	Feb 02, 2012

METHACHOLINE CHLORIDE

FOR SOLUTION;INHALATION

PROVOCHOLINE

	+	METHAPHARM	100MG/VIAL	N019193	001	Oct 31, 1986
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METHADONE HYDROCHLORIDE

CONCENTRATE;ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>		HIKMA	<u>10MG/ML</u>	<u>A040180</u>	<u>001</u>	Apr 30, 1998
<u>AA</u>		LANNETT CO INC	<u>10MG/ML</u>	<u>A212093</u>	<u>001</u>	Nov 02, 2020
<u>AA</u>		SPECGX LLC	<u>10MG/ML</u>	<u>A207368</u>	<u>001</u>	Aug 22, 2019
<u>AA</u>		VISTAPHARM	<u>10MG/ML</u>	<u>A040088</u>	<u>001</u>	Nov 30, 1994

METHADONE HYDROCHLORIDE INTENSOL

<u>AA</u>		HIKMA	<u>10MG/ML</u>	<u>A089897</u>	<u>001</u>	Sep 06, 1988
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METHADOSE

<u>AA</u>	+	SPECGX LLC	<u>10MG/ML</u>	<u>N017116</u>	<u>002</u>	
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INJECTABLE;INJECTION

METHADONE HYDROCHLORIDE

<u>AP</u>		LONG GROVE PHARMS	<u>10MG/ML</u>	<u>A208306</u>	<u>001</u>	Oct 27, 2017
<u>AP</u>	+	MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>N021624</u>	<u>001</u>	

PRESCRIPTION DRUG PRODUCT LIST

METHADONE HYDROCHLORIDE

SOLUTION;ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	+	HIKMA	<u>5MG/5ML</u>	<u>A087393</u>	<u>001</u>	
<u>AA</u>	+		<u>10MG/5ML</u>	<u>A087997</u>	<u>001</u>	Aug 30, 1982
<u>AA</u>		SPECGX LLC	<u>5MG/5ML</u>	<u>A207537</u>	<u>001</u>	Oct 02, 2019
<u>AA</u>			<u>10MG/5ML</u>	<u>A207537</u>	<u>002</u>	Oct 01, 2019
<u>AA</u>		VISTAPHARM	<u>5MG/5ML</u>	<u>A090707</u>	<u>001</u>	Jun 30, 2010
<u>AA</u>			<u>10MG/5ML</u>	<u>A090707</u>	<u>002</u>	Jun 30, 2010

TABLET;ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>		ASCENT PHARMS INC	<u>5MG</u>	<u>A211228</u>	<u>001</u>	Jan 03, 2019
<u>AA</u>			<u>10MG</u>	<u>A211228</u>	<u>002</u>	Jan 03, 2019
<u>AA</u>		AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A203502</u>	<u>001</u>	Aug 31, 2015
<u>AA</u>			<u>10MG</u>	<u>A203502</u>	<u>002</u>	Aug 31, 2015
<u>AA</u>		EPIC PHARMA LLC	<u>5MG</u>	<u>A090065</u>	<u>001</u>	Aug 18, 2015
<u>AA</u>			<u>10MG</u>	<u>A090065</u>	<u>002</u>	Aug 18, 2015
<u>AA</u>		ROXANE	<u>5MG</u>	<u>A088108</u>	<u>001</u>	Mar 08, 1983
<u>AA</u>			<u>10MG</u>	<u>A088109</u>	<u>001</u>	Mar 08, 1983
<u>AA</u>	!	SPECGX LLC	<u>5MG</u>	<u>A040517</u>	<u>001</u>	Apr 27, 2004
<u>AA</u>	!		<u>10MG</u>	<u>A040517</u>	<u>002</u>	Apr 27, 2004
<u>AA</u>		THEPHARMANETWORK LLC	<u>5MG</u>	<u>A090635</u>	<u>002</u>	Sep 22, 2020
<u>AA</u>			<u>10MG</u>	<u>A090635</u>	<u>001</u>	Nov 25, 2009
<u>AA</u>		VISTAPHARM	<u>10MG</u>	<u>A040241</u>	<u>002</u>	May 29, 1998
<u>AA</u>			<u>10MG</u>	<u>A204166</u>	<u>001</u>	Mar 16, 2020

TABLET, FOR SUSPENSION;ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	+	HIKMA	<u>40MG</u>	<u>N017058</u>	<u>001</u>	
<u>AA</u>		SPECGX LLC	<u>40MG</u>	<u>A077142</u>	<u>001</u>	Jul 12, 2005
<u>AA</u>		VISTAPHARM	<u>40MG</u>	<u>A075082</u>	<u>001</u>	Mar 25, 1998
<u>METHADOSE</u>						
<u>AA</u>		SPECGX LLC	<u>40MG</u>	<u>A074184</u>	<u>001</u>	Apr 29, 1993

METHAMPHETAMINE HYDROCHLORIDE

TABLET;ORAL

DESOXYN

<u>AA</u>	+	AJENAT PHARMS	<u>5MG</u>	<u>N005378</u>	<u>002</u>	
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METHAMPHETAMINE HYDROCHLORIDE

<u>AA</u>		DR REDDYS LABS SA	<u>5MG</u>	<u>A091189</u>	<u>001</u>	Apr 21, 2010
<u>AA</u>		HIKMA	<u>5MG</u>	<u>A203846</u>	<u>001</u>	Nov 17, 2015

METHAZOLAMIDE

TABLET;ORAL

METHAZOLAMIDE

<u>AB</u>		ANI PHARMS	<u>25MG</u>	<u>A040001</u>	<u>001</u>	Jun 30, 1993
<u>AB</u>			<u>50MG</u>	<u>A040001</u>	<u>002</u>	Jun 30, 1993
<u>AB</u>		BAUSCH AND LOMB INC	<u>25MG</u>	<u>A207438</u>	<u>001</u>	Oct 05, 2018
<u>AB</u>			<u>50MG</u>	<u>A207438</u>	<u>002</u>	Oct 05, 2018
<u>AB</u>		MIKART	<u>25MG</u>	<u>A040062</u>	<u>001</u>	Jan 27, 1994
<u>AB</u>	!		<u>50MG</u>	<u>A040062</u>	<u>002</u>	Jan 27, 1994
<u>AB</u>		SANDOZ	<u>25MG</u>	<u>A040036</u>	<u>001</u>	Jun 30, 1993
<u>AB</u>			<u>50MG</u>	<u>A040036</u>	<u>002</u>	Jun 30, 1993
<u>AB</u>		TAGI	<u>25MG</u>	<u>A215615</u>	<u>001</u>	Oct 18, 2022
<u>AB</u>			<u>50MG</u>	<u>A215615</u>	<u>002</u>	Oct 18, 2022

METHENAMINE HIPPURATE

TABLET;ORAL

HIPREX

<u>AB</u>	+	SS PHARMA	<u>1GM</u>	<u>N017681</u>	<u>001</u>	
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METHENAMINE HIPPURATE

<u>AB</u>	!	AUROBINDO PHARMA LTD	<u>1GM</u>	<u>A205661</u>	<u>001</u>	Jul 05, 2016
<u>AB</u>		LYRUS LIFE SCIENCES	<u>1GM</u>	<u>A217675</u>	<u>001</u>	Dec 01, 2023
<u>AB</u>		MICRO LABS	<u>1GM</u>	<u>A212172</u>	<u>001</u>	Aug 01, 2019
<u>AB</u>		NOVAST LABS	<u>1GM</u>	<u>A210068</u>	<u>001</u>	Nov 27, 2020

UREX

<u>AB</u>		ALVOGEN	<u>1GM</u>	<u>N016151</u>	<u>001</u>	
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PRESCRIPTION DRUG PRODUCT LIST

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

<u>AB</u>	BIONPHARMA	<u>5MG</u>	<u>A218149 001</u>	Sep 25, 2023
<u>AB</u>		<u>10MG</u>	<u>A218149 002</u>	Sep 25, 2023
<u>AB</u>	CHARTWELL RX	<u>5MG</u>	<u>A040411 001</u>	Mar 27, 2001
<u>AB</u>		<u>10MG</u>	<u>A040411 002</u>	Mar 27, 2001
<u>AB</u>	HERITAGE PHARMA	<u>5MG</u>	<u>A040734 001</u>	Dec 14, 2007
<u>AB</u>		<u>10MG</u>	<u>A040734 002</u>	Dec 14, 2007
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A209827 001</u>	May 24, 2023
<u>AB</u>		<u>10MG</u>	<u>A209827 002</u>	May 24, 2023
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A040350 001</u>	Mar 29, 2000
<u>AB</u>	!	<u>10MG</u>	<u>A040350 002</u>	Mar 29, 2000
<u>AB</u>	QINGDAO BAHEAL PHARM	<u>5MG</u>	<u>A040547 001</u>	Feb 18, 2005
<u>AB</u>		<u>10MG</u>	<u>A040547 002</u>	Feb 18, 2005
<u>AB</u>	RISING	<u>5MG</u>	<u>A202068 001</u>	Mar 07, 2012
<u>AB</u>		<u>10MG</u>	<u>A202068 002</u>	Mar 07, 2012

METHOCARBAMOL

SOLUTION; IM-IV

METHOCARBAMOL

<u>AP</u>	AM REGENT	<u>1GM/10ML (100MG/ML)</u>	<u>A207496 001</u>	Jun 22, 2017
<u>AP</u>	EUGIA PHARMA	<u>1GM/10ML (100MG/ML)</u>	<u>A206128 001</u>	May 27, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>1GM/10ML (100MG/ML)</u>	<u>A209331 001</u>	Apr 17, 2018
<u>AP</u>	GLAND PHARMA LTD	<u>1GM/10ML (100MG/ML)</u>	<u>A211504 001</u>	Oct 26, 2018
<u>AP</u>	MONTEREY PHARMS LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A205354 001</u>	Oct 27, 2016
<u>AP</u>	NAVINTA LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A206071 001</u>	Nov 24, 2017
<u>AP</u>	SAGENT PHARMS INC	<u>1GM/10ML (100MG/ML)</u>	<u>A205404 001</u>	Jul 18, 2017
<u>AP</u>	SLATE RUN PHARMA	<u>1GM/10ML (100MG/ML)</u>	<u>A208116 001</u>	Jan 19, 2017
<u>AP</u>	SOMERSET THERAPS LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A207522 001</u>	Jul 31, 2017

ROBAXIN

<u>AP</u>	+!	HIKMA	<u>1GM/10ML (100MG/ML)</u>	<u>N011790 001</u>
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TABLET; ORAL

METHOCARBAMOL

<u>AA</u>	ACME LABS	<u>500MG</u>	<u>A203550 001</u>	Feb 08, 2017	
<u>AA</u>		<u>750MG</u>	<u>A203550 002</u>	Feb 08, 2017	
<u>AA</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A213967 002</u>	Feb 09, 2023	
<u>AA</u>		<u>750MG</u>	<u>A213967 001</u>	Aug 12, 2020	
<u>AA</u>	AUSTARPHARMA LLC	<u>500MG</u>	<u>A200958 001</u>	Oct 21, 2011	
<u>AA</u>		<u>750MG</u>	<u>A200958 002</u>	Oct 21, 2011	
<u>AA</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A208507 001</u>	Jul 21, 2017	
<u>AA</u>		<u>750MG</u>	<u>A208507 002</u>	Jul 21, 2017	
<u>AA</u>	!	GRANULES	<u>500MG</u>	<u>A209312 001</u>	May 07, 2018
<u>AA</u>	!		<u>750MG</u>	<u>A209312 002</u>	May 07, 2018
<u>AA</u>	HETERO LABS LTD III	<u>500MG</u>	<u>A090200 001</u>	Nov 06, 2009	
<u>AA</u>		<u>750MG</u>	<u>A090200 002</u>	Nov 06, 2009	
<u>AA</u>	MLV	<u>500MG</u>	<u>A212623 001</u>	Apr 30, 2021	
<u>AA</u>		<u>750MG</u>	<u>A212623 002</u>	Apr 30, 2021	
<u>AA</u>	OXFORD PHARMS	<u>500MG</u>	<u>A040489 001</u>	Jan 29, 2003	
<u>AA</u>		<u>750MG</u>	<u>A040489 002</u>	Jan 29, 2003	
<u>AA</u>	PRINSTON INC	<u>500MG</u>	<u>A086988 002</u>		
<u>AA</u>		<u>750MG</u>	<u>A086988 001</u>		
<u>AB</u>	!	AUSTARPHARMA LLC	<u>1GM</u>	<u>A200958 003</u>	Dec 06, 2021
<u>AB</u>	MIKART	<u>1GM</u>	<u>A212707 001</u>	Jun 12, 2023	

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

	+!	PAR STERILE PRODUCTS	500MG/VIAL	N011559 001
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METHOTREXATE

SOLUTION; ORAL

JYLAMVO

	+!	SHORLA	2MG/ML	N212479 001	Nov 29, 2022
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SOLUTION; SUBCUTANEOUS

OTREXUP

	+!	OTTER PHARMS	10MG/0.4ML (10MG/0.4ML)	N204824 001	Oct 11, 2013
	+!		12.5MG/0.4ML (12.5MG/0.4ML)	N204824 006	Mar 24, 2016
	+!		15MG/0.4ML (15MG/0.4ML)	N204824 002	Oct 11, 2013
	+!		17.5MG/0.4ML (17.5MG/0.4ML)	N204824 007	Mar 24, 2016
	+!		20MG/0.4ML (20MG/0.4ML)	N204824 003	Oct 11, 2013

PRESCRIPTION DRUG PRODUCT LIST

METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP

+	!		22.5MG/0.4ML (22.5MG/0.4ML)	N204824	008	Mar 24, 2016
+	!		25MG/0.4ML (25MG/0.4ML)	N204824	004	Oct 11, 2013

RASUVO

+	!	MEDEXUS	7.5MG/0.15ML (7.5MG/0.15ML)	N205776	001	Jul 10, 2014
+	!		10MG/0.20ML (10MG/0.20ML)	N205776	002	Jul 10, 2014
+	!		12.5MG/0.25ML (12.5MG/0.25ML)	N205776	003	Jul 10, 2014
+	!		15MG/0.30ML (15MG/0.30ML)	N205776	004	Jul 10, 2014
+	!		17.5MG/0.35ML (17.5MG/0.35ML)	N205776	005	Jul 10, 2014
+	!		20MG/0.4ML (20MG/0.4ML)	N205776	006	Jul 10, 2014
+	!		22.5MG/0.45ML (22.5MG/0.45ML)	N205776	007	Jul 10, 2014
+	!		25MG/0.5ML (25MG/0.5ML)	N205776	008	Jul 10, 2014
+	!		30MG/0.6ML (30MG/0.6ML)	N205776	010	Jul 10, 2014

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE PRESERVATIVE FREE

AP	!	FRESENIUS KABI USA	<u>EQ 1GM BASE/VIAL</u>	A040266	001	Feb 26, 1999
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METHOTREXATE SODIUM

AP	!	FRESENIUS KABI USA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A040263	001	Feb 26, 1999
AP	!	HIKMA	<u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u>	A089341	001	Sep 16, 1986
AP	+	HOSPIRA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	N011719	010	Dec 15, 2004

METHOTREXATE SODIUM PRESERVATIVE FREE

AP	!	ACCORD HLTHCARE	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A040767	001	Apr 30, 2007
AP	!		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A040768	001	Apr 30, 2007
AP	!		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A040716	001	Apr 30, 2007
AP	!	HIKMA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A089340	001	Sep 16, 1986
AP			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A089343	001	Sep 16, 1986
AP	+	HOSPIRA	<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	N011719	012	Apr 13, 2005
AP		PHARMACHEMIE BV	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A040843	002	Jan 11, 2010
AP			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A040843	004	Jan 11, 2010
AP			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A040843	001	Jan 11, 2010
AP		SAGENT PHARMS INC	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A203407	001	Aug 09, 2018
AP			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A203407	002	Aug 09, 2018
AP			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A203407	003	Aug 09, 2018
AP		SANDOZ	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A090039	001	Mar 31, 2009
AP			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A090039	002	Mar 31, 2009
AP			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A090029	001	Mar 31, 2009

METHOTREXATE SODIUM

!	!	HIKMA	EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A089342	001	Sep 16, 1986
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METHOTREXATE SODIUM PRESERVATIVE FREE

!	!	HIKMA	EQ 1GM BASE/VIAL	A040632	001	Aug 12, 2005
	!	PHARMACHEMIE BV	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A040843	003	Feb 27, 2012

SOLUTION; ORAL

XATMEP

+	!	AZURITY	EQ 2.5MG BASE/ML	N208400	001	Apr 25, 2017
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TABLET; ORAL

METHOTREXATE SODIUM

AB		ACCORD HLTHCARE	<u>EQ 2.5MG BASE</u>	A213343	001	Jan 24, 2020
AB		AMNEAL PHARMS	<u>EQ 2.5MG BASE</u>	A210040	001	Dec 22, 2017
AB		BARR	<u>EQ 2.5MG BASE</u>	A081099	001	Oct 15, 1990
AB		DAITO	<u>EQ 2.5MG BASE</u>	A213362	001	Aug 07, 2023
AB		EUGIA PHARMA	<u>EQ 2.5MG BASE</u>	A210454	001	Jan 30, 2020
AB	!	HIKMA	<u>EQ 2.5MG BASE</u>	A040054	001	Aug 01, 1994
AB		LOTUS PHARM CO LTD	<u>EQ 2.5MG BASE</u>	A209787	001	Apr 23, 2021
AB		MYLAN	<u>EQ 2.5MG BASE</u>	A081235	001	May 15, 1992
AB		SUN PHARM	<u>EQ 2.5MG BASE</u>	A201749	001	May 21, 2015
AB		ZYDUS PHARMS	<u>EQ 2.5MG BASE</u>	A207812	001	Jan 13, 2017

TREXALL

		BARR	EQ 5MG BASE	A040385	001	Mar 21, 2001
			EQ 7.5MG BASE	A040385	002	Mar 21, 2001
			EQ 10MG BASE	A040385	003	Mar 21, 2001
		!	EQ 15MG BASE	A040385	004	Mar 21, 2001

METHOXSALEN

CAPSULE; ORAL

METHOXSALEN

AB		STRIDES PHARMA	<u>10MG</u>	A202687	001	Jun 05, 2014
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OXSORALEN-ULTRA

AB	+	BAUSCH	<u>10MG</u>	N019600	001	Oct 30, 1986
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PRESCRIPTION DRUG PRODUCT LIST

METHOXSALLEN

INJECTABLE; INJECTION

UVADEX

+! MALLINCKRODT HOSP 0.02MG/ML N020969 001 Feb 25, 1999

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDEAA BAYSHORE PHARMS LLC 2.5MG A200602 001 Sep 24, 2012AA ! 5MG A200602 002 Sep 24, 2012AA BRECKENRIDGE PHARM 2.5MG A040642 001 Dec 06, 2011AA 5MG A040642 002 Dec 06, 2011METHSUXIMIDE

CAPSULE; ORAL

CELONTINAB +! PARKE DAVIS 300MG N010596 008METHSUXIMIDEAB NOVITIUM PHARMA 300MG A217213 001 May 01, 2023METHYLENE BLUE

SOLUTION; INTRAVENOUS

METHYLENE BLUEAP ZYDUS 10MG/2ML (5MG/ML) A215636 001 Dec 05, 2023AP 50MG/10ML (5MG/ML) A215636 002 Dec 05, 2023PROVAYBLUEAP +! PROVEPHARM SAS 10MG/2ML (5MG/ML) N204630 002 Jul 18, 2019AP +! 50MG/10ML (5MG/ML) N204630 001 Apr 08, 2016METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHERGINEAP +! EDISON THERAPS LLC 0.2MG/ML N006035 004METHYLERGONOVINE MALEATEAP AM REGENT 0.2MG/ML A090193 001 Nov 24, 2008AP BRECKENRIDGE 0.2MG/ML A040889 001 Sep 13, 2010

TABLET; ORAL

METHYLERGONOVINE MALEATEAB AMNEAL PHARMS 0.2MG A211483 001 Sep 10, 2018AB ! CHARTWELL RX 0.2MG A091577 001 May 02, 2011AB GRANULES 0.2MG A210424 001 May 15, 2018AB RISING 0.2MG A211919 001 Jan 15, 2021AB TEVA PHARMS USA 0.2MG A211455 001 Mar 20, 2019AB TULEX PHARMS INC 0.2MG A212233 001 May 01, 2020METHYLNALTREXONE BROMIDE

SOLUTION; SUBCUTANEOUS

RELISTOR

+! SALIX PHARMS 8MG/0.4ML (8MG/0.4ML) N021964 002 Sep 27, 2010

+! 12MG/0.6ML (12MG/0.6ML) N021964 001 Apr 24, 2008

+! 12MG/0.6ML (12MG/0.6ML) N021964 003 Sep 27, 2010

TABLET; ORAL

RELISTOR

+! SALIX 150MG N208271 001 Jul 19, 2016

METHYLPHENIDATE

FILM, EXTENDED RELEASE; TRANSDERMAL

DAYTRANAAB + NOVEN PHARMS INC 10MG/9HR (1.1MG/HR) N021514 001 Apr 06, 2006AB + 15MG/9HR (1.6MG/HR) N021514 002 Apr 06, 2006AB + 20MG/9HR (2.2MG/HR) N021514 003 Apr 06, 2006AB +! 30MG/9HR (3.3MG/HR) N021514 004 Apr 06, 2006METHYLPHENIDATEAB MYLAN TECH VIATRIS 10MG/9HR (1.1MG/HR) A206497 001 Mar 14, 2022AB 15MG/9HR (1.6MG/HR) A206497 002 Mar 14, 2022AB 20MG/9HR (2.2MG/HR) A206497 003 Mar 14, 2022AB 30MG/9HR (3.3MG/HR) A206497 004 Mar 14, 2022

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE; ORAL

COTEMPLA XR-ODT

+ NEOS THERAPS INC 8.6MG N205489 001 Jun 19, 2017

+ 17.3MG N205489 002 Jun 19, 2017

+! 25.9MG N205489 003 Jun 19, 2017

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB1</u>	DR REDDYS LABS SA	<u>10MG</u>	<u>A200886 001</u>	Feb 26, 2018
<u>AB1</u>		<u>20MG</u>	<u>A078458 001</u>	Dec 01, 2011
<u>AB1</u>		<u>30MG</u>	<u>A078458 002</u>	Dec 01, 2011
<u>AB1</u>		<u>40MG</u>	<u>A078458 003</u>	Dec 01, 2011
<u>AB1</u>	!	<u>60MG</u>	<u>A078458 004</u>	Jun 23, 2016
<u>AB1</u>	GRANULES	<u>10MG</u>	<u>A211796 001</u>	May 23, 2019
<u>AB1</u>		<u>20MG</u>	<u>A211796 002</u>	May 23, 2019
<u>AB1</u>		<u>30MG</u>	<u>A211796 003</u>	May 23, 2019
<u>AB1</u>		<u>40MG</u>	<u>A211796 004</u>	May 23, 2019
<u>AB1</u>		<u>60MG</u>	<u>A211796 005</u>	May 23, 2019

RITALIN LA

<u>AB1</u>	+	SANDOZ	<u>10MG</u>	<u>N021284 004</u>	Apr 10, 2004
<u>AB1</u>	+		<u>20MG</u>	<u>N021284 001</u>	Jun 05, 2002
<u>AB1</u>	+		<u>30MG</u>	<u>N021284 002</u>	Jun 05, 2002
<u>AB1</u>	+		<u>40MG</u>	<u>N021284 003</u>	Jun 05, 2002

METADATE CD

<u>AB2</u>	+	AYTU BIOPHARMA	<u>10MG</u>	<u>N021259 003</u>	May 27, 2003
<u>AB2</u>	+		<u>20MG</u>	<u>N021259 001</u>	Apr 03, 2001
<u>AB2</u>	+		<u>30MG</u>	<u>N021259 002</u>	Jun 19, 2003
<u>AB2</u>	+		<u>40MG</u>	<u>N021259 004</u>	Feb 19, 2006
<u>AB2</u>	+		<u>50MG</u>	<u>N021259 005</u>	Feb 19, 2006
<u>AB2</u>	+		<u>60MG</u>	<u>N021259 006</u>	Feb 19, 2006

METHYLPHENIDATE HYDROCHLORIDE

<u>AB2</u>	IMPAX LABS INC	<u>10MG</u>	<u>A205105 001</u>	Jul 28, 2016
<u>AB2</u>		<u>20MG</u>	<u>A205105 002</u>	Jul 28, 2016
<u>AB2</u>		<u>30MG</u>	<u>A205105 003</u>	Jul 28, 2016
<u>AB2</u>		<u>40MG</u>	<u>A205105 004</u>	Jul 28, 2016
<u>AB2</u>		<u>50MG</u>	<u>A205105 005</u>	Jul 28, 2016
<u>AB2</u>		<u>60MG</u>	<u>A205105 006</u>	Jul 28, 2016
<u>AB2</u>	SPECGX LLC	<u>10MG</u>	<u>A203583 001</u>	Sep 29, 2015
<u>AB2</u>		<u>20MG</u>	<u>A203583 002</u>	Sep 29, 2015
<u>AB2</u>		<u>30MG</u>	<u>A203583 003</u>	Sep 29, 2015
<u>AB2</u>		<u>40MG</u>	<u>A203583 004</u>	Sep 29, 2015
<u>AB2</u>		<u>50MG</u>	<u>A203583 005</u>	Sep 29, 2015
<u>AB2</u>		<u>60MG</u>	<u>A203583 006</u>	Sep 29, 2015
<u>AB2</u>	TEVA PHARMS	<u>10MG</u>	<u>A077707 001</u>	Jul 19, 2012
<u>AB2</u>		<u>20MG</u>	<u>A077707 002</u>	Jul 19, 2012
<u>AB2</u>		<u>30MG</u>	<u>A077707 003</u>	Jul 19, 2012
<u>AB2</u>		<u>40MG</u>	<u>A078873 001</u>	Jul 19, 2012
<u>AB2</u>		<u>50MG</u>	<u>A078873 002</u>	Jul 19, 2012
<u>AB2</u>		<u>60MG</u>	<u>A078873 003</u>	Jul 19, 2012

APTENSIO XR

<u>AB3</u>	+	RHODES PHARMS	<u>10MG</u>	<u>N205831 001</u>	Apr 17, 2015
<u>AB3</u>	+		<u>15MG</u>	<u>N205831 002</u>	Apr 17, 2015
<u>AB3</u>	+		<u>20MG</u>	<u>N205831 003</u>	Apr 17, 2015
<u>AB3</u>	+		<u>30MG</u>	<u>N205831 004</u>	Apr 17, 2015
<u>AB3</u>	+		<u>40MG</u>	<u>N205831 005</u>	Apr 17, 2015
<u>AB3</u>	+		<u>50MG</u>	<u>N205831 006</u>	Apr 17, 2015
<u>AB3</u>	+		<u>60MG</u>	<u>N205831 007</u>	Apr 17, 2015

METHYLPHENIDATE HYDROCHLORIDE

<u>AB3</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A208861 001</u>	Dec 13, 2018
<u>AB3</u>		<u>15MG</u>	<u>A208861 002</u>	Dec 13, 2018
<u>AB3</u>		<u>20MG</u>	<u>A208861 003</u>	Dec 13, 2018
<u>AB3</u>		<u>30MG</u>	<u>A208861 004</u>	Dec 13, 2018
<u>AB3</u>		<u>40MG</u>	<u>A208861 005</u>	Dec 13, 2018
<u>AB3</u>		<u>50MG</u>	<u>A208861 006</u>	Dec 13, 2018
<u>AB3</u>		<u>60MG</u>	<u>A208861 007</u>	Dec 13, 2018

JORNAY PM

+	IRONSHORE PHARMS	20MG	N209311 001	Aug 08, 2018
+		40MG	N209311 002	Aug 08, 2018
+		60MG	N209311 003	Aug 08, 2018
+		80MG	N209311 004	Aug 08, 2018
+		100MG	N209311 005	Aug 08, 2018

FOR SUSPENSION, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>5MG/ML</u>	<u>A206049 001</u>	May 17, 2018
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QUILLIVANT XR

<u>AB</u>	+	NEXTWAVE	<u>5MG/ML</u>	<u>N202100 001</u>	Sep 27, 2012
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PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

SOLUTION;ORAL

METHYLIN

<u>AA</u>	+	SPECGX LLC	<u>5MG/5ML</u>	<u>N021419</u>	<u>001</u>	Dec 19, 2002
<u>AA</u>	+		<u>10MG/5ML</u>	<u>N021419</u>	<u>002</u>	Dec 19, 2002

METHYLPHENIDATE HYDROCHLORIDE

<u>AA</u>		ABHAI LLC	<u>5MG/5ML</u>	<u>A207485</u>	<u>001</u>	Nov 18, 2016
<u>AA</u>			<u>10MG/5ML</u>	<u>A207485</u>	<u>002</u>	Nov 18, 2016
<u>AA</u>		ALKEM LABS LTD	<u>5MG/5ML</u>	<u>A211647</u>	<u>001</u>	Mar 30, 2020
<u>AA</u>			<u>10MG/5ML</u>	<u>A211647</u>	<u>002</u>	Mar 30, 2020
<u>AA</u>		ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A207417</u>	<u>001</u>	Jan 29, 2021
<u>AA</u>			<u>10MG/5ML</u>	<u>A207417</u>	<u>002</u>	Jan 29, 2021
<u>AA</u>		BRECKENRIDGE	<u>5MG/5ML</u>	<u>A201466</u>	<u>001</u>	Nov 12, 2013
<u>AA</u>			<u>10MG/5ML</u>	<u>A201466</u>	<u>002</u>	Nov 12, 2013
<u>AA</u>		NOVEL LABS INC	<u>5MG/5ML</u>	<u>A204602</u>	<u>001</u>	Aug 14, 2015
<u>AA</u>			<u>10MG/5ML</u>	<u>A204602</u>	<u>002</u>	Aug 14, 2015
<u>AA</u>		PATRIN	<u>5MG/5ML</u>	<u>A210764</u>	<u>001</u>	Apr 10, 2020
<u>AA</u>			<u>10MG/5ML</u>	<u>A210764</u>	<u>002</u>	Apr 10, 2020
<u>AA</u>		QUAGEN	<u>5MG/5ML</u>	<u>A213567</u>	<u>001</u>	Jun 04, 2020
<u>AA</u>			<u>10MG/5ML</u>	<u>A213567</u>	<u>002</u>	Jun 04, 2020
<u>AA</u>		TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A091601</u>	<u>001</u>	Jul 23, 2010
<u>AA</u>			<u>10MG/5ML</u>	<u>A091601</u>	<u>002</u>	Jul 23, 2010
<u>AA</u>		WES PHARMA INC	<u>5MG/5ML</u>	<u>A210139</u>	<u>001</u>	Oct 03, 2018
<u>AA</u>			<u>10MG/5ML</u>	<u>A210139</u>	<u>002</u>	Oct 03, 2018

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		ABHAI INC	<u>5MG</u>	<u>A206932</u>	<u>001</u>	May 11, 2017
<u>AB</u>			<u>10MG</u>	<u>A206932</u>	<u>002</u>	May 11, 2017
<u>AB</u>			<u>20MG</u>	<u>A206932</u>	<u>003</u>	May 11, 2017
<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A213936</u>	<u>001</u>	Oct 28, 2020
<u>AB</u>			<u>10MG</u>	<u>A213936</u>	<u>002</u>	Oct 28, 2020
<u>AB</u>			<u>20MG</u>	<u>A213936</u>	<u>003</u>	Oct 28, 2020
<u>AB</u>		ALKEM LABS LTD	<u>5MG</u>	<u>A211779</u>	<u>001</u>	Oct 04, 2019
<u>AB</u>			<u>10MG</u>	<u>A211779</u>	<u>002</u>	Oct 04, 2019
<u>AB</u>			<u>20MG</u>	<u>A211779</u>	<u>003</u>	Oct 04, 2019
<u>AB</u>		ASCENT PHARMS INC	<u>5MG</u>	<u>A207416</u>	<u>001</u>	Sep 22, 2015
<u>AB</u>			<u>10MG</u>	<u>A207416</u>	<u>002</u>	Sep 22, 2015
<u>AB</u>			<u>20MG</u>	<u>A207416</u>	<u>003</u>	Sep 22, 2015
<u>AB</u>		BIONPHARMA	<u>5MG</u>	<u>A209753</u>	<u>001</u>	Mar 02, 2018
<u>AB</u>			<u>10MG</u>	<u>A209753</u>	<u>002</u>	Mar 02, 2018
<u>AB</u>			<u>20MG</u>	<u>A209753</u>	<u>003</u>	Mar 02, 2018
<u>AB</u>		MOUNTAIN	<u>5MG</u>	<u>A091159</u>	<u>001</u>	Mar 12, 2014
<u>AB</u>			<u>10MG</u>	<u>A091159</u>	<u>002</u>	Mar 12, 2014
<u>AB</u>			<u>20MG</u>	<u>A091159</u>	<u>003</u>	Mar 12, 2014
<u>AB</u>		NOVEL LABS INC	<u>5MG</u>	<u>A207884</u>	<u>001</u>	Nov 13, 2015
<u>AB</u>			<u>10MG</u>	<u>A207884</u>	<u>002</u>	Nov 13, 2015
<u>AB</u>			<u>20MG</u>	<u>A207884</u>	<u>003</u>	Nov 13, 2015
<u>AB</u>		OXFORD PHARMS	<u>5MG</u>	<u>A202892</u>	<u>001</u>	Sep 23, 2014
<u>AB</u>			<u>10MG</u>	<u>A202892</u>	<u>002</u>	Sep 23, 2014
<u>AB</u>			<u>20MG</u>	<u>A202892</u>	<u>003</u>	Sep 23, 2014
<u>AB</u>		PRINSTON INC	<u>5MG</u>	<u>A212697</u>	<u>001</u>	Jul 23, 2020
<u>AB</u>			<u>10MG</u>	<u>A212697</u>	<u>002</u>	Jul 23, 2020
<u>AB</u>			<u>20MG</u>	<u>A212697</u>	<u>003</u>	Jul 23, 2020
<u>AB</u>		SPECGX LLC	<u>5MG</u>	<u>A040300</u>	<u>001</u>	Nov 27, 1998
<u>AB</u>			<u>10MG</u>	<u>A040300</u>	<u>002</u>	Nov 27, 1998
<u>AB</u>			<u>20MG</u>	<u>A040300</u>	<u>003</u>	Nov 27, 1998
<u>AB</u>		SUN PHARM INDS INC	<u>5MG</u>	<u>A090710</u>	<u>001</u>	Mar 15, 2012
<u>AB</u>			<u>10MG</u>	<u>A090710</u>	<u>002</u>	Mar 15, 2012
<u>AB</u>			<u>20MG</u>	<u>A090710</u>	<u>003</u>	Mar 15, 2012

RITALIN

<u>AB</u>	+	SANDOZ	<u>5MG</u>	<u>N010187</u>	<u>003</u>	
<u>AB</u>	+		<u>10MG</u>	<u>N010187</u>	<u>006</u>	
<u>AB</u>	+		<u>20MG</u>	<u>N010187</u>	<u>010</u>	

TABLET, CHEWABLE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>		ASCENT PHARMS INC	<u>2.5MG</u>	<u>A210354</u>	<u>001</u>	Dec 29, 2017
<u>AB</u>			<u>5MG</u>	<u>A210354</u>	<u>002</u>	Dec 29, 2017
<u>AB</u>	!		<u>10MG</u>	<u>A210354</u>	<u>003</u>	Dec 29, 2017
<u>AB</u>		RISING	<u>2.5MG</u>	<u>A205756</u>	<u>001</u>	Nov 07, 2016
<u>AB</u>			<u>5MG</u>	<u>A205756</u>	<u>002</u>	Nov 07, 2016
<u>AB</u>			<u>10MG</u>	<u>A205756</u>	<u>003</u>	Nov 07, 2016

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CONCERTA

AB	+	JANSSEN PHARMS	18MG	<u>N021121</u>	<u>001</u>	Aug 01, 2000
AB	+		27MG	<u>N021121</u>	<u>004</u>	Apr 01, 2002
AB	+		36MG	<u>N021121</u>	<u>002</u>	Aug 01, 2000
AB	+		54MG	<u>N021121</u>	<u>003</u>	Dec 08, 2000

METHYLIN ER

AB		SPECGX LLC	10MG	<u>A075629</u>	<u>001</u>	May 09, 2000
AB			20MG	<u>A075629</u>	<u>002</u>	May 09, 2000

METHYLPHENIDATE HYDROCHLORIDE

AB		ABHAI LLC	10MG	<u>A207488</u>	<u>001</u>	Jun 09, 2015
AB	!		20MG	<u>A207488</u>	<u>002</u>	Jun 09, 2015
AB		ACTAVIS LABS FL	18MG	<u>A076772</u>	<u>001</u>	Mar 22, 2018
AB			27MG	<u>A076772</u>	<u>002</u>	Mar 22, 2018
AB			36MG	<u>A076772</u>	<u>003</u>	Mar 22, 2018
AB			54MG	<u>A076655</u>	<u>001</u>	Mar 21, 2018
AB			72MG	<u>A076655</u>	<u>002</u>	Feb 28, 2022
AB		ALKEM LABS LTD	18MG	<u>A214447</u>	<u>001</u>	May 23, 2023
AB			20MG	<u>A212288</u>	<u>001</u>	Oct 06, 2020
AB			27MG	<u>A214447</u>	<u>002</u>	May 23, 2023
AB			36MG	<u>A214447</u>	<u>003</u>	May 23, 2023
AB			54MG	<u>A214447</u>	<u>004</u>	May 23, 2023
AB		ANDOR PHARMS	18MG	<u>A211918</u>	<u>001</u>	Apr 24, 2019
AB			27MG	<u>A211918</u>	<u>002</u>	Apr 24, 2019
AB			36MG	<u>A211918</u>	<u>003</u>	Apr 24, 2019
AB			54MG	<u>A211918</u>	<u>004</u>	Apr 24, 2019
AB		ASCENT PHARMS INC	18MG	<u>A211009</u>	<u>001</u>	Sep 03, 2019
AB			27MG	<u>A211009</u>	<u>002</u>	Sep 03, 2019
AB			36MG	<u>A211009</u>	<u>003</u>	Sep 03, 2019
AB			54MG	<u>A211009</u>	<u>004</u>	Sep 03, 2019
AB		AUROLIFE PHARMA LLC	18MG	<u>A206726</u>	<u>001</u>	Oct 21, 2016
AB			27MG	<u>A206726</u>	<u>002</u>	Oct 21, 2016
AB			36MG	<u>A206726</u>	<u>003</u>	Oct 21, 2016
AB			54MG	<u>A206726</u>	<u>004</u>	Oct 21, 2016
AB		DR REDDYS	18MG	<u>A213473</u>	<u>001</u>	Jul 29, 2020
AB			27MG	<u>A213473</u>	<u>002</u>	Jul 29, 2020
AB			36MG	<u>A213473</u>	<u>003</u>	Jul 29, 2020
AB			54MG	<u>A213473</u>	<u>004</u>	Jul 29, 2020
AB		GRANULES	10MG	<u>A210992</u>	<u>001</u>	Nov 21, 2018
AB			20MG	<u>A210992</u>	<u>002</u>	Nov 21, 2018
AB		OSMOTICA PHARM US	18MG	<u>A205327</u>	<u>001</u>	Jul 28, 2017
AB			27MG	<u>A205327</u>	<u>002</u>	Jul 28, 2017
AB			36MG	<u>A205327</u>	<u>003</u>	Jul 28, 2017
AB			54MG	<u>A205327</u>	<u>004</u>	Jul 28, 2017
AB	!		72MG	<u>A205327</u>	<u>005</u>	Jul 28, 2017
AB		SUN PHARM INDS INC	18MG	<u>A205135</u>	<u>001</u>	Aug 19, 2020
AB			27MG	<u>A205135</u>	<u>002</u>	Aug 19, 2020
AB			36MG	<u>A205135</u>	<u>003</u>	Aug 19, 2020
AB			54MG	<u>A205135</u>	<u>004</u>	Aug 19, 2020
AB			72MG	<u>A217229</u>	<u>001</u>	Aug 25, 2023
BX		LANNETT CO INC	18MG	A091695	001	Jul 09, 2013
BX			27MG	A091695	002	Jul 09, 2013
BX			36MG	A091695	003	Sep 23, 2013
BX			54MG	A091695	004	Sep 23, 2013
BX		SPECGX LLC	27MG	A202608	001	Dec 28, 2012
BX			36MG	A202608	002	Dec 28, 2012
BX			54MG	A202608	003	Dec 28, 2012

RELEXXII

+		OSMOTICA PHARM US	18MG	N216117	001	Jun 23, 2022
+			27MG	N216117	002	Jun 23, 2022
+			36MG	N216117	003	Jun 23, 2022
+			45MG	N216117	004	Jun 23, 2022
+			54MG	N216117	005	Jun 23, 2022
+			63MG	N216117	006	Jun 23, 2022
+	!		72MG	N216117	007	Jun 23, 2022

TABLET, EXTENDED RELEASE, CHEWABLE;ORAL

QUILLICHEW ER

+		NEXTWAVE PHARMS	20MG	N207960	001	Dec 04, 2015
+			30MG	N207960	002	Dec 04, 2015
+	!		40MG	N207960	003	Dec 04, 2015

PRESCRIPTION DRUG PRODUCT LIST

METHYLPREDNISOLONE

TABLET; ORAL

MEDROL

<u>AB</u>	+	PFIZER	<u>4MG</u>	<u>N011153</u>	<u>001</u>	
<u>AB</u>	+		<u>8MG</u>	<u>N011153</u>	<u>004</u>	
<u>AB</u>	+		<u>16MG</u>	<u>N011153</u>	<u>003</u>	
<u>AB</u>	+	!	<u>32MG</u>	<u>N011153</u>	<u>006</u>	

METHYLPREDNISOLONE

<u>AB</u>		JUBILANT CADISTA	<u>4MG</u>	<u>A040189</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>			<u>8MG</u>	<u>A040189</u>	<u>002</u>	Oct 31, 1997
<u>AB</u>			<u>16MG</u>	<u>A040189</u>	<u>003</u>	Jul 20, 2007
<u>AB</u>			<u>32MG</u>	<u>A040189</u>	<u>004</u>	Jul 20, 2007
<u>AB</u>		PRAXGEN	<u>4MG</u>	<u>A212262</u>	<u>001</u>	Jun 27, 2019
<u>AB</u>		SANDOZ	<u>4MG</u>	<u>A040194</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>		TIANJIN TIANYAO	<u>4MG</u>	<u>A204072</u>	<u>001</u>	May 14, 2018
<u>AB</u>		VINTAGE PHARMS	<u>4MG</u>	<u>A040183</u>	<u>001</u>	Dec 22, 1998
<u>AB</u>		WATSON LABS	<u>4MG</u>	<u>A040232</u>	<u>001</u>	Oct 16, 1997
<u>AB</u>		ZYDUS PHARMS	<u>4MG</u>	<u>A206751</u>	<u>001</u>	Apr 23, 2018
<u>AB</u>			<u>8MG</u>	<u>A206751</u>	<u>002</u>	Apr 23, 2018
<u>AB</u>			<u>16MG</u>	<u>A206751</u>	<u>003</u>	Apr 23, 2018
<u>AB</u>			<u>32MG</u>	<u>A206751</u>	<u>004</u>	Apr 23, 2018

MEDROL

+	PFIZER	2MG	N011153	002	
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METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

DEPO-MEDROL

<u>AB</u>	+	PFIZER	<u>40MG/ML</u>	<u>N011757</u>	<u>001</u>	
<u>AB</u>	+		<u>80MG/ML</u>	<u>N011757</u>	<u>004</u>	

METHYLPREDNISOLONE ACETATE

<u>AB</u>	!	AMNEAL	<u>40MG/ML</u>	<u>A210043</u>	<u>001</u>	May 20, 2019
<u>AB</u>			<u>40MG/ML</u>	<u>A216502</u>	<u>001</u>	Nov 03, 2023
<u>AB</u>	!		<u>80MG/ML</u>	<u>A210043</u>	<u>002</u>	May 20, 2019
<u>AB</u>			<u>80MG/ML</u>	<u>A216502</u>	<u>002</u>	Nov 03, 2023
<u>AB</u>		EUGIA PHARMA	<u>40MG/ML</u>	<u>A211930</u>	<u>001</u>	Apr 13, 2023
<u>AB</u>			<u>80MG/ML</u>	<u>A211930</u>	<u>002</u>	Apr 13, 2023
<u>AB</u>		MEITHEAL	<u>40MG/ML</u>	<u>A040557</u>	<u>001</u>	Feb 23, 2005
<u>AB</u>			<u>80MG/ML</u>	<u>A040557</u>	<u>002</u>	Feb 23, 2005
<u>AB</u>		PAR STERILE PRODUCTS	<u>40MG/ML</u>	<u>A214297</u>	<u>001</u>	Jan 21, 2022
<u>AB</u>			<u>80MG/ML</u>	<u>A214297</u>	<u>002</u>	Jan 21, 2022
<u>AB</u>	!	SAGENT PHARMS INC	<u>40MG/ML</u>	<u>A201835</u>	<u>002</u>	Jun 27, 2018
<u>AB</u>	!		<u>80MG/ML</u>	<u>A201835</u>	<u>003</u>	Jun 27, 2018
<u>AB</u>		SANDOZ	<u>40MG/ML</u>	<u>A040719</u>	<u>001</u>	Jan 29, 2009
<u>AB</u>			<u>40MG/ML</u>	<u>A040794</u>	<u>001</u>	Mar 05, 2009
<u>AB</u>			<u>80MG/ML</u>	<u>A040719</u>	<u>002</u>	Jan 29, 2009
<u>AB</u>			<u>80MG/ML</u>	<u>A040794</u>	<u>002</u>	Mar 05, 2009
<u>AB</u>		SLAYBACK PHARMA LLC	<u>40MG/ML</u>	<u>A214870</u>	<u>002</u>	May 10, 2023
<u>AB</u>			<u>80MG/ML</u>	<u>A214870</u>	<u>001</u>	Jan 03, 2023
<u>AB</u>		TEVA PHARMS USA	<u>40MG/ML</u>	<u>A040620</u>	<u>001</u>	Oct 27, 2006
<u>AB</u>			<u>80MG/ML</u>	<u>A040620</u>	<u>002</u>	Oct 27, 2006

DEPO-MEDROL

+	!	PFIZER	20MG/ML	N011757	002
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METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

<u>AP</u>		AMNEAL	<u>EQ 40MG BASE/VIAL</u>	<u>A207549</u>	<u>001</u>	Nov 09, 2016
<u>AP</u>			<u>EQ 125MG BASE/VIAL</u>	<u>A207549</u>	<u>002</u>	Nov 09, 2016
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 40MG BASE/VIAL</u>	<u>A040583</u>	<u>001</u>	Jul 30, 2004
<u>AP</u>			<u>EQ 125MG BASE/VIAL</u>	<u>A040583</u>	<u>002</u>	Jul 30, 2004
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A040612</u>	<u>001</u>	Aug 12, 2004
<u>AP</u>		HIKMA	<u>EQ 40MG BASE/VIAL</u>	<u>A203125</u>	<u>001</u>	Sep 26, 2022
<u>AP</u>			<u>EQ 125MG BASE/VIAL</u>	<u>A203125</u>	<u>002</u>	Sep 26, 2022
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A202691</u>	<u>001</u>	Feb 16, 2016
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A202691</u>	<u>002</u>	Feb 16, 2016
<u>AP</u>		SAGENT PHARMS INC	<u>EQ 40MG BASE/VIAL</u>	<u>A040888</u>	<u>001</u>	Jul 18, 2011
<u>AP</u>			<u>EQ 125MG BASE/VIAL</u>	<u>A040888</u>	<u>002</u>	Jul 18, 2011
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A040888</u>	<u>003</u>	Jul 18, 2011
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A040888</u>	<u>004</u>	Jul 18, 2011
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A040888</u>	<u>005</u>	Jul 18, 2011
<u>AP</u>		TIANJIN KINGYORK	<u>EQ 40MG BASE/VIAL</u>	<u>A212396</u>	<u>001</u>	Apr 20, 2021
<u>AP</u>			<u>EQ 125MG BASE/VIAL</u>	<u>A212396</u>	<u>002</u>	Apr 20, 2021
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>A212396</u>	<u>003</u>	Apr 20, 2021

PRESCRIPTION DRUG PRODUCT LIST

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A212396 004</u>	Apr 20, 2021
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A212396 005</u>	Apr 20, 2021
<u>SOLU-MEDROL</u>				
<u>AP</u>	+!	PHARMACIA AND UPJOHN	<u>EQ 40MG BASE/VIAL</u>	<u>N011856 003</u>
<u>AP</u>	+!		<u>EQ 125MG BASE/VIAL</u>	<u>N011856 004</u>
<u>AP</u>	+!		<u>EQ 500MG BASE/VIAL</u>	<u>N011856 005</u>
<u>AP</u>	+!		<u>EQ 1GM BASE/VIAL</u>	<u>N011856 006</u>
<u>AP</u>	+!		<u>EQ 2GM BASE/VIAL</u>	<u>N011856 007</u> Feb 27, 1985

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

<u>AB</u>	!	IMPAX LABS INC	<u>10MG</u>	<u>A204851 001</u>	Sep 21, 2015
<u>AB</u>		NOVITIUM PHARMA	<u>10MG</u>	<u>A215270 001</u>	Feb 18, 2022
TABLET; ORAL					
ANDROID 25					
BP		VALEANT PHARM INTL	25MG	A087147 001	
METHYLTESTOSTERONE					
BP		IMPAX LABS	10MG	A080767 002	

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE

<u>AP</u>		AVET LIFESCIENCES	<u>EQ 5MG BASE/ML</u>	<u>A204756 001</u>	Dec 20, 2013
<u>METOCLOPRAMIDE HYDROCHLORIDE</u>					
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A091392 001</u>	Apr 19, 2013
<u>AP</u>	!	HOSPIRA	<u>EQ 5MG BASE/ML</u>	<u>A073118 001</u>	Jan 17, 1991
<u>AP</u>		TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<u>A073135 001</u>	Nov 27, 1991

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

<u>AA</u>		ANI PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A071402 001</u>	Jun 25, 1993
<u>AA</u>	!	PHARM ASSOC	<u>EQ 5MG BASE/5ML</u>	<u>A072744 001</u>	May 28, 1991
SPRAY, METERED; NASAL					
GIMOTI					
	+!	EVOKE PHARMA INC	EQ 15MG BASE/SPRAY	N209388 001	Jun 19, 2020
TABLET; ORAL					

METOCLOPRAMIDE HYDROCHLORIDE

<u>AB</u>		IMPAX LABS INC	<u>EQ 5MG BASE</u>	<u>A071250 002</u>	Dec 28, 1995
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A071250 001</u>	Feb 03, 1988
<u>AB</u>		IPCA LABS LTD	<u>EQ 5MG BASE</u>	<u>A078807 001</u>	Jun 12, 2008
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A078807 002</u>	Jun 12, 2008
<u>AB</u>		STRIDES PHARMA	<u>EQ 5MG BASE</u>	<u>A077878 001</u>	Aug 28, 2006
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A070581 001</u>	Oct 17, 1985
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A077878 002</u>	Aug 28, 2006
<u>AB</u>		TEVA	<u>EQ 5MG BASE</u>	<u>A072801 001</u>	Jun 15, 1993
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A070184 001</u>	Jul 29, 1985

REGLAN

<u>AB</u>	+	ANI PHARMS	<u>EQ 5MG BASE</u>	<u>N017854 002</u>	May 05, 1987
<u>AB</u>	+!		<u>EQ 10MG BASE</u>	<u>N017854 001</u>	

TABLET, ORALLY DISINTEGRATING; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

		NOVEL LABS INC	EQ 5MG BASE	A202191 001	Aug 15, 2014
	!		EQ 10MG BASE	A202191 002	Aug 15, 2014

METOLAZONE

TABLET; ORAL

METOLAZONE

<u>AB</u>		ALEMBIC	<u>2.5MG</u>	<u>A213251 001</u>	Dec 02, 2020
<u>AB</u>			<u>5MG</u>	<u>A213251 002</u>	Dec 02, 2020
<u>AB</u>			<u>10MG</u>	<u>A213251 003</u>	Dec 02, 2020
<u>AB</u>		BAYSHORE PHARMS LLC	<u>2.5MG</u>	<u>A214799 001</u>	Mar 30, 2021
<u>AB</u>			<u>5MG</u>	<u>A214799 002</u>	Mar 30, 2021
<u>AB</u>			<u>10MG</u>	<u>A214799 003</u>	Mar 30, 2021
<u>AB</u>		INNOGENIX	<u>2.5MG</u>	<u>A213827 001</u>	Mar 30, 2021
<u>AB</u>			<u>5MG</u>	<u>A213827 002</u>	Mar 30, 2021
<u>AB</u>			<u>10MG</u>	<u>A213827 003</u>	Mar 30, 2021
<u>AB</u>		MYLAN	<u>2.5MG</u>	<u>A076698 001</u>	Dec 23, 2003
<u>AB</u>			<u>5MG</u>	<u>A076698 002</u>	Oct 19, 2004
<u>AB</u>			<u>10MG</u>	<u>A076698 003</u>	Oct 19, 2004
<u>AB</u>		NE RX PHARMA	<u>2.5MG</u>	<u>A216216 001</u>	Oct 25, 2022

PRESCRIPTION DRUG PRODUCT LIST

METOLAZONE

TABLET; ORAL

METOLAZONE

AB		5MG	<u>A216216 002</u>	Oct 25, 2022
AB		10MG	<u>A216216 003</u>	Oct 25, 2022
AB	RENATA	2.5MG	<u>A215616 001</u>	Oct 24, 2022
AB		5MG	<u>A215616 002</u>	Oct 24, 2022
AB		10MG	<u>A215616 003</u>	Oct 24, 2022
AB	RUBICON	2.5MG	<u>A215184 001</u>	Aug 20, 2021
AB		5MG	<u>A215184 002</u>	Aug 20, 2021
AB		10MG	<u>A215184 003</u>	Aug 20, 2021
AB	SANDOZ	2.5MG	<u>A076732 001</u>	Dec 19, 2003
AB	!	5MG	<u>A076466 001</u>	Dec 19, 2003
AB	!	10MG	<u>A076466 002</u>	Dec 19, 2003

METOPROLOL SUCCINATE

CAPSULE, EXTENDED RELEASE; ORAL

KAPSPARGO SPRINKLE

+	SPIL	EQ 25MG TARTRATE	N210428 001	Jan 26, 2018
+		EQ 50MG TARTRATE	N210428 002	Jan 26, 2018
+		EQ 100MG TARTRATE	N210428 003	Jan 26, 2018
+	!	EQ 200MG TARTRATE	N210428 004	Jan 26, 2018

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

AB	ACTAVIS ELIZABETH	<u>EQ 25MG TARTRATE</u>	<u>A204161 001</u>	Nov 25, 2016
AB		<u>EQ 50MG TARTRATE</u>	<u>A204161 002</u>	Nov 25, 2016
AB		<u>EQ 100MG TARTRATE</u>	<u>A204161 003</u>	Nov 25, 2016
AB		<u>EQ 200MG TARTRATE</u>	<u>A204161 004</u>	Nov 25, 2016
AB	ACTAVIS LABS FL INC	<u>EQ 50MG TARTRATE</u>	<u>A076862 001</u>	Aug 03, 2009
AB	ALKEM LABS LTD	<u>EQ 25MG TARTRATE</u>	<u>A211143 001</u>	Nov 25, 2020
AB		<u>EQ 50MG TARTRATE</u>	<u>A211143 002</u>	Nov 25, 2020
AB		<u>EQ 100MG TARTRATE</u>	<u>A211143 003</u>	Nov 25, 2020
AB		<u>EQ 200MG TARTRATE</u>	<u>A211143 004</u>	Nov 25, 2020
AB	CIPLA	<u>EQ 50MG TARTRATE</u>	<u>A207465 001</u>	Oct 26, 2018
AB		<u>EQ 100MG TARTRATE</u>	<u>A207465 002</u>	Oct 26, 2018
AB		<u>EQ 200MG TARTRATE</u>	<u>A207465 003</u>	Oct 26, 2018
AB	DR REDDYS LABS LTD	<u>EQ 25MG TARTRATE</u>	<u>A090617 001</u>	Aug 01, 2012
AB		<u>EQ 50MG TARTRATE</u>	<u>A090617 002</u>	Aug 01, 2012
AB	GRANULES	<u>EQ 25MG TARTRATE</u>	<u>A216509 001</u>	Aug 07, 2023
AB		<u>EQ 25MG TARTRATE</u>	<u>A216916 001</u>	Jun 12, 2023
AB		<u>EQ 50MG TARTRATE</u>	<u>A216509 002</u>	Aug 07, 2023
AB		<u>EQ 50MG TARTRATE</u>	<u>A216916 002</u>	Jun 12, 2023
AB		<u>EQ 100MG TARTRATE</u>	<u>A216509 003</u>	Aug 07, 2023
AB		<u>EQ 100MG TARTRATE</u>	<u>A216916 003</u>	Jun 12, 2023
AB		<u>EQ 200MG TARTRATE</u>	<u>A216509 004</u>	Aug 07, 2023
AB		<u>EQ 200MG TARTRATE</u>	<u>A216916 004</u>	Jun 12, 2023
AB	HETERO LABS LTD III	<u>EQ 25MG TARTRATE</u>	<u>A205541 001</u>	Nov 06, 2020
AB		<u>EQ 50MG TARTRATE</u>	<u>A205541 002</u>	Nov 06, 2020
AB		<u>EQ 100MG TARTRATE</u>	<u>A205541 003</u>	Nov 06, 2020
AB		<u>EQ 200MG TARTRATE</u>	<u>A205541 004</u>	Nov 06, 2020
AB	MYLAN PHARMS INC	<u>EQ 25MG TARTRATE</u>	<u>A202033 001</u>	Dec 15, 2011
AB		<u>EQ 50MG TARTRATE</u>	<u>A202033 002</u>	Dec 15, 2011
AB		<u>EQ 100MG TARTRATE</u>	<u>A202033 003</u>	Dec 15, 2011
AB		<u>EQ 200MG TARTRATE</u>	<u>A202033 004</u>	Dec 15, 2011
AB	NOVAST LABS	<u>EQ 25MG TARTRATE</u>	<u>A204106 001</u>	Feb 06, 2018
AB		<u>EQ 50MG TARTRATE</u>	<u>A204106 002</u>	Feb 06, 2018
AB		<u>EQ 100MG TARTRATE</u>	<u>A204106 003</u>	Feb 06, 2018
AB		<u>EQ 200MG TARTRATE</u>	<u>A204106 004</u>	Feb 06, 2018
AB	PHARMADAX INC	<u>EQ 25MG TARTRATE</u>	<u>A203028 001</u>	Mar 31, 2020
AB		<u>EQ 50MG TARTRATE</u>	<u>A203028 002</u>	Mar 31, 2020
AB		<u>EQ 100MG TARTRATE</u>	<u>A203699 001</u>	Mar 30, 2020
AB		<u>EQ 200MG TARTRATE</u>	<u>A203699 002</u>	Mar 30, 2020
AB	REDDYS	<u>EQ 100MG TARTRATE</u>	<u>A078889 001</u>	Aug 15, 2012
AB		<u>EQ 200MG TARTRATE</u>	<u>A078889 002</u>	Aug 15, 2012
AB	SUNSHINE	<u>EQ 50MG TARTRATE</u>	<u>A214004 001</u>	Sep 26, 2022
AB		<u>EQ 100MG TARTRATE</u>	<u>A214004 002</u>	Sep 26, 2022
AB		<u>EQ 200MG TARTRATE</u>	<u>A214004 003</u>	Sep 26, 2022
AB	VISUM PHARM	<u>EQ 25MG TARTRATE</u>	<u>A207206 001</u>	Dec 19, 2018
AB		<u>EQ 50MG TARTRATE</u>	<u>A207206 002</u>	Dec 19, 2018
AB		<u>EQ 100MG TARTRATE</u>	<u>A207206 003</u>	Dec 19, 2018
AB		<u>EQ 200MG TARTRATE</u>	<u>A207206 004</u>	Dec 19, 2018
AB	WOCKHARDT	<u>EQ 25MG TARTRATE</u>	<u>A090615 001</u>	Jul 22, 2010
AB		<u>EQ 50MG TARTRATE</u>	<u>A090615 002</u>	Jul 22, 2010

PRESCRIPTION DRUG PRODUCT LIST

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

METOPROLOL SUCCINATE

<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A090615 003</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A090615 004</u>	Jul 22, 2010
<u>AB</u>	YICHANG HUMANWELL	<u>EQ 25MG TARTRATE</u>	<u>A213854 004</u>	Aug 01, 2022
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A213854 003</u>	Nov 08, 2021
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A213854 001</u>	Feb 12, 2021
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A213854 002</u>	Feb 12, 2021
<u>AB</u>	ZYDUS PHARMS	<u>EQ 25MG TARTRATE</u>	<u>A203894 001</u>	Mar 23, 2018
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A203894 002</u>	Mar 23, 2018
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A203894 003</u>	Mar 23, 2018
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A203894 004</u>	Mar 23, 2018

TOPROL-XL

<u>AB</u>	+	TOPROL	<u>EQ 25MG TARTRATE</u>	<u>N019962 004</u>	Feb 05, 2001
<u>AB</u>	+	!	<u>EQ 50MG TARTRATE</u>	<u>N019962 001</u>	Jan 10, 1992
<u>AB</u>	+		<u>EQ 100MG TARTRATE</u>	<u>N019962 002</u>	Jan 10, 1992
<u>AB</u>	+	!	<u>EQ 200MG TARTRATE</u>	<u>N019962 003</u>	Jan 10, 1992

METOPROLOL TARTRATE

INJECTABLE; INJECTION

METOPROLOL TARTRATE

<u>AP</u>	BAXTER HLTHCARE CORP	<u>1MG/ML</u>	<u>A078950 001</u>	Apr 29, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A091045 001</u>	Oct 25, 2010
<u>AP</u>	GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A204205 001</u>	Aug 27, 2014
<u>AP</u>	HIKMA	<u>1MG/ML</u>	<u>A076495 001</u>	Jul 07, 2003
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A077761 001</u>	May 30, 2007
<u>AP</u>	HOSPIRA	<u>1MG/ML</u>	<u>A074133 001</u>	Dec 21, 1993
<u>AP</u>	!	<u>1MG/ML</u>	<u>A078085 001</u>	Apr 29, 2008
<u>AP</u>	SANDOZ	<u>1MG/ML</u>	<u>A077360 001</u>	Oct 02, 2007

TABLET; ORAL

LOPRESSOR

<u>AB</u>	+	VALIDUS PHARMS	<u>50MG</u>	<u>N017963 001</u>
<u>AB</u>	+		<u>100MG</u>	<u>N017963 002</u>

METOPROLOL TARTRATE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A202871 001</u>	May 28, 2013
<u>AB</u>		<u>50MG</u>	<u>A202871 002</u>	May 28, 2013
<u>AB</u>		<u>100MG</u>	<u>A202871 003</u>	May 28, 2013
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A077739 001</u>	Sep 11, 2007
<u>AB</u>		<u>37.5MG</u>	<u>A077739 004</u>	Dec 07, 2023
<u>AB</u>		<u>50MG</u>	<u>A077739 002</u>	Sep 11, 2007
<u>AB</u>		<u>75MG</u>	<u>A077739 005</u>	Dec 07, 2023
<u>AB</u>		<u>100MG</u>	<u>A077739 003</u>	Sep 11, 2007
<u>AB</u>	IPCA LABS LTD	<u>25MG</u>	<u>A078459 001</u>	Jun 17, 2008
<u>AB</u>		<u>50MG</u>	<u>A078459 002</u>	Jun 17, 2008
<u>AB</u>		<u>100MG</u>	<u>A078459 003</u>	Jun 17, 2008
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A076704 001</u>	Jan 16, 2004
<u>AB</u>		<u>37.5MG</u>	<u>A076704 004</u>	Mar 18, 2015
<u>AB</u>		<u>50MG</u>	<u>A076704 002</u>	Jan 16, 2004
<u>AB</u>		<u>75MG</u>	<u>A076704 005</u>	Mar 18, 2015
<u>AB</u>	!	<u>100MG</u>	<u>A076704 003</u>	Jan 16, 2004
<u>AB</u>	RENATA	<u>50MG</u>	<u>A074453 001</u>	Apr 27, 1995
<u>AB</u>		<u>100MG</u>	<u>A074453 002</u>	Apr 27, 1995
<u>AB</u>	RUBICON	<u>25MG</u>	<u>A200981 001</u>	Oct 28, 2014
<u>AB</u>		<u>37.5MG</u>	<u>A200981 004</u>	Aug 21, 2019
<u>AB</u>		<u>50MG</u>	<u>A200981 002</u>	Oct 28, 2014
<u>AB</u>		<u>75MG</u>	<u>A200981 005</u>	Aug 21, 2019
<u>AB</u>		<u>100MG</u>	<u>A200981 003</u>	Oct 28, 2014
<u>AB</u>	SUN PHARM INDS INC	<u>25MG</u>	<u>A076670 001</u>	Jan 15, 2004
<u>AB</u>		<u>50MG</u>	<u>A074644 001</u>	Dec 10, 1996
<u>AB</u>		<u>100MG</u>	<u>A074644 002</u>	Dec 10, 1996
<u>AB</u>	YOUNGTECH PHARMS INC	<u>25MG</u>	<u>A208955 001</u>	Feb 05, 2020
<u>AB</u>		<u>50MG</u>	<u>A208955 002</u>	Feb 05, 2020
<u>AB</u>		<u>100MG</u>	<u>A208955 003</u>	Feb 05, 2020

METRONIDAZOLE

CAPSULE; ORAL

FLAGYL

<u>AB</u>	+	PFIZER	<u>375MG</u>	<u>N020334 001</u>	May 03, 1995
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METRONIDAZOLE

<u>AB</u>	ALEMBIC	<u>375MG</u>	<u>A079065 001</u>	Jun 23, 2009
<u>AB</u>	CHARTWELL RX	<u>375MG</u>	<u>A076522 001</u>	Jan 29, 2004

PRESCRIPTION DRUG PRODUCT LIST

METRONIDAZOLE

CREAM; TOPICAL

METROCREAM

AB	+ !	GALDERMA LABS LP	0.75%	N020531	001	Sep 20, 1995
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METRONIDAZOLE

AB		COSETTE	0.75%	A077549	001	Dec 19, 2007
AB		FOUGERA PHARMS	0.75%	A076408	001	May 28, 2004
AB		ZYDUS LIFESCIENCES	0.75%	A217128	001	Apr 21, 2023

NORITATE

+ !	BAUSCH	1%	N020743	001	Sep 26, 1997
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GEL; TOPICAL

METROGEL

AB	+ !	GALDERMA LABS LP	0.75%	N019737	001	Nov 22, 1988
AB	+ !		1%	N021789	001	Jun 30, 2005

METRONIDAZOLE

AB		ALEMBIC	1%	A212646	001	Sep 03, 2021
AB		COSETTE	0.75%	A078178	001	Jan 19, 2011
AB			1%	A216692	001	Jan 23, 2023
AB		ENCUBE	0.75%	A077547	001	Jul 13, 2006
AB		FOUGERA PHARMS	0.75%	A077018	001	Jun 06, 2006
AB		TARO	0.75%	A077819	001	Jul 18, 2006
AB			1%	A204651	001	Mar 14, 2017

GEL; VAGINAL

METROGEL-VAGINAL

AB	+ !	BAUSCH	0.75%	N020208	001	Aug 17, 1992
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METRONIDAZOLE

AB		COSETTE	0.75%	A216323	001	Jul 12, 2023
AB		ENCUBE	0.75%	A215610	001	Jun 16, 2023
AB		GLENMARK PHARMS LTD	0.75%	A215794	001	Jan 27, 2022
AB		PADAGIS ISRAEL	0.75%	A211786	001	Jul 02, 2019
AB		SOLARIS PHARMA CORP	0.75%	A213648	001	Oct 14, 2021

VANDAZOLE

BX	!	TEVA PHARMS	0.75%	N021806	001	May 20, 2005
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NUVESSA

+ !	CHEMO RESEARCH SL	1.3%	N205223	001	Mar 24, 2014
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INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

AP	+ !	BAXTER HLTHCARE	500MG/100ML	N018657	001	
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METRO I.V. IN PLASTIC CONTAINER

AP	+ !	B BRAUN	500MG/100ML	N018900	001	Sep 29, 1983
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METRONIDAZOLE IN PLASTIC CONTAINER

AP		AMNEAL	500MG/100ML	A217665	001	May 24, 2023
AP		BAXTER HLTHCARE	500MG/100ML	A078084	001	Mar 31, 2008
		CORP				
AP		GLAND PHARMA LTD	500MG/100ML	A212435	001	Aug 03, 2020
AP	+ !	HOSPIRA	500MG/100ML	N018890	002	Nov 18, 1983
AP		INFORLIFE	500MG/100ML	A206191	001	Feb 25, 2019

LOTION; TOPICAL

METROLOTION

AB	+ !	GALDERMA LABS LP	0.75%	N020901	001	Nov 24, 1998
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METRONIDAZOLE

AB		FOUGERA PHARMS	0.75%	A077197	001	May 24, 2006
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SUSPENSION; ORAL

LIKMEZ

+ !	SAPTALIS PHARMS	500MG/5ML	N216755	001	Sep 22, 2023
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TABLET; ORAL

METRONIDAZOLE

AB		ALEMBIC	250MG	A208681	001	Jun 20, 2017
AB			500MG	A208681	002	Jun 20, 2017
AB		ALEMBIC PHARMS LTD	250MG	A079067	001	Mar 13, 2009
AB			500MG	A079067	002	Mar 13, 2009
AB		AUROBINDO PHARMA	250MG	A203974	001	May 29, 2015
		LTD				
AB	!		500MG	A203974	002	May 29, 2015
AB		CADILA	250MG	A206560	001	Nov 16, 2016
AB			500MG	A206560	002	Nov 16, 2016
AB		CADILA PHARMS LTD	250MG	A209794	001	Dec 12, 2017
AB			500MG	A209794	002	Dec 12, 2017
AB		FLAMINGO PHARMS	250MG	A207309	001	May 16, 2016
AB			500MG	A207309	002	May 16, 2016
AB		HERITAGE PHARMS INC	250MG	A205245	001	Sep 23, 2015
AB			500MG	A205245	002	Sep 23, 2015
AB		INNOGENIX	250MG	A070772	001	Jul 16, 1986

PRESCRIPTION DRUG PRODUCT LIST

METRONIDAZOLE

TABLET; ORAL

METRONIDAZOLE

<u>AB</u>		<u>500MG</u>	<u>A070772</u>	<u>002</u>	Jul 16, 1986
<u>AB</u>	LUPIN LTD	<u>250MG</u>	<u>A209096</u>	<u>001</u>	Sep 12, 2017
<u>AB</u>		<u>500MG</u>	<u>A209096</u>	<u>002</u>	Sep 12, 2017
<u>AB</u>	PLIVA	<u>500MG</u>	<u>A070033</u>	<u>001</u>	Dec 06, 1984
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A070040</u>	<u>001</u>	Jan 29, 1985
<u>AB</u>		<u>250MG</u>	<u>A208162</u>	<u>001</u>	May 25, 2016
<u>AB</u>		<u>500MG</u>	<u>A070039</u>	<u>001</u>	Jan 29, 1985
<u>AB</u>		<u>500MG</u>	<u>A208162</u>	<u>002</u>	May 25, 2016
<u>AB</u>	TEVA PHARMS USA	<u>250MG</u>	<u>A070027</u>	<u>001</u>	Nov 06, 1984
<u>AB</u>	UNICHEM	<u>250MG</u>	<u>A203458</u>	<u>001</u>	Jan 22, 2014
<u>AB</u>		<u>500MG</u>	<u>A203458</u>	<u>002</u>	Jan 22, 2014
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A070035</u>	<u>001</u>	Dec 20, 1984
<u>AB</u>	WATSON LABS INC	<u>500MG</u>	<u>A070044</u>	<u>001</u>	Feb 08, 1985

METYRAPONE

CAPSULE; ORAL

METOPIRONE

+! HRA PHARMA 250MG N012911 002 Aug 09, 1996

METYROSINE

CAPSULE; ORAL

DEMSEERAB +! BAUSCH 250MG N017871 001METYROSINEAB AMNEAL 250MG A213734 001 Jul 24, 2020MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>150MG</u>	<u>A074450</u>	<u>001</u>	May 16, 1996
<u>AB</u>		<u>200MG</u>	<u>A074450</u>	<u>002</u>	May 16, 1996
<u>AB</u>		<u>250MG</u>	<u>A074450</u>	<u>003</u>	May 16, 1996
<u>AB</u>	ANNORA PHARMA	<u>150MG</u>	<u>A216463</u>	<u>001</u>	Nov 09, 2022
<u>AB</u>		<u>200MG</u>	<u>A216463</u>	<u>002</u>	Nov 09, 2022
<u>AB</u>		<u>250MG</u>	<u>A216463</u>	<u>003</u>	Nov 09, 2022
<u>AB</u>	CARNEGIE	<u>150MG</u>	<u>A215315</u>	<u>001</u>	Aug 26, 2022
<u>AB</u>		<u>200MG</u>	<u>A215315</u>	<u>002</u>	Aug 26, 2022
<u>AB</u>		<u>250MG</u>	<u>A215315</u>	<u>003</u>	Aug 26, 2022
<u>AB</u>	CHEMISTRY HLTH	<u>150MG</u>	<u>A215876</u>	<u>001</u>	Feb 27, 2023
<u>AB</u>		<u>200MG</u>	<u>A215876</u>	<u>002</u>	Feb 27, 2023
<u>AB</u>		<u>250MG</u>	<u>A215876</u>	<u>003</u>	Feb 27, 2023
<u>AB</u>	CROSSMEDIKA SA	<u>150MG</u>	<u>A213500</u>	<u>001</u>	Jul 22, 2020
<u>AB</u>		<u>200MG</u>	<u>A213500</u>	<u>002</u>	Jul 22, 2020
<u>AB</u>		<u>250MG</u>	<u>A213500</u>	<u>003</u>	Jul 22, 2020
<u>AB</u>	NOVAST LABS	<u>150MG</u>	<u>A214352</u>	<u>001</u>	Jan 25, 2021
<u>AB</u>		<u>200MG</u>	<u>A214352</u>	<u>002</u>	Jan 25, 2021
<u>AB</u>		<u>250MG</u>	<u>A214352</u>	<u>003</u>	Jan 25, 2021
<u>AB</u>	SENORES PHARMS	<u>150MG</u>	<u>A214089</u>	<u>001</u>	Oct 01, 2021
<u>AB</u>		<u>200MG</u>	<u>A214089</u>	<u>002</u>	Oct 01, 2021
<u>AB</u>		<u>250MG</u>	<u>A214089</u>	<u>003</u>	Oct 01, 2021
<u>AB</u>	TEVA	<u>150MG</u>	<u>A074377</u>	<u>001</u>	May 16, 1995
<u>AB</u>		<u>200MG</u>	<u>A074377</u>	<u>002</u>	May 16, 1995
<u>AB</u>	!	<u>250MG</u>	<u>A074377</u>	<u>003</u>	May 16, 1995

MICAFUNGIN SODIUM

INJECTABLE; INTRAVENOUS

MICAFUNGIN SODIUM

<u>AP</u>	APOTEX	<u>EQ 50MG BASE/VIAL</u>	<u>A208366</u>	<u>001</u>	Nov 05, 2020
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A208366</u>	<u>002</u>	Nov 05, 2020
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 50MG BASE/VIAL</u>	<u>A207344</u>	<u>001</u>	May 17, 2019
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A207344</u>	<u>002</u>	May 17, 2019
<u>AP</u>	HIKMA	<u>EQ 50MG BASE/VIAL</u>	<u>A213261</u>	<u>001</u>	Jul 09, 2021
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A213261</u>	<u>002</u>	Jul 09, 2021
<u>AP</u>	JIANGSU HANSOH PHARM	<u>EQ 50MG BASE/VIAL</u>	<u>A213363</u>	<u>001</u>	Jul 09, 2021
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A213363</u>	<u>002</u>	Jul 09, 2021
<u>AP</u>	MEITHEAL	<u>EQ 50MG BASE/VIAL</u>	<u>A215381</u>	<u>001</u>	Sep 28, 2022
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A215381</u>	<u>002</u>	Sep 28, 2022
<u>AP</u>	XELLIA PHARMS APS	<u>EQ 50MG BASE/VIAL</u>	<u>A211713</u>	<u>001</u>	Jun 02, 2021
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A211713</u>	<u>002</u>	Jun 02, 2021
<u>AP</u>	ZYDUS PHARMS	<u>EQ 50MG BASE/VIAL</u>	<u>A215241</u>	<u>001</u>	Oct 24, 2022
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A215241</u>	<u>002</u>	Oct 24, 2022

PRESCRIPTION DRUG PRODUCT LIST

MICAFUNGIN SODIUM

INJECTABLE; INTRAVENOUS

MYCAMINE

<u>AP</u>	+	ASTELLAS	<u>EQ 50MG BASE/VIAL</u>	<u>N021506 002</u>	Mar 16, 2005
<u>AP</u>	+		<u>EQ 100MG BASE/VIAL</u>	<u>N021506 003</u>	Jun 27, 2006

POWDER; INTRAVENOUS

MICAFUNGIN

+	!	PAR STERILE PRODUCTS	EQ 50MG BASE/VIAL	N212156 001	Jun 16, 2021
+	!		EQ 100MG BASE/VIAL	N212156 002	Jun 16, 2021

SOLUTION; INTRAVENOUS

MICAFUNGIN IN SODIUM CHLORIDE 0.9%

+	!	BAXTER HLTHCARE CORP	EQ 50MG BASE/50ML (EQ 1MG BASE/ML)	N216142 001	Sep 29, 2023
+	!		EQ 100MG BASE/100ML (EQ 1MG BASE/ML)	N216142 002	Sep 29, 2023
+	!		EQ 150MG BASE/150ML (EQ 1MG BASE/ML)	N216142 003	Sep 29, 2023

MICONAZOLE

TABLET; BUCCAL

ORAVIG

+	!	GALT PHARMS	50MG	N022404 001	Apr 16, 2010
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MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

<u>AB</u>		ACTAVIS PHARMA	<u>200MG</u>	<u>A073508 001</u>	Nov 19, 1993
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MONISTAT 3

<u>AB</u>	+	MEDTECH PRODUCTS	<u>200MG</u>	<u>N018888 001</u>	Aug 15, 1984
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MICONAZOLE NITRATE; WHITE PETROLATUM; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+	!	MYLAN	0.25%; 81.35%; 15%	N021026 001	Feb 16, 2006
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MIDAZOLAM

SOLUTION; INTRAVENOUS

MIDAZOLAM IN 0.9% SODIUM CHLORIDE

<u>AP</u>		HIKMA	<u>50MG/50ML (1MG/ML)</u>	<u>A216159 001</u>	Apr 17, 2023
<u>AP</u>			<u>100MG/100ML (1MG/ML)</u>	<u>A216159 002</u>	Apr 17, 2023
<u>AP</u>	+	INFORLIFE	<u>50MG/50ML (1MG/ML)</u>	<u>N211844 001</u>	Mar 22, 2021
<u>AP</u>	+		<u>100MG/100ML (1MG/ML)</u>	<u>N211844 002</u>	Mar 22, 2021

MIDAZOLAM IN 0.8% SODIUM CHLORIDE

+	!	EXELA PHARMA	50MG/50ML (1MG/ML)	N215868 001	Jul 20, 2022
+	!		100MG/100ML (1MG/ML)	N215868 002	Jul 20, 2022

SPRAY; NASAL

NAYZILAM

+	!	UCB INC	5MG/SPRAY	N211321 001	May 17, 2019
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MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

<u>AP</u>		AKORN	<u>EQ 1MG BASE/ML</u>	<u>A075494 001</u>	Jun 30, 2000
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A075494 002</u>	Jun 30, 2000
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A075154 002</u>	Jun 20, 2000
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A075154 001</u>	Jun 20, 2000
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 1MG BASE/ML</u>	<u>A090696 001</u>	Feb 29, 2012
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A090850 001</u>	Jan 25, 2012
<u>AP</u>		HIKMA	<u>EQ 1MG BASE/ML</u>	<u>A075243 001</u>	Jun 20, 2000
<u>AP</u>			<u>EQ 1MG BASE/ML</u>	<u>A075247 002</u>	Jun 23, 2000
<u>AP</u>			<u>EQ 1MG BASE/ML</u>	<u>A075324 001</u>	Jun 20, 2000
<u>AP</u>			<u>EQ 1MG BASE/ML</u>	<u>A075421 002</u>	Jun 20, 2000
<u>AP</u>			<u>EQ 1MG BASE/ML</u>	<u>A212847 001</u>	Dec 11, 2020
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A075243 002</u>	Jun 20, 2000
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A075247 001</u>	Jun 23, 2000
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A075324 002</u>	Jun 20, 2000
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A075421 001</u>	Jun 20, 2000
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A212847 002</u>	Dec 11, 2020
<u>AP</u>	!	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075293 001</u>	Jun 20, 2000
<u>AP</u>	!		<u>EQ 5MG BASE/ML</u>	<u>A075293 002</u>	Jun 20, 2000
<u>AP</u>		MICRO LABS	<u>EQ 1MG BASE/ML</u>	<u>A217504 001</u>	Aug 21, 2023
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A217504 002</u>	Aug 21, 2023

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>		FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A203460 001</u>	Aug 22, 2014
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A203460 002</u>	Aug 22, 2014
<u>AP</u>	!	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075857 001</u>	Jul 22, 2002

PRESCRIPTION DRUG PRODUCT LIST

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	!		<u>EQ 5MG BASE/ML</u>	<u>A075857 002</u>	Jul 22, 2002
<u>AP</u>		STERISCIENCE	<u>EQ 1MG BASE/ML</u>	<u>A090315 001</u>	Nov 29, 2010
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A090315 002</u>	Nov 29, 2010

MIDOZALAM HYDROCHLORIDE

<u>AP</u>		STERISCIENCE	<u>EQ 1MG BASE/ML</u>	<u>A090316 001</u>	May 04, 2011
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A090316 002</u>	May 04, 2011

MIDAZOLAM HYDROCHLORIDE

		FRESENIUS KABI USA	EQ 5MG BASE/ML	A208878 001	Mar 28, 2017
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SOLUTION; INTRAMUSCULAR

SEIZALAM

+! MMT

			EQ 50MG BASE/10ML (EQ 5MG BASE/ML)	N209566 001	Sep 14, 2018
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SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

<u>AA</u>	!	HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A075873 001</u>	Apr 30, 2002
<u>AA</u>		PADAGIS US	<u>EQ 2MG BASE/ML</u>	<u>A076379 001</u>	May 02, 2005

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

<u>AB</u>		ALEM BIC	<u>2.5MG</u>	<u>A214734 001</u>	Jan 21, 2021
<u>AB</u>			<u>5MG</u>	<u>A214734 002</u>	Jan 21, 2021
<u>AB</u>			<u>10MG</u>	<u>A214734 003</u>	Jan 21, 2021
<u>AB</u>		APOTEX	<u>2.5MG</u>	<u>A077746 001</u>	Sep 12, 2006
<u>AB</u>			<u>5MG</u>	<u>A077746 002</u>	Sep 12, 2006
<u>AB</u>			<u>10MG</u>	<u>A077746 003</u>	Sep 12, 2006
<u>AB</u>		AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A212774 001</u>	Aug 10, 2020
<u>AB</u>			<u>5MG</u>	<u>A212774 002</u>	Aug 10, 2020
<u>AB</u>			<u>10MG</u>	<u>A212774 003</u>	Aug 10, 2020
<u>AB</u>		IMPAX PHARMS	<u>2.5MG</u>	<u>A076449 001</u>	May 27, 2004
<u>AB</u>			<u>5MG</u>	<u>A076449 002</u>	May 27, 2004
<u>AB</u>			<u>10MG</u>	<u>A076449 003</u>	Dec 16, 2005
<u>AB</u>		MANKIND PHARMA	<u>2.5MG</u>	<u>A217271 001</u>	Nov 01, 2023
<u>AB</u>			<u>5MG</u>	<u>A217271 002</u>	Nov 01, 2023
<u>AB</u>			<u>10MG</u>	<u>A217271 003</u>	Nov 01, 2023
<u>AB</u>		MYLAN PHARMS INC	<u>2.5MG</u>	<u>A076577 001</u>	Sep 10, 2003
<u>AB</u>			<u>5MG</u>	<u>A076577 002</u>	Sep 10, 2003
<u>AB</u>			<u>10MG</u>	<u>A076577 003</u>	Sep 10, 2003
<u>AB</u>		NOVUGEN	<u>2.5MG</u>	<u>A211973 001</u>	Oct 17, 2023
<u>AB</u>			<u>5MG</u>	<u>A211973 002</u>	Oct 17, 2023
<u>AB</u>			<u>10MG</u>	<u>A211973 003</u>	Oct 17, 2023
<u>AB</u>		PAR PHARM INC	<u>2.5MG</u>	<u>A207169 001</u>	Oct 29, 2018
<u>AB</u>			<u>5MG</u>	<u>A207169 002</u>	Oct 29, 2018
<u>AB</u>			<u>10MG</u>	<u>A207169 003</u>	Oct 29, 2018
<u>AB</u>		RUBICON	<u>2.5MG</u>	<u>A212543 001</u>	Aug 19, 2019
<u>AB</u>			<u>5MG</u>	<u>A212543 002</u>	Aug 19, 2019
<u>AB</u>			<u>10MG</u>	<u>A212543 003</u>	Aug 19, 2019
<u>AB</u>		THINQ PHARM-CRO PVT	<u>2.5MG</u>	<u>A207613 001</u>	Nov 02, 2018
<u>AB</u>			<u>5MG</u>	<u>A207613 002</u>	Nov 02, 2018
<u>AB</u>			<u>10MG</u>	<u>A207613 003</u>	Nov 02, 2018
<u>AB</u>		XIROMED	<u>2.5MG</u>	<u>A207849 001</u>	Oct 01, 2020
<u>AB</u>			<u>5MG</u>	<u>A207849 002</u>	Oct 01, 2020
<u>AB</u>			<u>10MG</u>	<u>A207849 003</u>	Oct 01, 2020
<u>AB</u>		ZYDUS	<u>2.5MG</u>	<u>A213055 001</u>	Sep 01, 2020
<u>AB</u>			<u>5MG</u>	<u>A213055 002</u>	Sep 01, 2020
<u>AB</u>			<u>10MG</u>	<u>A213055 003</u>	Sep 01, 2020

ORVATEN

<u>AB</u>		UPSHER SMITH LABS	<u>2.5MG</u>	<u>A076725 001</u>	Nov 03, 2004
<u>AB</u>	!		<u>5MG</u>	<u>A076725 002</u>	Nov 03, 2004
<u>AB</u>			<u>10MG</u>	<u>A076725 003</u>	Nov 03, 2004

MIDOSTAURIN

CAPSULE; ORAL

RYDAPT

	+	NOVARTIS	25MG	N207997 001	Apr 28, 2017
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PRESCRIPTION DRUG PRODUCT LIST

MIFEPRISTONE

TABLET; ORAL

KORLYM

AB	+ !	CORCEPT THERAP	300MG	N202107	001	Feb 17, 2012
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MIFEPREX

AB	+ !	DANCO LABS LLC	200MG	N020687	001	Sep 28, 2000
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MIFEPRISTONE

AB		GENBIOPRO	200MG	A091178	001	Apr 11, 2019
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AB		TEVA PHARMS USA INC	300MG	A211436	001	Aug 03, 2020
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MIGALASTAT HYDROCHLORIDE

CAPSULE; ORAL

GALAFOLD

	+ !	AMICUS THERAP US	EQ 123MG BASE	N208623	001	Aug 10, 2018
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MIGLITOL

TABLET; ORAL

GLYSET

AB	+ !	PFIZER	25MG	N020682	001	Dec 18, 1996
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AB	+		50MG	N020682	002	Dec 18, 1996
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AB	+		100MG	N020682	003	Dec 18, 1996
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MIGLITOL

AB		WESTMINSTER PHARMS	25MG	A203965	001	Feb 24, 2015
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AB			50MG	A203965	002	Feb 24, 2015
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AB			100MG	A203965	003	Feb 24, 2015
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MIGLUSTAT

CAPSULE; ORAL

MIGLUSTAT

AB		ANI PHARMS	100MG	A208342	001	Apr 17, 2018
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AB		BRECKENRIDGE	100MG	A209325	001	Feb 03, 2022
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YARGESA

AB		EDENBRIDGE PHARMS	100MG	A209821	001	Aug 06, 2020
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ZAVESCA

AB	+ !	ACTELION	100MG	N021348	001	Jul 31, 2003
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OPFOLDA

	+ !	AMICUS THERAP US	65MG	N215211	001	Sep 28, 2023
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MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

SAVELLA

	+	ABBVIE	12.5MG	N022256	001	Jan 14, 2009
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	+		25MG	N022256	002	Jan 14, 2009
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	+ !		50MG	N022256	003	Jan 14, 2009
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	+		100MG	N022256	004	Jan 14, 2009
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MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

AP		CAPLIN	EQ 1MG BASE/ML	A214380	001	Apr 16, 2021
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AP		FRESENIUS KABI USA	EQ 1MG BASE/ML	A075936	001	May 28, 2002
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AP		HIKMA	EQ 1MG BASE/ML	A075530	001	May 28, 2002
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AP			EQ 1MG BASE/ML	A075660	001	May 28, 2002
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AP	!	HIKMA FARMACEUTICA	EQ 1MG BASE/ML	A077966	001	Dec 03, 2010
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AP		HOSPIRA	EQ 1MG BASE/ML	A203280	001	Sep 03, 2014
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AP		MEITHEAL	EQ 1MG BASE/ML	A211671	001	Mar 24, 2020
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AP		SHANDONG	EQ 1MG BASE/ML	A216373	001	Jan 23, 2023
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MILRINONE LACTATE IN DEXTROSE 5%

AP		EUGIA PHARMA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A209666	001	Sep 03, 2020
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AP			EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A209666	002	Sep 03, 2020
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AP		WOODWARD	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A077151	002	Jul 20, 2005
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MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP	!	BAXTER HLTHCARE	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075834	001	May 28, 2002
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AP	!		EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A075834	002	May 28, 2002
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AP		GLAND PHARMA LTD	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A213585	001	Jul 16, 2020
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AP			EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A213585	002	Jul 16, 2020
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AP		WEST-WARD PHARMS	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A078113	001	May 21, 2008
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AP		INT	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A078113	002	May 21, 2008
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MILRINONE LACTATE IN PLASTIC CONTAINER

AP		HIKMA FARMACEUTICA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A090038	001	Jan 21, 2010
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AP			EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A090038	002	Jan 21, 2010
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PRESCRIPTION DRUG PRODUCT LIST

MILTEFOSINE

CAPSULE;ORAL

IMPAVIDO

+! KNIGHT THERAPS 50MG N204684 001 Mar 19, 2014

MINOCYCLINE HYDROCHLORIDE

AEROSOL, FOAM;TOPICAL

AMZEEQ

+! JOURNEY EQ 4% BASE N212379 001 Oct 18, 2019

ZILXI

+! JOURNEY EQ 1.5% BASE N213690 001 May 28, 2020

CAPSULE;ORAL

MINOCINAB + BAUSCH EQ 50MG BASE N050649 001 May 31, 1990AB + EQ 100MG BASE N050649 002 May 31, 1990MINOCYCLINE HYDROCHLORIDEAB AUROBINDO PHARMA EQ 50MG BASE A065470 001 Mar 11, 2008AB EQ 75MG BASE A065470 002 Mar 11, 2008AB EQ 100MG BASE A065470 003 Mar 11, 2008AB IMPAX LABS EQ 50MG BASE A065005 001 Mar 23, 1999AB EQ 75MG BASE A065005 003 Apr 18, 2001AB EQ 100MG BASE A065005 002 Mar 23, 1999AB SUN PHARM INDS INC EQ 50MG BASE A090867 001 May 13, 2013AB EQ 75MG BASE A090867 002 May 13, 2013AB EQ 100MG BASE A090867 003 May 13, 2013AB TORRENT EQ 50MG BASE A065062 001 Nov 30, 2000AB EQ 75MG BASE A065062 002 Nov 30, 2000AB EQ 100MG BASE A065062 003 Nov 30, 2000AB WATSON LABS EQ 75MG BASE A063065 002 Jun 10, 1999AB EQ 100MG BASE A063065 001 Dec 30, 1991AB WATSON LABS TEVA EQ 50MG BASE A063181 001 Dec 30, 1991AB ZYDUS EQ 50MG BASE A063011 001 Mar 02, 1992AB EQ 75MG BASE A063009 002 Aug 12, 2003AB ! EQ 100MG BASE A063009 001 Mar 02, 1992

INJECTABLE;INJECTION

MINOCIN

+! REMPEX EQ 100MG BASE/VIAL N050444 001

POWDER, EXTENDED RELEASE;DENTAL

ARESTIN

+! ORAPHARMA EQ 1MG BASE N050781 001 Feb 16, 2001

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDEAB AUROBINDO PHARMA EQ 50MG BASE A213662 001 May 01, 2020AB LTD EQ 75MG BASE A213662 002 May 01, 2020AB EQ 100MG BASE A213662 003 May 01, 2020AB BEXIMCO PHARMS USA EQ 50MG BASE A215466 001 May 27, 2022AB EQ 75MG BASE A215466 002 May 27, 2022AB EQ 100MG BASE A215466 003 May 27, 2022AB DR REDDYS LABS LTD EQ 50MG BASE A065436 001 Dec 26, 2007AB EQ 75MG BASE A065436 002 Dec 26, 2007AB EQ 100MG BASE A065436 003 Dec 26, 2007AB STRIDES PHARMA EQ 50MG BASE A065131 001 Apr 16, 2003AB EQ 75MG BASE A065131 002 Apr 16, 2003AB ! EQ 100MG BASE A065131 003 Apr 16, 2003AB SUN PHARM EQ 50MG BASE A090217 001 Jan 29, 2016AB INDUSTRIES EQ 75MG BASE A090217 002 Jan 29, 2016AB EQ 100MG BASE A090217 003 Jan 29, 2016AB TORRENT EQ 50MG BASE A065156 001 Jan 06, 2004AB EQ 75MG BASE A065156 002 Jan 06, 2004AB EQ 100MG BASE A065156 003 Jan 06, 2004

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDEAB ALKEM LABS LTD EQ 45MG BASE A204453 001 Sep 28, 2016AB EQ 55MG BASE A204453 008 Dec 19, 2019AB EQ 65MG BASE A204453 006 Mar 16, 2018AB EQ 80MG BASE A204453 002 Sep 28, 2016AB EQ 90MG BASE A204453 003 Sep 28, 2016AB EQ 105MG BASE A204453 004 Sep 28, 2016AB EQ 115MG BASE A204453 007 Mar 16, 2018AB EQ 135MG BASE A204453 005 Sep 28, 2016AB AUROBINDO PHARMA EQ 45MG BASE A202261 001 Nov 19, 2012

PRESCRIPTION DRUG PRODUCT LIST

MINOCYCLINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDE

LTD

<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A202261 008</u>	Aug 21, 2019
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A202261 002</u>	Sep 28, 2018
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A202261 006</u>	Jun 13, 2016
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A202261 003</u>	Nov 19, 2012
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A202261 007</u>	Jun 13, 2016
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A202261 004</u>	Sep 28, 2018
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A202261 005</u>	Nov 19, 2012
<u>AB</u>	LUPIN LTD	<u>EQ 45MG BASE</u>	<u>A091424 001</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A091424 002</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A091424 003</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A091424 004</u>	Nov 30, 2011
<u>AB</u>	SANDOZ	<u>EQ 45MG BASE</u>	<u>A090422 001</u>	Aug 13, 2009
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A090422 002</u>	Aug 13, 2009
<u>AB</u>	!	<u>EQ 135MG BASE</u>	<u>A090422 003</u>	Aug 13, 2009
<u>AB</u>	SIDMAK LABS INDIA	<u>EQ 45MG BASE</u>	<u>A204394 001</u>	Dec 30, 2015
<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A204394 002</u>	Oct 07, 2022
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A204394 003</u>	Oct 07, 2022
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204394 004</u>	Dec 30, 2015
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A204394 005</u>	Dec 30, 2015
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A204394 006</u>	Oct 07, 2022
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A204394 007</u>	Dec 30, 2015
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 45MG BASE</u>	<u>A091118 001</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A091118 003</u>	Dec 03, 2019
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A091118 004</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A091118 005</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A091118 006</u>	Sep 25, 2014
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A091118 007</u>	Dec 03, 2019
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A091118 008</u>	Sep 25, 2014
<u>AB</u>	ZYDUS PHARMS	<u>EQ 45MG BASE</u>	<u>A203553 001</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 55MG BASE</u>	<u>A203553 002</u>	Jun 16, 2023
<u>AB</u>		<u>EQ 65MG BASE</u>	<u>A203553 003</u>	Jun 16, 2023
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A203553 004</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A203553 005</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A203553 006</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 115MG BASE</u>	<u>A203553 007</u>	Jun 16, 2023
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A203553 008</u>	Nov 16, 2017

SOLODYN

<u>AB</u>	+	BAUSCH	<u>EQ 55MG BASE</u>	<u>N050808 008</u>	Aug 27, 2010
<u>AB</u>	+		<u>EQ 65MG BASE</u>	<u>N050808 004</u>	Jul 23, 2009
<u>AB</u>	+		<u>EQ 80MG BASE</u>	<u>N050808 007</u>	Aug 27, 2010
<u>AB</u>	+		<u>EQ 105MG BASE</u>	<u>N050808 006</u>	Aug 27, 2010
<u>AB</u>	+	!	<u>EQ 115MG BASE</u>	<u>N050808 005</u>	Jul 23, 2009

MINOLIRA

		EPI HLTH	EQ 105MG BASE	N209269 001	May 08, 2017
	!		EQ 135MG BASE	N209269 002	May 08, 2017

MINOXIDIL

TABLET;ORAL

MINOXIDIL

<u>AB</u>		PAR PHARM	<u>2.5MG</u>	<u>A071826 001</u>	Nov 14, 1988
<u>AB</u>			<u>10MG</u>	<u>A071839 001</u>	Nov 14, 1988
<u>AB</u>		SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A072709 002</u>	Dec 14, 1995
<u>AB</u>			<u>10MG</u>	<u>A072709 001</u>	Dec 14, 1995
<u>AB</u>		WATSON LABS	<u>2.5MG</u>	<u>A071344 001</u>	Mar 03, 1987
<u>AB</u>	!		<u>10MG</u>	<u>A071345 001</u>	Mar 03, 1987

MIRABEGRON

FOR SUSPENSION, EXTENDED RELEASE;ORAL

MYRBETRIQ GRANULES

	+	APGDI	8MG/ML	N213801 001	Mar 25, 2021
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TABLET, EXTENDED RELEASE;ORAL

MIRABEGRON

<u>AB</u>		LUPIN LTD	<u>25MG</u>	<u>A209485 001</u>	Sep 28, 2022
<u>AB</u>			<u>50MG</u>	<u>A209485 002</u>	Sep 28, 2022

MYRBETRIQ

<u>AB</u>	+	APGDI	<u>25MG</u>	<u>N202611 001</u>	Jun 28, 2012
<u>AB</u>	+		<u>50MG</u>	<u>N202611 002</u>	Jun 28, 2012

PRESCRIPTION DRUG PRODUCT LIST

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

<u>AB</u>	APOTEX INC	<u>15MG</u>	<u>A077666 001</u>	Aug 22, 2007
<u>AB</u>		<u>30MG</u>	<u>A077666 002</u>	Aug 22, 2007
<u>AB</u>		<u>45MG</u>	<u>A077666 003</u>	Aug 22, 2007
<u>AB</u>	AUROBINDO	<u>7.5MG</u>	<u>A076921 001</u>	Oct 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076921 002</u>	Oct 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076921 003</u>	Oct 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076921 004</u>	Oct 22, 2004
<u>AB</u>	MYLAN	<u>15MG</u>	<u>A076122 001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076122 002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076122 003</u>	Jun 19, 2003
<u>AB</u>	PRASCO	<u>7.5MG</u>	<u>A216751 001</u>	Jan 18, 2023
<u>AB</u>		<u>15MG</u>	<u>A216751 002</u>	Jan 18, 2023
<u>AB</u>		<u>30MG</u>	<u>A216751 003</u>	Jan 18, 2023
<u>AB</u>		<u>45MG</u>	<u>A216751 004</u>	Jan 18, 2023
<u>AB</u>	! SUN PHARM INDS INC	<u>7.5MG</u>	<u>A076541 004</u>	Apr 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076541 001</u>	Apr 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076541 002</u>	Apr 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076541 003</u>	Apr 22, 2004
<u>AB</u>	TEVA	<u>15MG</u>	<u>A076119 001</u>	Jan 24, 2003
<u>AB</u>		<u>30MG</u>	<u>A076119 002</u>	Jan 24, 2003
<u>AB</u>		<u>45MG</u>	<u>A076119 003</u>	Jun 19, 2003
<u>AB</u>	UPSHER SMITH LABS	<u>15MG</u>	<u>A076219 001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076219 002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076219 003</u>	Jun 19, 2003

REMERON

<u>AB</u>	+! ORGANON	<u>15MG</u>	<u>N020415 001</u>	Jun 14, 1996
<u>AB</u>	+	<u>30MG</u>	<u>N020415 002</u>	Jun 14, 1996

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

<u>AB</u>	AUROBINDO PHARMA	<u>15MG</u>	<u>A077376 002</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>A077376 003</u>	Dec 08, 2005
<u>AB</u>		<u>45MG</u>	<u>A077376 004</u>	Feb 28, 2006
<u>AB</u>	SQUARE PHARMS	<u>15MG</u>	<u>A205798 001</u>	Jun 01, 2017
<u>AB</u>		<u>30MG</u>	<u>A205798 002</u>	Jun 01, 2017
<u>AB</u>		<u>45MG</u>	<u>A205798 003</u>	Jun 01, 2017

REMERON SOLTAB

<u>AB</u>	+! ORGANON USA ORGANON	<u>15MG</u>	<u>N021208 001</u>	Jan 12, 2001
<u>AB</u>	+	<u>30MG</u>	<u>N021208 002</u>	Jan 12, 2001
<u>AB</u>	+	<u>45MG</u>	<u>N021208 003</u>	Jan 12, 2001

MISOPROSTOL

TABLET; ORAL

CYTOTEK

<u>AB</u>	+ PFIZER	<u>0.1MG</u>	<u>N019268 003</u>	Sep 21, 1990
<u>AB</u>	+!	<u>0.2MG</u>	<u>N019268 001</u>	Dec 27, 1988

MISOPROSTOL

<u>AB</u>	ANI PHARMS	<u>0.1MG</u>	<u>A076095 001</u>	Jul 10, 2002
<u>AB</u>		<u>0.2MG</u>	<u>A076095 002</u>	Jul 10, 2002
<u>AB</u>	MICRO LABS	<u>0.1MG</u>	<u>A216872 001</u>	Oct 17, 2022
<u>AB</u>		<u>0.2MG</u>	<u>A216872 002</u>	Oct 17, 2022

MITAPIVAT SULFATE

TABLET; ORAL

PYRUKYND

+	AGIOS PHARMS INC	EQ 5MG BASE	N216196 001	Feb 17, 2022
+		EQ 20MG BASE	N216196 002	Feb 17, 2022
+	!	EQ 50MG BASE	N216196 003	Feb 17, 2022

MITOMYCIN

FOR SOLUTION; TOPICAL

MITOSOL

+	MOBIUS THERAP	0.2MG/VIAL	N022572 001	Feb 07, 2012
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INJECTABLE; INJECTION

MITOMYCIN

<u>AP</u>	! ACCORD HLTHCARE	<u>5MG/VIAL</u>	<u>A064144 001</u>	Apr 30, 1998
<u>AP</u>	!	<u>20MG/VIAL</u>	<u>A064144 002</u>	Apr 30, 1998
<u>AP</u>	!	<u>40MG/VIAL</u>	<u>A064144 003</u>	Aug 11, 2009
<u>AP</u>	EUGIA PHARMA	<u>5MG/VIAL</u>	<u>A216732 001</u>	Oct 30, 2023
<u>AP</u>		<u>20MG/VIAL</u>	<u>A216732 002</u>	Oct 30, 2023
<u>AP</u>		<u>40MG/VIAL</u>	<u>A216732 003</u>	Oct 30, 2023
<u>AP</u>	GLAND PHARMA LTD	<u>5MG/VIAL</u>	<u>A215687 001</u>	Oct 20, 2021

PRESCRIPTION DRUG PRODUCT LIST

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

<u>AP</u>		<u>20MG/VIAL</u>	<u>A215687 002</u>	Oct 20, 2021
<u>AP</u>		<u>40MG/VIAL</u>	<u>A216648 001</u>	Nov 22, 2022
<u>AP</u>	HIKMA	<u>5MG/VIAL</u>	<u>A064180 001</u>	Dec 23, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A064117 002</u>	Apr 19, 1995
<u>AP</u>		<u>20MG/VIAL</u>	<u>A064180 002</u>	Dec 23, 1999
<u>AP</u>		<u>40MG/VIAL</u>	<u>A064117 003</u>	Jun 02, 1999
<u>AP</u>	MEITHEAL	<u>5MG/VIAL</u>	<u>A214505 001</u>	Sep 08, 2022
<u>AP</u>		<u>20MG/VIAL</u>	<u>A214505 002</u>	Sep 08, 2022
<u>AP</u>		<u>40MG/VIAL</u>	<u>A214504 001</u>	Sep 06, 2022
<u>AP</u>	RK PHARMA	<u>5MG/VIAL</u>	<u>A202670 001</u>	Oct 13, 2017
<u>AP</u>		<u>20MG/VIAL</u>	<u>A202670 002</u>	Oct 13, 2017
<u>AP</u>		<u>40MG/VIAL</u>	<u>A203386 001</u>	Oct 13, 2017
	POWDER; PYELOALYCEAL JELMYTO			
	+! UROGEN PHARMA	40MG/VIAL	N211728 001	Apr 15, 2020

MITOTANE

TABLET; ORAL

LYSODREN

+! HRA PHARMA

500MG

N016885 001

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077496 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077496 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077496 003</u>	Apr 11, 2006
<u>AP</u>	HIKMA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076611 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076611 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076611 003</u>	Apr 11, 2006
<u>AP</u>	! HOSPIRA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076871 001</u>	Apr 11, 2006
<u>AP</u>	!	<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076871 002</u>	Apr 11, 2006
<u>AP</u>	!	<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076871 003</u>	Apr 11, 2006
<u>AP</u>	MEITHEAL	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077356 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077356 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077356 003</u>	Apr 11, 2006

MIVACURIUM CHLORIDE

SOLUTION; INTRAVENOUS

MIVACURIUM CHLORIDE

WOODWARD

EQ 10MG BASE/5ML (EQ 2MG BASE/ML)

A209708 001 Oct 12, 2021

EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

A209708 002 Oct 12, 2021

MOBOCERTINIB SUCCINATE

CAPSULE; ORAL

EXKIVITY

+! TAKEDA PHARMS USA

EQ 40MG BASE

N215310 001 Sep 15, 2021

MODAFINIL

TABLET; ORAL

MODAFINIL

<u>AB</u>	ALEMBIC	<u>100MG</u>	<u>A202700 001</u>	Oct 18, 2012
<u>AB</u>		<u>200MG</u>	<u>A202700 002</u>	Oct 18, 2012
<u>AB</u>	APOTEX	<u>100MG</u>	<u>A077667 001</u>	Feb 03, 2014
<u>AB</u>		<u>200MG</u>	<u>A077667 002</u>	Feb 03, 2014
<u>AB</u>	APPCO	<u>100MG</u>	<u>A207196 001</u>	Aug 16, 2017
<u>AB</u>		<u>200MG</u>	<u>A207196 002</u>	Aug 16, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A202566 001</u>	Sep 27, 2012
<u>AB</u>		<u>200MG</u>	<u>A202566 002</u>	Sep 27, 2012
<u>AB</u>	CADILA	<u>100MG</u>	<u>A209966 001</u>	Sep 14, 2017
<u>AB</u>		<u>200MG</u>	<u>A209966 002</u>	Sep 14, 2017
<u>AB</u>	ORBION PHARMS	<u>100MG</u>	<u>A078963 001</u>	Sep 26, 2012
<u>AB</u>		<u>200MG</u>	<u>A078963 002</u>	Sep 26, 2012
<u>AB</u>	WATSON LABS INC	<u>100MG</u>	<u>A076715 001</u>	Nov 01, 2012
<u>AB</u>		<u>200MG</u>	<u>A076715 002</u>	Nov 01, 2012
	<u>PROVIGIL</u>			
<u>AB</u>	+ CEPHALON	<u>100MG</u>	<u>N020717 001</u>	Dec 24, 1998
<u>AB</u>	+!	<u>200MG</u>	<u>N020717 002</u>	Dec 24, 1998

PRESCRIPTION DRUG PRODUCT LIST

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>7.5MG</u>	<u>A078454</u>	<u>001</u>	Jun 02, 2008
<u>AB</u>		<u>15MG</u>	<u>A078454</u>	<u>002</u>	Jun 02, 2008
<u>AB</u>	CHARTWELL RX	<u>7.5MG</u>	<u>A077536</u>	<u>001</u>	Nov 30, 2006
<u>AB</u>		<u>15MG</u>	<u>A077536</u>	<u>002</u>	Nov 30, 2006
<u>AB</u>	GLENMARK GENERICS	<u>7.5MG</u>	<u>A090416</u>	<u>001</u>	Mar 30, 2010
<u>AB</u>		<u>15MG</u>	<u>A090416</u>	<u>002</u>	Mar 30, 2010
<u>AB</u>	TEVA	<u>7.5MG</u>	<u>A076204</u>	<u>001</u>	May 08, 2003
<u>AB</u>	!	<u>15MG</u>	<u>A076204</u>	<u>002</u>	May 08, 2003

MOLINDONE HYDROCHLORIDE

TABLET; ORAL

MOLINDONE HYDROCHLORIDE

EPIC PHARMA LLC

5MG

A090453 001 Mar 20, 2015

10MG

A090453 002 Mar 20, 2015

!

25MG

A090453 003 Mar 20, 2015

MOMELOTINIB DIHYDROCHLORIDE

TABLET; ORAL

OJJAARA

+	GLAXOSMITHKLINE	EQ 100MG BASE	N216873 001	Sep 15, 2023
+		EQ 150MG BASE	N216873 002	Sep 15, 2023
+	!	EQ 200MG BASE	N216873 003	Sep 15, 2023

MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

ASMANEX HFA

+	ORGANON LLC	0.05MG/INH	N205641 003	Aug 12, 2019
+		0.10MG/INH	N205641 001	Apr 25, 2014
+	!	0.20MG/INH	N205641 002	Apr 25, 2014

CREAM; TOPICAL

MOMETASONE FUROATE

<u>AB</u>	COSETTE	<u>0.1%</u>	<u>A077447</u>	<u>001</u>	May 22, 2006
<u>AB</u>	!	<u>0.1%</u>	<u>A078541</u>	<u>001</u>	May 28, 2008
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076679</u>	<u>001</u>	Dec 21, 2004

IMPLANT; IMPLANTATION

SINUVA

+	INTERSECT ENT INC	1.35MG	N209310 001	Dec 08, 2017
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LOTION; TOPICAL

MOMETASONE FUROATE

<u>AB</u>	COSETTE	<u>0.1%</u>	<u>A077678</u>	<u>001</u>	Nov 21, 2007
<u>AB</u>	FOUGERA PHARMS	<u>0.1%</u>	<u>A075919</u>	<u>001</u>	Nov 29, 2007
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A090506</u>	<u>001</u>	Aug 09, 2010
<u>AB</u>	!	<u>0.1%</u>	<u>A077180</u>	<u>001</u>	Apr 06, 2005
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076788</u>	<u>001</u>	Mar 15, 2006

OINTMENT; TOPICAL

MOMETASONE FUROATE

<u>AB</u>	COSETTE	<u>0.1%</u>	<u>A077401</u>	<u>001</u>	Jun 20, 2006
<u>AB</u>	FOUGERA PHARMS	<u>0.1%</u>	<u>A077061</u>	<u>001</u>	Mar 28, 2005
<u>AB</u>	!	<u>0.1%</u>	<u>A078571</u>	<u>001</u>	May 28, 2008
<u>AB</u>	PADAGIS US	<u>0.1%</u>	<u>A076067</u>	<u>001</u>	Mar 18, 2002

POWDER; INHALATION

ASMANEX TWISTHALER

+	ORGANON LLC	0.11MG/INH	N021067 002	Feb 01, 2008
+	!	0.22MG/INH	N021067 001	Mar 30, 2005

SPRAY, METERED; NASAL

MOMETASONE FUROATE

<u>AB</u>	AMNEAL PHARMS	<u>0.05MG/SPRAY</u>	<u>A207989</u>	<u>001</u>	Apr 03, 2017
<u>AB</u>	!	<u>0.05MG/SPRAY</u>	<u>A091161</u>	<u>001</u>	Mar 22, 2016

MOMETASONE FUROATE; OLOPATADINE HYDROCHLORIDE

SPRAY, METERED; NASAL

RYALTRIS

+	GLENMARK SPECIALTY	0.025MG/SPRAY; 0.665MG/SPRAY	N211746 001	Jan 13, 2022
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MONOMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL

BAFIERTAM

+	BANNER LIFE SCIENCES	95MG	N210296 001	Apr 28, 2020
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PRESCRIPTION DRUG PRODUCT LIST

MONTELUKAST SODIUM

GRANULE; ORAL

MONTELUKAST SODIUM

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A203438 001</u>	Jul 31, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A213471 001</u>	Feb 18, 2020
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A202906 001</u>	Sep 17, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 4MG BASE/PACKET</u>	<u>A090955 001</u>	Aug 03, 2012
<u>AB</u>	TORRENT	<u>EQ 4MG BASE/PACKET</u>	<u>A210431 001</u>	Jul 31, 2018

SINGULAIR

<u>AB</u>	+! ORGANON	<u>EQ 4MG BASE/PACKET</u>	<u>N021409 001</u>	Jul 26, 2002
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TABLET; ORAL

MONTELUKAST SODIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 10MG BASE</u>	<u>A202717 001</u>	Sep 21, 2012
<u>AB</u>	AMNEAL PHARMS	<u>EQ 10MG BASE</u>	<u>A204604 001</u>	Sep 04, 2015
<u>AB</u>	ANBISON LAB	<u>EQ 10MG BASE</u>	<u>A205683 001</u>	Jan 12, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE</u>	<u>A202468 001</u>	Aug 03, 2012
<u>AB</u>	CHARTWELL MOLECULAR	<u>EQ 10MG BASE</u>	<u>A201522 001</u>	Aug 03, 2012
<u>AB</u>	CIPLA	<u>EQ 10MG BASE</u>	<u>A207463 001</u>	Oct 28, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A201582 001</u>	Aug 06, 2012
<u>AB</u>	GLENMARK GENERICS	<u>EQ 10MG BASE</u>	<u>A090926 001</u>	Aug 03, 2012
<u>AB</u>	GRAVITI PHARMS	<u>EQ 10MG BASE</u>	<u>A209012 001</u>	Apr 24, 2017
<u>AB</u>	HETERO LABS LTD V	<u>EQ 10MG BASE</u>	<u>A202843 001</u>	Sep 10, 2014
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A203366 001</u>	Sep 11, 2014
<u>AB</u>	SANDOZ INC	<u>EQ 10MG BASE</u>	<u>A200889 001</u>	Aug 03, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 10MG BASE</u>	<u>A078605 001</u>	Aug 03, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A201515 001</u>	Aug 03, 2012
<u>AB</u>	UNICHEM	<u>EQ 10MG BASE</u>	<u>A204290 001</u>	Oct 08, 2015
<u>AB</u>	UNIMARK REMEDIES LTD	<u>EQ 10MG BASE</u>	<u>A202859 001</u>	Oct 30, 2014

SINGULAIR

<u>AB</u>	+! ORGANON	<u>EQ 10MG BASE</u>	<u>N020829 002</u>	Feb 20, 1998
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TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>EQ 4MG BASE</u>	<u>A205107 001</u>	Sep 04, 2020
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A205107 002</u>	Sep 04, 2020
<u>AB</u>	ANBISON LAB	<u>EQ 4MG BASE</u>	<u>A205695 001</u>	Nov 05, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A205695 002</u>	Nov 05, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A202096 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A202096 002</u>	Aug 03, 2012
<u>AB</u>	CHARTWELL MOLECULAR	<u>EQ 4MG BASE</u>	<u>A207464 001</u>	Dec 06, 2018
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A207464 002</u>	Dec 06, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A201581 001</u>	Aug 06, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A201581 002</u>	Aug 06, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 4MG BASE</u>	<u>A204093 001</u>	May 22, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A204093 002</u>	May 22, 2015
<u>AB</u>	LANNETT CO INC	<u>EQ 4MG BASE</u>	<u>A200405 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A200405 002</u>	Aug 03, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 4MG BASE</u>	<u>A203582 001</u>	Mar 12, 2015
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203582 002</u>	Mar 12, 2015
<u>AB</u>	RISING	<u>EQ 4MG BASE</u>	<u>A209011 001</u>	Apr 18, 2017
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A209011 002</u>	Apr 18, 2017
<u>AB</u>	SANDOZ INC	<u>EQ 4MG BASE</u>	<u>A091414 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A091414 002</u>	Aug 03, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 4MG BASE</u>	<u>A078723 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078723 002</u>	Aug 03, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 4MG BASE</u>	<u>A090984 001</u>	Aug 03, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090984 002</u>	Aug 03, 2012
<u>AB</u>	UNIMARK REMEDIES LTD	<u>EQ 4MG BASE</u>	<u>A203037 001</u>	Oct 30, 2014
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A203037 002</u>	Oct 30, 2014

SINGULAIR

<u>AB</u>	+ ORGANON	<u>EQ 4MG BASE</u>	<u>N020830 002</u>	Mar 03, 2000
<u>AB</u>	+!	<u>EQ 5MG BASE</u>	<u>N020830 001</u>	Feb 20, 1998

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

ACTAVIS ELIZABETH	30MG	A079040 001	Jan 16, 2013
	45MG	A079040 002	Jan 16, 2013
	60MG	A079040 003	Jan 16, 2013
	75MG	A079040 004	Jan 16, 2013

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

		90MG	A079040 005	Jan 16, 2013
!		120MG	A079040 006	Jan 16, 2013
!	UPSHER SMITH LABS	10MG	A202104 001	Jun 03, 2013
		20MG	A202104 002	Jun 03, 2013
		30MG	A202104 003	Jun 03, 2013
		50MG	A202104 004	Jun 03, 2013
		60MG	A202104 005	Jun 03, 2013
		80MG	A202104 006	Jun 03, 2013
!		100MG	A202104 007	Jun 03, 2013

INJECTABLE; INJECTION

DURAMORPH PF

<u>AP</u>	+!	HIKMA	<u>0.5MG/ML</u>	<u>N018565 001</u>	Sep 18, 1984
<u>AP</u>	+!		<u>1MG/ML</u>	<u>N018565 002</u>	Sep 18, 1984

INFUMORPH

<u>AP</u>	+!	HIKMA	<u>10MG/ML</u>	<u>N018565 003</u>	Jul 19, 1991
<u>AP</u>	+!		<u>25MG/ML</u>	<u>N018565 004</u>	Jul 19, 1991

MITIGO

<u>AP</u>		PIRAMAL CRITICAL	<u>10MG/ML</u>	<u>A204393 001</u>	Jul 16, 2018
<u>AP</u>			<u>25MG/ML</u>	<u>A204393 002</u>	Jul 16, 2018

MORPHINE SULFATE

<u>AP</u>		HIKMA	<u>2MG/ML</u>	<u>A211452 001</u>	Jan 12, 2023
<u>AP</u>	!		<u>4MG/ML</u>	<u>A205758 001</u>	May 21, 2015
<u>AP</u>			<u>4MG/ML</u>	<u>A211452 002</u>	Jan 12, 2023
<u>AP</u>	!		<u>8MG/ML</u>	<u>A205758 002</u>	May 21, 2015
<u>AP</u>			<u>8MG/ML</u>	<u>A211452 003</u>	Jan 12, 2023
<u>AP</u>			<u>10MG/ML</u>	<u>A205758 003</u>	May 21, 2015
<u>AP</u>			<u>10MG/ML</u>	<u>A211452 004</u>	Jan 12, 2023
<u>AP</u>		HOSPIRA	<u>0.5MG/ML</u>	<u>A071849 001</u>	May 11, 1988
<u>AP</u>			<u>0.5MG/ML</u>	<u>A073509 001</u>	Sep 30, 1992
<u>AP</u>			<u>1MG/ML</u>	<u>A073510 001</u>	Sep 30, 1992
<u>AP</u>	+!	HOSPIRA INC	<u>2MG/ML</u>	<u>N202515 001</u>	Nov 14, 2011
<u>AP</u>	+		<u>4MG/ML</u>	<u>N202515 002</u>	Nov 14, 2011
<u>AP</u>	+		<u>8MG/ML</u>	<u>N202515 003</u>	Nov 14, 2011
<u>AP</u>	+		<u>10MG/ML</u>	<u>N202515 004</u>	Nov 14, 2011
		HIKMA	15MG/ML	A211452 005	Jan 12, 2023
	+!	HOSPIRA INC	50MG/ML	N202515 006	Apr 29, 2021
		INTL MEDICATION SYS	1MG/ML	A202861 001	Apr 29, 2021

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MORPHINE SULFATE

+!	FRESENIUS KABI USA	2MG/ML (2MG/ML)	N204223 001	Oct 30, 2013
+!		4MG/ML (4MG/ML)	N204223 002	Oct 30, 2013
+!		5MG/ML (5MG/ML)	N204223 003	Oct 30, 2013
+!		8MG/ML (8MG/ML)	N204223 004	Oct 30, 2013
+!		10MG/ML (10MG/ML)	N204223 005	Oct 30, 2013

SOLUTION; ORAL

MORPHINE SULFATE

<u>AA</u>	+	HIKMA	<u>10MG/5ML</u>	<u>N022195 001</u>	Mar 17, 2008
<u>AA</u>	+		<u>20MG/5ML</u>	<u>N022195 002</u>	Mar 17, 2008
<u>AA</u>			<u>100MG/5ML</u>	<u>A208809 001</u>	Jul 06, 2017
<u>AA</u>	+!		<u>100MG/5ML</u>	<u>N022195 003</u>	Jan 25, 2010
<u>AA</u>		PADAGIS US	<u>100MG/5ML</u>	<u>A201574 001</u>	Aug 06, 2012
<u>AA</u>		PHARM ASSOC	<u>10MG/5ML</u>	<u>A206573 002</u>	Sep 12, 2023
<u>AA</u>			<u>100MG/5ML</u>	<u>A206573 001</u>	Nov 14, 2016
<u>AA</u>		RHODES PHARMS	<u>10MG/5ML</u>	<u>A206308 001</u>	Jun 22, 2017
<u>AA</u>			<u>20MG/5ML</u>	<u>A206420 001</u>	Jul 12, 2016
<u>AA</u>			<u>100MG/5ML</u>	<u>A206308 002</u>	Jun 22, 2017
<u>AA</u>		SPECGX LLC	<u>100MG/5ML</u>	<u>A202348 001</u>	Jul 15, 2011
<u>AA</u>		TRIS PHARMA INC	<u>10MG/5ML</u>	<u>A203518 001</u>	May 12, 2015
<u>AA</u>			<u>100MG/5ML</u>	<u>A203518 002</u>	May 12, 2015

TABLET; ORAL

MORPHINE SULFATE

<u>AB</u>		ALKEM LABS LTD	<u>15MG</u>	<u>A212451 001</u>	Dec 03, 2020
<u>AB</u>			<u>30MG</u>	<u>A212451 002</u>	Dec 03, 2020
<u>AB</u>	+	HIKMA	<u>15MG</u>	<u>N022207 001</u>	Mar 17, 2008
<u>AB</u>	+!		<u>30MG</u>	<u>N022207 002</u>	Mar 17, 2008
<u>AB</u>		SPECGX LLC	<u>15MG</u>	<u>A215194 001</u>	Aug 21, 2023
<u>AB</u>			<u>30MG</u>	<u>A215194 002</u>	Aug 21, 2023
<u>AB</u>		UPSHER SMITH LABS	<u>15MG</u>	<u>A210610 001</u>	Jul 22, 2019
<u>AB</u>			<u>30MG</u>	<u>A210610 002</u>	Jul 22, 2019

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

MORPHINE SULFATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>15MG</u>	<u>A203849 001</u>	Apr 06, 2015
<u>AB</u>		<u>30MG</u>	<u>A203849 002</u>	Apr 06, 2015
<u>AB</u>		<u>60MG</u>	<u>A203849 003</u>	Apr 06, 2015
<u>AB</u>		<u>100MG</u>	<u>A203849 004</u>	Apr 06, 2015
<u>AB</u>		<u>200MG</u>	<u>A203849 005</u>	Apr 06, 2015
<u>AB</u>	DAVA PHARMS INC	<u>15MG</u>	<u>A075407 001</u>	Jan 28, 2000
<u>AB</u>	NOVEL LABS INC	<u>15MG</u>	<u>A203602 001</u>	Dec 16, 2015
<u>AB</u>		<u>30MG</u>	<u>A203602 002</u>	Dec 16, 2015
<u>AB</u>		<u>60MG</u>	<u>A203602 003</u>	Dec 16, 2015
<u>AB</u>		<u>100MG</u>	<u>A203602 004</u>	Dec 16, 2015
<u>AB</u>		<u>200MG</u>	<u>A203602 005</u>	Dec 16, 2015
<u>AB</u>	RHODES PHARMS	<u>15MG</u>	<u>A074862 001</u>	Jul 07, 1998
<u>AB</u>		<u>30MG</u>	<u>A074862 002</u>	Jul 07, 1998
<u>AB</u>		<u>60MG</u>	<u>A074862 003</u>	Jul 07, 1998
<u>AB</u>		<u>100MG</u>	<u>A074769 001</u>	Jul 02, 1998
<u>AB</u>		<u>200MG</u>	<u>A074769 002</u>	Jul 02, 1998
<u>AB</u>	SPECGX LLC	<u>15MG</u>	<u>A076412 001</u>	Jul 31, 2003
<u>AB</u>		<u>30MG</u>	<u>A076412 002</u>	Jul 31, 2003
<u>AB</u>		<u>60MG</u>	<u>A076412 003</u>	Jul 31, 2003
<u>AB</u>		<u>100MG</u>	<u>A076438 001</u>	Jul 03, 2003
<u>AB</u>		<u>200MG</u>	<u>A076438 002</u>	Jul 03, 2003
<u>AB</u>	STRIDES PHARMA	<u>15MG</u>	<u>A075295 001</u>	Oct 28, 1998
<u>AB</u>		<u>30MG</u>	<u>A075295 002</u>	Oct 28, 1998
<u>AB</u>		<u>60MG</u>	<u>A075295 003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>A075295 004</u>	Sep 15, 2000
<u>AB</u>		<u>200MG</u>	<u>A075295 005</u>	Sep 15, 2000
<u>AB</u>	SUN PHARM INDS LTD	<u>15MG</u>	<u>A078761 001</u>	May 11, 2012
<u>AB</u>		<u>30MG</u>	<u>A078761 002</u>	May 11, 2012
<u>AB</u>		<u>60MG</u>	<u>A078761 003</u>	May 11, 2012
<u>AB</u>		<u>100MG</u>	<u>A078761 004</u>	May 11, 2012
<u>AB</u>		<u>200MG</u>	<u>A078761 005</u>	May 11, 2012

MS CONTIN

<u>AB</u>	+	PURDUE PHARMA LP	<u>15MG</u>	<u>N019516 003</u>	Sep 12, 1989
<u>AB</u>	+		<u>30MG</u>	<u>N019516 001</u>	May 29, 1987
<u>AB</u>	+		<u>60MG</u>	<u>N019516 002</u>	Apr 08, 1988
<u>AB</u>	+	!	<u>100MG</u>	<u>N019516 004</u>	Jan 16, 1990
<u>AB</u>	+		<u>200MG</u>	<u>N019516 005</u>	Nov 08, 1993

MOTIXAFORTIDE ACETATE

POWDER; SUBCUTANEOUS

APHEXDA

+	!	BIOLINERX LTD	EQ 62MG BASE/VIAL	N217159 001	Sep 08, 2023
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MOXIDECTIN

TABLET; ORAL

MOXIDECTIN

+	!	MDGH	2MG	N210867 001	Jun 13, 2018
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MOXIFLOXACIN HYDROCHLORIDE

SOLUTION; INTRAVENOUS

MOXIFLOXACIN HYDROCHLORIDE

+	!	FRESENIUS KABI USA	EQ 400MG BASE/250ML (EQ 1.6MG BASE/ML)	N205572 001	Apr 03, 2015
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MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

!		MYLAN LABS LTD	400MG/250ML (1.6MG/ML)	A205833 001	May 05, 2017
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SOLUTION/DROPS; OPHTHALMIC

MOXIFLOXACIN HYDROCHLORIDE

<u>AT1</u>		ALEMBIC	<u>EQ 0.5% BASE</u>	<u>A209469 001</u>	Feb 13, 2019
<u>AT1</u>		APOTEX	<u>EQ 0.5% BASE</u>	<u>A090080 001</u>	Jun 30, 2017
<u>AT1</u>		EUGIA PHARMA	<u>EQ 0.5% BASE</u>	<u>A206242 001</u>	Oct 04, 2017
<u>AT1</u>		GLAND PHARMA LTD	<u>EQ 0.5% BASE</u>	<u>A208778 001</u>	Mar 30, 2020
<u>AT1</u>		INDOCO	<u>EQ 0.5% BASE</u>	<u>A202525 001</u>	Mar 06, 2015
<u>AT1</u>		LUPIN LTD	<u>EQ 0.5% BASE</u>	<u>A202867 001</u>	Sep 04, 2014
<u>AT1</u>		MYLAN	<u>EQ 0.5% BASE</u>	<u>A206447 001</u>	Mar 30, 2020
<u>AT1</u>		UPSHER SMITH LABS	<u>EQ 0.5% BASE</u>	<u>A212616 001</u>	Feb 10, 2021

VIGAMOX

<u>AT1</u>	+	!	NOVARTIS	<u>EQ 0.5% BASE</u>	<u>N021598 001</u>	Apr 15, 2003
		!	LUPIN LTD	EQ 0.5% BASE	A204079 001	May 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

MOXIFLOXACIN HYDROCHLORIDE

TABLET; ORAL

MOXIFLOXACIN HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 400MG BASE</u>	<u>A202632 001</u>	Mar 04, 2014
<u>AB</u>	CHARTWELL RX	<u>EQ 400MG BASE</u>	<u>A207285 001</u>	Feb 13, 2017
<u>AB</u>	CROSSMEDIKA SA	<u>EQ 400MG BASE</u>	<u>A205348 001</u>	Jan 14, 2016
<u>AB</u>	! DR REDDYS	<u>EQ 400MG BASE</u>	<u>A076938 001</u>	Mar 04, 2014
<u>AB</u>	HETERO LABS LTD V	<u>EQ 400MG BASE</u>	<u>A204836 001</u>	Mar 02, 2023
<u>AB</u>	MSN	<u>EQ 400MG BASE</u>	<u>A208682 001</u>	Sep 22, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 400MG BASE</u>	<u>A077437 001</u>	Feb 18, 2014

MUPIROCIN

OINTMENT; TOPICAL

MUPIROCIN

<u>AB</u>	FOUGERA PHARMS	<u>2%</u>	<u>A065192 001</u>	Nov 30, 2005
<u>AB</u>	GLENMARK PHARMS	<u>2%</u>	<u>A090480 001</u>	Jun 08, 2011
<u>AB</u>	! PADAGIS ISRAEL	<u>2%</u>	<u>A065123 001</u>	Nov 07, 2003
<u>AB</u>	TARO	<u>2%</u>	<u>A065170 001</u>	Sep 23, 2005
<u>AB</u>	TEVA	<u>2%</u>	<u>A065085 001</u>	Nov 07, 2003

CENTANY

BX	PADAGIS US	2%	N050788 001	Dec 04, 2002
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MUPIROCIN CALCIUM

CREAM; TOPICAL

MUPIROCIN

<u>AB</u>	ALEMBIC	<u>EQ 2% BASE</u>	<u>A213053 001</u>	Nov 16, 2021
<u>AB</u>	AMNEAL	<u>EQ 2% BASE</u>	<u>A214811 001</u>	Nov 15, 2022
<u>AB</u>	ENCUBE	<u>EQ 2% BASE</u>	<u>A213076 001</u>	Aug 31, 2021
<u>AB</u>	! GLENMARK PHARMS INC	<u>EQ 2% BASE</u>	<u>A201587 001</u>	Jan 24, 2013
<u>AB</u>	TARO	<u>EQ 2% BASE</u>	<u>A207116 001</u>	Apr 27, 2020

MYCOPHENOLATE MOFETIL

CAPSULE; ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>250MG</u>	<u>N050722 001</u>	May 03, 1995
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MYCOPHENOLATE MOFETIL

<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>	<u>A090253 001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>250MG</u>	<u>A200197 001</u>	Jun 13, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A217828 001</u>	Jan 05, 2024
<u>AB</u>	CONCORD BIOTECH LTD	<u>250MG</u>	<u>A210181 001</u>	Jan 08, 2019
<u>AB</u>	HIKMA	<u>250MG</u>	<u>A065410 001</u>	Jul 29, 2008
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A065520 001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065379 001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A090055 001</u>	Jun 10, 2010
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>A065491 001</u>	May 06, 2009
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>250MG</u>	<u>A204077 001</u>	Nov 13, 2017

FOR SUSPENSION; ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>200MG/ML</u>	<u>N050759 001</u>	Oct 01, 1998
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MYCOPHENOLATE MOFETIL

<u>AB</u>	ALKEM LABS LTD	<u>200MG/ML</u>	<u>A203005 001</u>	Nov 14, 2014
<u>AB</u>	AMNEAL	<u>200MG/ML</u>	<u>A214871 001</u>	Nov 02, 2021
<u>AB</u>	HETERO LABS LTD V	<u>200MG/ML</u>	<u>A213955 001</u>	Sep 11, 2023
<u>AB</u>	LANNETT CO INC	<u>200MG/ML</u>	<u>A214525 001</u>	Jul 29, 2021
<u>AB</u>	STRIDES PHARMA	<u>200MG/ML</u>	<u>A212634 001</u>	Aug 29, 2023
<u>AB</u>	TEVA PHARMS USA	<u>200MG/ML</u>	<u>A211272 001</u>	Jan 25, 2022
<u>AB</u>	VISTAPHARM	<u>200MG/ML</u>	<u>A210370 001</u>	Feb 12, 2019

TABLET; ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>500MG</u>	<u>N050723 001</u>	Jun 19, 1997
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MYCOPHENOLATE MOFETIL

<u>AB</u>	ACCORD HLTHCARE	<u>500MG</u>	<u>A065416 001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A091249 001</u>	Nov 04, 2011
<u>AB</u>	AMNEAL	<u>500MG</u>	<u>A090606 001</u>	Jul 16, 2010
<u>AB</u>	CONCORD BIOTECH LTD	<u>500MG</u>	<u>A212087 001</u>	Jul 31, 2020
<u>AB</u>	HIKMA	<u>500MG</u>	<u>A065413 001</u>	Jul 29, 2008
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A065521 001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065451 001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>500MG</u>	<u>A090456 001</u>	Jun 10, 2010
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>500MG</u>	<u>A204076 001</u>	Nov 16, 2017

PRESCRIPTION DRUG PRODUCT LIST

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

CELLCEPT

<u>AP</u>	<u>+!</u> ROCHE PALO	<u>500MG/VIAL</u>	<u>N050758</u>	<u>001</u>	Aug 12, 1998
<u>MYCOPHENOLATE MOFETIL HYDROCHLORIDE</u>					
<u>AP</u>	BPI LABS	<u>500MG/VIAL</u>	<u>A214283</u>	<u>001</u>	Jun 01, 2023
<u>AP</u>	MEITHEAL	<u>500MG/VIAL</u>	<u>A212130</u>	<u>001</u>	Jan 15, 2021
<u>AP</u>	MYLAN LABS LTD	<u>500MG/VIAL</u>	<u>A203859</u>	<u>001</u>	Mar 31, 2017
<u>AP</u>	PAR STERILE PRODUCTS	<u>500MG/VIAL</u>	<u>A203575</u>	<u>001</u>	Oct 28, 2016
<u>AP</u>	RISING	<u>500MG/VIAL</u>	<u>A204043</u>	<u>001</u>	Feb 28, 2017
<u>AP</u>	STERISCIENCE	<u>500MG/VIAL</u>	<u>A216390</u>	<u>001</u>	Dec 23, 2022
<u>AP</u>	ZYDUS PHARMS	<u>500MG/VIAL</u>	<u>A204473</u>	<u>001</u>	Aug 31, 2017

MYCOPHENOLIC SODIUM

TABLET, DELAYED RELEASE; ORAL

MYCOPHENOLIC SODIUM

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 180MG BASE</u>	<u>A202555</u>	<u>001</u>	Aug 23, 2017
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A202555</u>	<u>002</u>	Aug 23, 2017
<u>AB</u>	AIRIS PHARMA PVT LTD	<u>EQ 180MG BASE</u>	<u>A217031</u>	<u>001</u>	Nov 29, 2023
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A217031</u>	<u>002</u>	Nov 29, 2023
<u>AB</u>	ALKEM LABS LTD	<u>EQ 180MG BASE</u>	<u>A208315</u>	<u>001</u>	Sep 23, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A208315</u>	<u>002</u>	Sep 23, 2021
<u>AB</u>	AMTA	<u>EQ 180MG BASE</u>	<u>A214376</u>	<u>001</u>	Feb 10, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A214376</u>	<u>002</u>	Feb 10, 2021
<u>AB</u>	APOTEX INC	<u>EQ 180MG BASE</u>	<u>A091558</u>	<u>001</u>	Aug 21, 2012
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A091558</u>	<u>002</u>	Aug 19, 2014
<u>AB</u>	BIOCON PHARMA	<u>EQ 180MG BASE</u>	<u>A214630</u>	<u>001</u>	Nov 29, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A214630</u>	<u>002</u>	Nov 29, 2021
<u>AB</u>	CONCORD BIOTECH LTD	<u>EQ 180MG BASE</u>	<u>A211173</u>	<u>001</u>	Dec 13, 2019
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A211173</u>	<u>002</u>	Dec 13, 2019
<u>AB</u>	RK PHARMA	<u>EQ 180MG BASE</u>	<u>A091248</u>	<u>002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A091248</u>	<u>001</u>	Jan 08, 2014
<u>AB</u>	TWI PHARMS	<u>EQ 180MG BASE</u>	<u>A214289</u>	<u>001</u>	Nov 03, 2021
<u>AB</u>		<u>EQ 360MG BASE</u>	<u>A214289</u>	<u>002</u>	Nov 03, 2021

MYFORTIC

<u>AB</u>	<u>+</u> NOVARTIS	<u>EQ 180MG BASE</u>	<u>N050791</u>	<u>001</u>	Feb 27, 2004
<u>AB</u>	<u>+!</u>	<u>EQ 360MG BASE</u>	<u>N050791</u>	<u>002</u>	Feb 27, 2004

NABILONE

CAPSULE; ORAL

CESAMET

<u>+!</u>	BAUSCH	1MG	N018677	001	Dec 26, 1985
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NABUMETONE

TABLET; ORAL

NABUMETONE

<u>AB</u>	ANNORA PHARMA	<u>500MG</u>	<u>A090445</u>	<u>001</u>	Jan 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A090445</u>	<u>002</u>	Jan 12, 2011
<u>AB</u>	CHARTWELL MOLECULES	<u>500MG</u>	<u>A076009</u>	<u>001</u>	Jan 24, 2003
<u>AB</u>		<u>750MG</u>	<u>A076009</u>	<u>002</u>	Jan 24, 2003
<u>AB</u>	CHARTWELL RX	<u>500MG</u>	<u>A075280</u>	<u>001</u>	Feb 25, 2002
<u>AB</u>		<u>750MG</u>	<u>A075280</u>	<u>002</u>	Feb 25, 2002
<u>AB</u>	INVAGEN PHARMS	<u>500MG</u>	<u>A078671</u>	<u>001</u>	Mar 07, 2008
<u>AB</u>	<u>!</u>	<u>750MG</u>	<u>A078671</u>	<u>002</u>	Mar 07, 2008
<u>AB</u>	LGM PHARMA	<u>500MG</u>	<u>A203166</u>	<u>001</u>	Aug 30, 2019
<u>AB</u>		<u>750MG</u>	<u>A203166</u>	<u>002</u>	Aug 30, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A078420</u>	<u>001</u>	Sep 24, 2008
<u>AB</u>		<u>750MG</u>	<u>A078420</u>	<u>002</u>	Sep 24, 2008
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>A091083</u>	<u>001</u>	Jun 13, 2011
<u>AB</u>		<u>750MG</u>	<u>A091083</u>	<u>002</u>	Jun 13, 2011
	LGM PHARMA	1GM	A203166	003	Aug 30, 2019

NADOLOL

TABLET; ORAL

CORGARD

<u>AB</u>	<u>+</u> USWM	<u>20MG</u>	<u>N018063</u>	<u>005</u>	Oct 28, 1986
<u>AB</u>	<u>+</u>	<u>40MG</u>	<u>N018063</u>	<u>001</u>	
<u>AB</u>	<u>+!</u>	<u>80MG</u>	<u>N018063</u>	<u>002</u>	

NADOLOL

<u>AB</u>	ALEMBIC	<u>20MG</u>	<u>A211763</u>	<u>001</u>	Jun 02, 2023
<u>AB</u>		<u>40MG</u>	<u>A211763</u>	<u>002</u>	Jun 02, 2023
<u>AB</u>		<u>80MG</u>	<u>A211763</u>	<u>003</u>	Jun 02, 2023
<u>AB</u>	AMNEAL PHARMS CO	<u>20MG</u>	<u>A208832</u>	<u>001</u>	Jun 02, 2017

PRESCRIPTION DRUG PRODUCT LIST

NADOLOL

TABLET; ORAL

NADOLOL

<u>AB</u>		<u>40MG</u>	<u>A208832</u>	<u>002</u>	Jun 02, 2017
<u>AB</u>		<u>80MG</u>	<u>A208832</u>	<u>003</u>	Jun 02, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>20MG</u>	<u>A201893</u>	<u>003</u>	Jun 07, 2022
<u>AB</u>		<u>40MG</u>	<u>A201893</u>	<u>001</u>	Sep 16, 2015
<u>AB</u>		<u>80MG</u>	<u>A201893</u>	<u>002</u>	Sep 16, 2015
<u>AB</u>	BEXIMCO PHARMS USA	<u>20MG</u>	<u>A210955</u>	<u>001</u>	Jul 23, 2018
<u>AB</u>		<u>40MG</u>	<u>A210955</u>	<u>002</u>	Jul 23, 2018
<u>AB</u>		<u>80MG</u>	<u>A210955</u>	<u>003</u>	Jul 23, 2018
<u>AB</u>	CHARTWELL RX	<u>20MG</u>	<u>A209309</u>	<u>001</u>	Oct 05, 2017
<u>AB</u>		<u>40MG</u>	<u>A209309</u>	<u>002</u>	Oct 05, 2017
<u>AB</u>		<u>80MG</u>	<u>A209309</u>	<u>003</u>	Oct 05, 2017
<u>AB</u>	INVAGEN PHARMS	<u>20MG</u>	<u>A203455</u>	<u>001</u>	Dec 18, 2015
<u>AB</u>		<u>40MG</u>	<u>A203455</u>	<u>002</u>	Dec 18, 2015
<u>AB</u>		<u>80MG</u>	<u>A203455</u>	<u>003</u>	Dec 18, 2015
<u>AB</u>	RISING	<u>20MG</u>	<u>A074172</u>	<u>001</u>	Oct 31, 1993
<u>AB</u>		<u>40MG</u>	<u>A074172</u>	<u>002</u>	Oct 31, 1993
<u>AB</u>		<u>80MG</u>	<u>A074172</u>	<u>003</u>	Oct 31, 1993
<u>AB</u>	RK PHARMA	<u>20MG</u>	<u>A212856</u>	<u>001</u>	Sep 13, 2019
<u>AB</u>		<u>40MG</u>	<u>A212856</u>	<u>002</u>	Sep 13, 2019
<u>AB</u>		<u>80MG</u>	<u>A212856</u>	<u>003</u>	Sep 13, 2019
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>A074501</u>	<u>001</u>	Nov 09, 1995
<u>AB</u>		<u>40MG</u>	<u>A074501</u>	<u>002</u>	Nov 09, 1995
<u>AB</u>		<u>80MG</u>	<u>A074501</u>	<u>003</u>	Nov 09, 1995
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A207761</u>	<u>001</u>	Jul 28, 2017
<u>AB</u>		<u>40MG</u>	<u>A207761</u>	<u>002</u>	Jul 28, 2017
<u>AB</u>		<u>80MG</u>	<u>A207761</u>	<u>003</u>	Jul 28, 2017

NAFARELIN ACETATE

SPRAY, METERED; NASAL

SYNAREL

+! PFIZER

EQ 0.2MG BASE/SPRAY

N019886 001 Feb 13, 1990

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

<u>AP</u>	ANTIBIOTICE	<u>EQ 1GM BASE/VIAL</u>	<u>A090560</u>	<u>001</u>	Oct 03, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090560</u>	<u>002</u>	Oct 03, 2011
<u>AP</u>	! EUGIA PHARMA SPECLTS	<u>EQ 1GM BASE/VIAL</u>	<u>A091613</u>	<u>001</u>	Dec 26, 2012
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A091613</u>	<u>002</u>	Dec 26, 2012
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A091614</u>	<u>001</u>	Dec 26, 2012
<u>AP</u>	FRESENIUS	<u>EQ 10GM BASE/VIAL</u>	<u>A206761</u>	<u>001</u>	Jun 02, 2020
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 1GM BASE/VIAL</u>	<u>A090002</u>	<u>001</u>	Jun 30, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090002</u>	<u>002</u>	Jun 30, 2011
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090005</u>	<u>001</u>	Apr 20, 2011
<u>AP</u>	SAGENT PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A090582</u>	<u>001</u>	Aug 24, 2012
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090582</u>	<u>002</u>	Aug 24, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090580</u>	<u>001</u>	Aug 24, 2012
<u>AP</u>	STERISCIENCE	<u>EQ 1GM BASE/VIAL</u>	<u>A200002</u>	<u>001</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A200002</u>	<u>002</u>	Apr 07, 2014
	NALLPEN IN PLASTIC CONTAINER				
	+! BAXTER HLTHCARE	EQ 20MG BASE/ML	N050655	001	Oct 31, 1989
	+!	EQ 2GM BASE/100ML	N050655	002	Oct 31, 1989

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIFINE HYDROCHLORIDE

<u>AB</u>	AMNEAL	<u>2%</u>	<u>A206960</u>	<u>001</u>	Apr 10, 2017
<u>AB</u>	TARO	<u>2%</u>	<u>A206901</u>	<u>001</u>	Jan 06, 2016
<u>AB</u>	XIROMED	<u>2%</u>	<u>A210038</u>	<u>001</u>	Sep 22, 2020
	<u>NAFTIN</u>				
<u>AB</u>	+! SEBELA IRELAND LTD	<u>2%</u>	<u>N019599</u>	<u>002</u>	Jan 13, 2012
	NAFTIFINE HYDROCHLORIDE				
	! TARO	1%	A205975	001	Sep 08, 2016
	GEL; TOPICAL				
	<u>NAFTIFINE HYDROCHLORIDE</u>				
<u>AB</u>	AMNEAL	<u>1%</u>	<u>A206165</u>	<u>001</u>	Mar 20, 2019
<u>AB</u>	TARO	<u>2%</u>	<u>A208201</u>	<u>001</u>	Apr 10, 2019
	<u>NAFTIN</u>				
<u>AB</u>	+! SEBELA IRELAND LTD	<u>1%</u>	<u>N019356</u>	<u>001</u>	Jun 18, 1990

PRESCRIPTION DRUG PRODUCT LIST

NAFTIFINE HYDROCHLORIDE

GEL; TOPICAL

NAFTIN

AB	+ !	2%	<u>N204286</u>	<u>001</u>	Jun 27, 2013
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NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

AP	!	HOSPIRA	<u>10MG/ML</u>	<u>A070914</u>	<u>001</u>	Feb 03, 1989
AP	!		<u>10MG/ML</u>	<u>A070915</u>	<u>001</u>	Feb 03, 1989
AP	!		<u>20MG/ML</u>	<u>A070916</u>	<u>001</u>	Feb 03, 1989
AP	!		<u>20MG/ML</u>	<u>A070918</u>	<u>001</u>	Feb 03, 1989

NALDEMEDINE TOSYLATE

TABLET; ORAL

SYMPROIC

+! BDSI

EQ 0.2MG BASE

N208854 001 Mar 23, 2017

NALMEFENE HYDROCHLORIDE

SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

NALMEFENE HYDROCHLORIDE

AP		CHENGDU SHUODE	<u>EQ 2MG BASE/2ML (EQ 1MG BASE/ML)</u>	<u>A216007</u>	<u>002</u>	Nov 15, 2023
AP	!	PURDUE PHARMA LP	<u>EQ 2MG BASE/2ML (EQ 1MG BASE/ML)</u>	<u>A212955</u>	<u>001</u>	Feb 08, 2022
		CHENGDU SHUODE	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	A216007	001	Nov 15, 2023
		SPRAY; NASAL				
		OPVEE				
	+ !	INDIVIOR	EQ 2.7MG BASE/SPRAY	N217470	001	May 22, 2023

NALOXEGOL OXALATE

TABLET; ORAL

MOVANTIK

+ VALINOR

EQ 12.5MG BASE

N204760 001 Sep 16, 2014

+!

EQ 25MG BASE

N204760 002 Sep 16, 2014

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

AP		HIKMA	<u>0.4MG/ML</u>	<u>A070299</u>	<u>001</u>	Sep 24, 1986
		<u>NALOXONE HYDROCHLORIDE</u>				
AP		ACCORD HLTHCARE	<u>0.4MG/ML</u>	<u>A216624</u>	<u>001</u>	Oct 26, 2022
AP			<u>1MG/ML</u>	<u>A216624</u>	<u>002</u>	Oct 26, 2022
AP		BAXTER HLTHCARE CORP	<u>0.4MG/ML</u>	<u>A214785</u>	<u>001</u>	Jan 29, 2021
AP			<u>0.4MG/ML</u>	<u>A214792</u>	<u>001</u>	Nov 07, 2022
AP		BPI LABS	<u>1MG/ML</u>	<u>A216977</u>	<u>001</u>	Oct 06, 2023
AP		DR REDDYS	<u>1MG/ML</u>	<u>A213209</u>	<u>001</u>	Mar 16, 2020
AP		EUGIA PHARMA	<u>0.4MG/ML</u>	<u>A212455</u>	<u>001</u>	Oct 15, 2019
AP			<u>1MG/ML</u>	<u>A213279</u>	<u>001</u>	Jan 14, 2021
AP			<u>0.4MG/ML</u>	<u>A212456</u>	<u>001</u>	Nov 04, 2019
AP		HIKMA	<u>1MG/ML</u>	<u>A212300</u>	<u>001</u>	Jun 10, 2022
AP	!	HOSPIRA	<u>0.4MG/ML</u>	<u>A070172</u>	<u>001</u>	Sep 24, 1986
AP	!		<u>0.4MG/ML</u>	<u>A070256</u>	<u>001</u>	Jan 07, 1987
AP	!		<u>0.4MG/ML</u>	<u>A070257</u>	<u>001</u>	Jan 07, 1987
AP	!	INTL MEDICATION	<u>1MG/ML</u>	<u>A072076</u>	<u>001</u>	Mar 24, 1988
AP		MYLAN INSTITUTIONAL	<u>0.4MG/ML</u>	<u>A204997</u>	<u>001</u>	Mar 06, 2014
AP			<u>0.4MG/ML</u>	<u>A205014</u>	<u>001</u>	Jun 29, 2016
AP		MYLAN LABS LTD	<u>1MG/ML</u>	<u>A213843</u>	<u>001</u>	Jun 09, 2022
AP		RISING	<u>0.4MG/ML</u>	<u>A208871</u>	<u>001</u>	Feb 28, 2017
AP			<u>0.4MG/ML</u>	<u>A208872</u>	<u>001</u>	Mar 14, 2017
AP		SOMERSET THERAPS LLC	<u>0.4MG/ML</u>	<u>A207633</u>	<u>001</u>	Aug 08, 2017
AP			<u>0.4MG/ML</u>	<u>A207634</u>	<u>001</u>	Jul 26, 2017
		SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS				
		ZIMHI				
	+ !	ADAMIS PHARMS CORP	5MG/0.5ML (5MG/0.5ML)	N212854	001	Oct 15, 2021
		SPRAY; NASAL				
		KLOXXADO				
	+ !	HIKMA	8MG/SPRAY	N212045	001	Apr 29, 2021
		SPRAY, METERED; NASAL				
		NALOXONE HYDROCHLORIDE				
		TEVA PHARMS USA	4MG/SPRAY	A209522	001	Apr 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE

AB	LUPIN	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	A075735 001	Jul 11, 2001
AB	SUN PHARM INDS LTD	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	A075523 001	Mar 17, 2000
AB	! WATSON LABS	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	A074736 001	Jan 21, 1997

NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

NALTREXONE

AP	TEVA PHARMS USA INC	380MG/VIAL	A213195 001	Jul 06, 2023
AP	+! ALKERMES	380MG/VIAL	N021897 001	Apr 13, 2006

VIVITROLNALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

AB	ACCORD HLTHCARE	50MG	A091205 001	Aug 17, 2011
AB	BARR	50MG	A074918 001	May 08, 1998
AB	CHARTWELL	50MG	A207905 001	Jul 21, 2017
AB	ELITE LABS	50MG	A075274 001	May 26, 1999
AB	! SPECGX LLC	50MG	A076264 002	Mar 22, 2002
AB	SUN PHARM	50MG	A090356 001	Feb 24, 2012
	SPECGX LLC	25MG	A076264 001	Mar 22, 2002
		100MG	A076264 003	Mar 22, 2002

NAPROXEN

SUSPENSION; ORAL

NAPROSYN

AB	+ ATNAHS PHARMA US	25MG/ML	N018965 001	Mar 23, 1987
AB	AMNEAL	25MG/ML	A212705 001	Jul 31, 2020
AB	HETERO LABS LTD III	25MG/ML	A215776 001	Jun 07, 2022
AB	! HIKMA	25MG/ML	A074190 001	Mar 30, 1994
AB	NOVITIUM PHARMA	25MG/ML	A211910 001	Mar 10, 2021

NAPROXEN

TABLET; ORAL

NAPROSYN

AB	+! ATNAHS PHARMA US	500MG	N017581 004	Apr 15, 1982
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NAPROXEN

AB	AMNEAL PHARMS NY	250MG	A075927 001	Dec 18, 2001
AB		375MG	A075927 002	Dec 18, 2001
AB		500MG	A075927 003	Dec 18, 2001
AB	AUROBINDO PHARMA	250MG	A200429 001	Nov 08, 2011
AB		375MG	A200429 002	Nov 08, 2011
AB		500MG	A200429 003	Nov 08, 2011
AB	GLENMARK GENERICS	250MG	A078250 001	Mar 28, 2007
AB		375MG	A078250 002	Mar 28, 2007
AB		500MG	A078250 003	Mar 28, 2007
AB	GRANULES	250MG	A074140 001	Dec 21, 1993
AB		375MG	A074140 002	Dec 21, 1993
AB		500MG	A074140 003	Dec 21, 1993
AB	INVAGEN PHARMS	250MG	A091305 001	Aug 24, 2011
AB		375MG	A091305 002	Aug 24, 2011
AB		500MG	A091305 003	Aug 24, 2011
AB	MARKSANS PHARMA	250MG	A091416 001	Feb 14, 2011
AB		375MG	A091416 002	Feb 14, 2011
AB		500MG	A091416 003	Feb 14, 2011
AB	SCIEGEN PHARMS INC	250MG	A212517 001	Feb 21, 2020
AB		375MG	A212517 002	Feb 21, 2020
AB		500MG	A212517 003	Feb 21, 2020
AB	TEVA	250MG	A074201 001	Dec 21, 1993
AB		375MG	A074201 002	Dec 21, 1993
AB		500MG	A074201 003	Dec 21, 1993
AB	ZYDUS PHARMS USA	250MG	A078620 001	Jun 07, 2007
AB		375MG	A078620 002	Jun 07, 2007
AB		500MG	A078620 003	Jun 07, 2007

TABLET, DELAYED RELEASE; ORAL

EC-NAPROSYN

AB	+! ATNAHS PHARMA US	375MG	N020067 002	Oct 14, 1994
AB	+!	500MG	N020067 003	Oct 14, 1994

NAPROXEN

AB	NUVO PHARMS INC	375MG	A091432 001	Sep 19, 2011
AB		500MG	A091432 002	Sep 19, 2011
AB	TEVA	375MG	A075227 001	Jun 30, 1998

PRESCRIPTION DRUG PRODUCT LIST

NAPROXEN

TABLET, DELAYED RELEASE;ORAL

NAPROXEN

<u>AB</u>		<u>500MG</u>	<u>A075227 002</u>	Jun 30, 1998
<u>AB</u>	TULEX PHARMS INC	<u>375MG</u>	<u>A216908 001</u>	May 31, 2023
<u>AB</u>		<u>500MG</u>	<u>A216908 002</u>	May 31, 2023

NAPROXEN SODIUM

TABLET;ORAL

ANAPROX DS

<u>AB</u>	<u>+</u> !	ATNAHS PHARMA US	<u>EQ 500MG BASE</u>	<u>N018164 003</u>	Sep 30, 1987
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NAPROXEN SODIUM

<u>AB</u>		AMNEAL PHARMS NY	<u>EQ 250MG BASE</u>	<u>A078432 001</u>	Apr 25, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078432 002</u>	Apr 25, 2007
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A200629 001</u>	Oct 31, 2011
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A200629 002</u>	Oct 31, 2011
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 250MG BASE</u>	<u>A078486 001</u>	Jul 26, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078486 002</u>	Jul 26, 2007
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 250MG BASE</u>	<u>A078314 001</u>	Apr 27, 2007
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A078314 002</u>	Apr 27, 2007
<u>AB</u>		SCIEGEN PHARMS INC	<u>EQ 250MG BASE</u>	<u>A212199 001</u>	Oct 30, 2019
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A212199 002</u>	Oct 30, 2019

TABLET, EXTENDED RELEASE;ORAL

NAPRELAN

<u>AB</u>	<u>+</u>	TWI PHARMS	<u>EQ 375MG BASE</u>	<u>N020353 001</u>	Jan 05, 1996
<u>AB</u>	<u>+</u>		<u>EQ 500MG BASE</u>	<u>N020353 002</u>	Jan 05, 1996
<u>AB</u>	<u>+</u> !		<u>EQ 750MG BASE</u>	<u>N020353 003</u>	Jan 05, 1996

NAPROXEN SODIUM

<u>AB</u>		ACTAVIS LABS FL INC	<u>EQ 375MG BASE</u>	<u>A075416 002</u>	Apr 23, 2003
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>A075416 001</u>	Aug 27, 2002
<u>AB</u>			<u>EQ 750MG BASE</u>	<u>A075416 003</u>	Aug 11, 2016

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET;ORAL

SUMATRIPTAN AND NAPROXEN SODIUM

<u>AB</u>		AUROBINDO PHARMA LTD	<u>500MG;EQ 85MG BASE</u>	<u>A207457 001</u>	Feb 15, 2018
<u>AB</u>		RISING	<u>500MG;EQ 85MG BASE</u>	<u>A090872 001</u>	Sep 04, 2018
<u>AB</u>		SUN PHARM	<u>500MG;EQ 85MG BASE</u>	<u>A202803 001</u>	Jul 20, 2018

TREXIMET

<u>AB</u>	<u>+</u> !	CURRAX	<u>500MG;EQ 85MG BASE</u>	<u>N021926 001</u>	Apr 15, 2008
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NARATRIPTAN HYDROCHLORIDE

TABLET;ORAL

NARATRIPTAN

<u>AB</u>		HERITAGE PHARMS	<u>EQ 1MG BASE</u>	<u>A200502 001</u>	Feb 28, 2011
<u>AB</u>	<u>!</u>		<u>EQ 2.5MG BASE</u>	<u>A200502 002</u>	Feb 28, 2011
<u>AB</u>		HIKMA	<u>EQ 1MG BASE</u>	<u>A090381 001</u>	Jul 07, 2010
<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A090381 002</u>	Jul 07, 2010
<u>AB</u>		ORBION PHARMS	<u>EQ 1MG BASE</u>	<u>A091441 001</u>	Apr 30, 2012
<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A091441 002</u>	Apr 30, 2012
<u>AB</u>		PADAGIS US	<u>EQ 1MG BASE</u>	<u>A091326 001</u>	Jul 08, 2010
<u>AB</u>			<u>EQ 2.5MG BASE</u>	<u>A091326 002</u>	Jul 08, 2010
<u>AB</u>		SUN PHARM INDS LTD	<u>EQ 2.5MG BASE</u>	<u>A091552 001</u>	Feb 14, 2011

NATAMYCIN

SUSPENSION;OPHTHALMIC

NATACYN

<u>+</u> !	HARROW EYE	5%	<u>N050514 001</u>
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NATEGLINIDE

TABLET;ORAL

NATEGLINIDE

<u>AB</u>		CADILA PHARMS LTD	<u>60MG</u>	<u>A206432 001</u>	Apr 19, 2019
<u>AB</u>			<u>120MG</u>	<u>A206432 002</u>	Apr 19, 2019
<u>AB</u>		DR REDDYS LABS LTD	<u>60MG</u>	<u>A077461 001</u>	Sep 09, 2009
<u>AB</u>			<u>120MG</u>	<u>A077461 002</u>	Sep 09, 2009
<u>AB</u>		RISING	<u>60MG</u>	<u>A205544 001</u>	Jun 18, 2018
<u>AB</u>			<u>120MG</u>	<u>A205544 002</u>	Jun 18, 2018
<u>AB</u>		STRIDES PHARMA	<u>60MG</u>	<u>A077463 001</u>	Sep 09, 2009
<u>AB</u>			<u>120MG</u>	<u>A077463 002</u>	Sep 09, 2009
<u>AB</u>		WATSON LABS	<u>60MG</u>	<u>A077462 001</u>	Mar 30, 2011
<u>AB</u>			<u>120MG</u>	<u>A077462 002</u>	Mar 30, 2011
<u>AB</u>		ZYDUS PHARMS	<u>60MG</u>	<u>A205248 001</u>	Jul 06, 2016

PRESCRIPTION DRUG PRODUCT LIST

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

<u>AB</u>	!	<u>120MG</u>	<u>A205248</u>	<u>002</u>	Jul 06, 2016
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NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

BYSTOLIC

<u>AB</u>	+	ALLERGAN	<u>EQ 2.5MG BASE</u>	<u>N021742</u>	<u>002</u>	Dec 17, 2007
<u>AB</u>	+		<u>EQ 5MG BASE</u>	<u>N021742</u>	<u>003</u>	Dec 17, 2007
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N021742</u>	<u>004</u>	Dec 17, 2007
<u>AB</u>	+	!	<u>EQ 20MG BASE</u>	<u>N021742</u>	<u>005</u>	Oct 08, 2008

NEBIVOLOL HYDROCHLORIDE

<u>AB</u>		ANI PHARMS	<u>EQ 2.5MG BASE</u>	<u>A203659</u>	<u>001</u>	Apr 16, 2015
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203659</u>	<u>002</u>	Apr 16, 2015
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203659</u>	<u>003</u>	Apr 16, 2015
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203659</u>	<u>004</u>	Apr 16, 2015
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 2.5MG BASE</u>	<u>A211053</u>	<u>001</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A211053</u>	<u>002</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A211053</u>	<u>003</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A211053</u>	<u>004</u>	Dec 17, 2021
<u>AB</u>		BEXIMCO PHARMS USA	<u>EQ 2.5MG BASE</u>	<u>A216568</u>	<u>001</u>	Mar 30, 2023
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A216568</u>	<u>002</u>	Mar 30, 2023
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A216568</u>	<u>003</u>	Mar 30, 2023
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A216568</u>	<u>004</u>	Mar 30, 2023
<u>AB</u>		CADILA PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A208717</u>	<u>001</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A208717</u>	<u>002</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A208717</u>	<u>003</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A208717</u>	<u>004</u>	Dec 17, 2021
<u>AB</u>		HETERO LABS LTD III	<u>EQ 2.5MG BASE</u>	<u>A203825</u>	<u>001</u>	Nov 03, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203825</u>	<u>002</u>	Nov 03, 2020
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203825</u>	<u>003</u>	Nov 03, 2020
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203825</u>	<u>004</u>	Nov 03, 2020
<u>AB</u>		INDCHEMIE HEALTH	<u>EQ 2.5MG BASE</u>	<u>A203828</u>	<u>001</u>	Jul 29, 2015
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203828</u>	<u>002</u>	Jul 29, 2015
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203828</u>	<u>003</u>	Jul 29, 2015
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203828</u>	<u>004</u>	Jul 29, 2015
<u>AB</u>		MANKIND PHARMA	<u>EQ 2.5MG BASE</u>	<u>A216172</u>	<u>001</u>	Nov 14, 2022
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A216172</u>	<u>002</u>	Nov 14, 2022
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A216172</u>	<u>003</u>	Nov 14, 2022
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A216172</u>	<u>004</u>	Nov 14, 2022
<u>AB</u>		PRINSTON INC	<u>EQ 2.5MG BASE</u>	<u>A212682</u>	<u>001</u>	Apr 07, 2022
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A212682</u>	<u>002</u>	Apr 07, 2022
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A212682</u>	<u>003</u>	Apr 07, 2022
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A212682</u>	<u>004</u>	Apr 07, 2022
<u>AB</u>		REYOUNG	<u>EQ 2.5MG BASE</u>	<u>A212917</u>	<u>001</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A212917</u>	<u>002</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A212917</u>	<u>003</u>	Dec 17, 2021
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A212917</u>	<u>004</u>	Dec 17, 2021
<u>AB</u>		TORRENT	<u>EQ 2.5MG BASE</u>	<u>A203966</u>	<u>001</u>	Mar 02, 2018
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203966</u>	<u>002</u>	Mar 02, 2018
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A203966</u>	<u>003</u>	Mar 02, 2018
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A203966</u>	<u>004</u>	Mar 02, 2018

NEBIVOLOL HYDROCHLORIDE; VALSARTAN

TABLET; ORAL

VYDUO

		PRINSTON INC	<u>EQ 5MG BASE; 80MG</u>	<u>A210596</u>	<u>001</u>	Sep 19, 2022
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NEDOSIRAN SODIUM

SOLUTION; INJECTION

RIVFLOZA

	+	NOVO	<u>EQ 80MG BASE/0.5ML (EQ 160MG BASE/ML)</u>	<u>N215842</u>	<u>001</u>	Sep 29, 2023
	+	!	<u>EQ 128MG BASE/0.8ML (EQ 160MG BASE/ML)</u>	<u>N215842</u>	<u>002</u>	Sep 29, 2023
	+	!	<u>EQ 160MG BASE/ML (EQ 160MG BASE/ML)</u>	<u>N215842</u>	<u>003</u>	Sep 29, 2023

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

TEVA

			<u>50MG</u>	<u>A076037</u>	<u>001</u>	Sep 16, 2003
			<u>100MG</u>	<u>A076037</u>	<u>002</u>	Sep 16, 2003
			<u>150MG</u>	<u>A076037</u>	<u>003</u>	Sep 16, 2003
			<u>200MG</u>	<u>A076037</u>	<u>004</u>	Sep 16, 2003
	!		<u>250MG</u>	<u>A076037</u>	<u>005</u>	Sep 16, 2003

PRESCRIPTION DRUG PRODUCT LIST

NELARABINE

INJECTABLE; INTRAVENOUS

ARRANON

<u>AP</u>	<u>+!</u>	<u>SANDOZ</u>	<u>250MG/50ML (5MG/ML)</u>	<u>N021877</u>	<u>001</u>	Oct 28, 2005
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NELARABINE

<u>AP</u>		<u>AMNEAL</u>	<u>250MG/50ML (5MG/ML)</u>	<u>A216346</u>	<u>001</u>	Apr 04, 2023
<u>AP</u>		<u>DR REDDYS</u>	<u>250MG/50ML (5MG/ML)</u>	<u>A216934</u>	<u>001</u>	Dec 23, 2022
<u>AP</u>		<u>GLAND PHARMA LTD</u>	<u>250MG/50ML (5MG/ML)</u>	<u>A212605</u>	<u>001</u>	Jan 03, 2024
<u>AP</u>		<u>KINDOS</u>	<u>250MG/50ML (5MG/ML)</u>	<u>A216038</u>	<u>001</u>	Jan 10, 2023
<u>AP</u>		<u>NEXUS</u>	<u>250MG/50ML (5MG/ML)</u>	<u>A215057</u>	<u>001</u>	Jun 02, 2023
<u>AP</u>		<u>SHORLA</u>	<u>250MG/50ML (5MG/ML)</u>	<u>A214809</u>	<u>001</u>	Mar 03, 2023
<u>AP</u>		<u>ZYDUS PHARMS</u>	<u>250MG/50ML (5MG/ML)</u>	<u>A215037</u>	<u>001</u>	Nov 17, 2021

NELFINAVIR MESYLATE

TABLET; ORAL

VIRACEPT

<u>+!</u>	<u>AGOURON PHARMS</u>	<u>EQ 250MG BASE</u>	<u>N020779</u>	<u>001</u>	Mar 14, 1997
<u>+!</u>		<u>EQ 625MG BASE</u>	<u>N021503</u>	<u>001</u>	Apr 30, 2003

NEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

<u>AA</u>	<u>!</u>	<u>TEVA</u>	<u>500MG</u>	<u>A060304</u>	<u>001</u>	
<u>AA</u>		<u>XGEN PHARMS</u>	<u>500MG</u>	<u>A065220</u>	<u>001</u>	Jul 28, 2006

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

<u>AT</u>		<u>WATSON LABS</u>	<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A062664</u>	<u>001</u>	Apr 08, 1986
<u>AT</u>		<u>XGEN PHARMS</u>	<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A065106</u>	<u>001</u>	Jan 31, 2006
<u>AT</u>			<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A065108</u>	<u>001</u>	Jan 31, 2006

NEOSPORIN G.U. IRRIGANT

<u>AT</u>	<u>!</u>	<u>MONARCH PHARMS</u>	<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A060707</u>	<u>001</u>	
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NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

BLOXIVERZ

<u>AP</u>	<u>+!</u>	<u>EXELA PHARMA</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>N204078</u>	<u>001</u>	May 31, 2013
<u>AP</u>	<u>+!</u>		<u>10MG/10ML (1MG/ML)</u>	<u>N204078</u>	<u>002</u>	May 31, 2013

NEOSTIGMINE METHYLSULFATE

<u>AP</u>		<u>AMNEAL</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210051</u>	<u>001</u>	Jun 15, 2018
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A210051</u>	<u>002</u>	Jun 15, 2018
<u>AP</u>		<u>AMPHASTAR PHARMS INC</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A209933</u>	<u>001</u>	Sep 25, 2017
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A209933</u>	<u>002</u>	Sep 25, 2017
<u>AP</u>		<u>AMRING PHARMS</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210989</u>	<u>001</u>	Aug 22, 2018
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A210989</u>	<u>002</u>	Aug 22, 2018
<u>AP</u>		<u>BE PHARMS</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212512</u>	<u>001</u>	May 13, 2019
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A212512</u>	<u>002</u>	May 13, 2019
<u>AP</u>		<u>CAPLIN</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A213074</u>	<u>001</u>	Apr 20, 2021
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A213074</u>	<u>002</u>	Apr 20, 2021
<u>AP</u>		<u>DR REDDYS</u>	<u>3MG/3ML (1MG/ML)</u>	<u>A216291</u>	<u>001</u>	Jul 06, 2022
<u>AP</u>			<u>5MG/10ML (0.5MG/ML)</u>	<u>A209135</u>	<u>001</u>	Jul 10, 2018
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A209135</u>	<u>002</u>	Jul 10, 2018
<u>AP</u>		<u>EUGIA PHARMA</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A213244</u>	<u>001</u>	Nov 02, 2023
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A213244</u>	<u>002</u>	Nov 02, 2023
<u>AP</u>	<u>+!</u>	<u>FRESENIUS KABI USA</u>	<u>3MG/3ML (1MG/ML)</u>	<u>N203629</u>	<u>003</u>	Sep 18, 2018
<u>AP</u>		<u>GLAND PHARMA LTD</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212968</u>	<u>001</u>	Oct 16, 2019
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A212968</u>	<u>002</u>	Oct 16, 2019
<u>AP</u>		<u>HIKMA</u>	<u>3MG/3ML (1MG/ML)</u>	<u>A216206</u>	<u>001</u>	Jul 13, 2022
<u>AP</u>			<u>5MG/10ML (0.5MG/ML)</u>	<u>A207042</u>	<u>001</u>	Dec 28, 2015
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A207042</u>	<u>002</u>	Dec 28, 2015
<u>AP</u>		<u>INDOCO</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A210652</u>	<u>001</u>	Oct 01, 2021
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A210652</u>	<u>002</u>	Oct 01, 2021
<u>AP</u>		<u>MEITHEAL</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212804</u>	<u>001</u>	Apr 05, 2021
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A212804</u>	<u>002</u>	Apr 05, 2021
<u>AP</u>		<u>PAR STERILE PRODUCTS</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A208405</u>	<u>001</u>	Apr 26, 2017
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A208405</u>	<u>002</u>	Apr 26, 2017
<u>AP</u>		<u>SAGENT PHARMS INC</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A216542</u>	<u>001</u>	Feb 17, 2023
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A216542</u>	<u>002</u>	Feb 17, 2023
<u>AP</u>		<u>UMEDICA</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A212627</u>	<u>001</u>	Nov 03, 2022
<u>AP</u>			<u>10MG/10ML (1MG/ML)</u>	<u>A212627</u>	<u>002</u>	Nov 03, 2022

PRESCRIPTION DRUG PRODUCT LIST

NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

NEOSTIGMINE METHYLSULFATE

AP	AVET LIFESCIENCES	5MG/10ML (0.5MG/ML)	A208230 001	Nov 25, 2022
AP		10MG/10ML (1MG/ML)	A208230 002	Nov 25, 2022
	BLOXIVERZ			
	+! EXELA PHARMA	5MG/5ML (1MG/ML)	N204078 003	Oct 27, 2023
	NEOSTIGMINE METHYLSULFATE			
	+! FRESENIUS KABI USA	5MG/10ML (0.5MG/ML)	N203629 001	Jan 08, 2015
	+!	10MG/10ML (1MG/ML)	N203629 002	Jan 08, 2015

NEPAFENAC

SUSPENSION/DROPS; OPHTHALMIC

ILEVRO

+! HARROW EYE 0.3% N203491 001 Oct 16, 2012

NEVANAC

+! HARROW EYE 0.1% N021862 001 Aug 19, 2005

NERATINIB MALEATE

TABLET; ORAL

NERLYNX

+! PUMA BIOTECH EQ 40MG BASE N208051 001 Jul 17, 2017

NETARSUDIL MESYLATE

SOLUTION/DROPS; OPHTHALMIC

RHOPRESSA

+! ALCON LABS INC EQ 0.02% BASE N208254 001 Dec 18, 2017

NETUPITANT; PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

AKYNZEO

+! HELSINN HLTHCARE 300MG; EQ 0.5MG BASE N205718 001 Oct 10, 2014

NEVIRAPINE

SUSPENSION; ORAL

NEVIRAPINE

AA	AUROBINDO	50MG/5ML	A077702 001	May 22, 2012
AA	CIPLA	50MG/5ML	A207684 001	Aug 03, 2017

VIRAMUNE

AA	+! BOEHRINGER INGELHEIM	50MG/5ML	N020933 001	Sep 11, 1998
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TABLET; ORAL

NEVIRAPINE

AB	AUROBINDO	200MG	A077521 001	May 22, 2012
AB	CIPLA	200MG	A077956 001	May 22, 2012
AB	HETERO LABS LTD III	200MG	A078584 001	May 22, 2012
AB	! MACLEODS PHARMS LTD	200MG	A090688 001	Jan 14, 2019
AB	MICRO LABS LTD	200MG	A203080 001	May 22, 2012
AB	MYLAN PHARMS INC	200MG	A202523 001	May 22, 2012
AB	STRIDES PHARMA	200MG	A078195 001	May 22, 2012

TABLET, EXTENDED RELEASE; ORAL

NEVIRAPINE

AB	MACLEODS PHARMS LTD	400MG	A206879 001	Oct 06, 2017
AB	MYLAN	400MG	A205651 001	Oct 27, 2014
AB	SANDOZ	400MG	A203411 001	Apr 03, 2014

VIRAMUNE XR

AB	+! BOEHRINGER INGELHEIM	400MG	N201152 001	Mar 25, 2011
	+	100MG	N201152 002	Nov 08, 2012

NIACIN

TABLET; ORAL

NIACOR

! AVONDALE PHARMS 500MG A040378 001 May 03, 2000

TABLET, EXTENDED RELEASE; ORAL

NIACIN

AB	AMNEAL PHARMS	500MG	A203578 001	Jul 24, 2015
AB		750MG	A204178 001	Dec 11, 2015
AB		1GM	A203578 002	Jul 24, 2015
AB	AUROBINDO PHARMA LTD	500MG	A209236 001	Feb 01, 2018
AB		750MG	A209236 002	Feb 01, 2018
AB		1GM	A209236 003	Feb 01, 2018
AB	BARR	500MG	A076378 001	Apr 26, 2005
AB		750MG	A076378 002	Apr 26, 2005
AB		1GM	A076250 001	Apr 14, 2005

PRESCRIPTION DRUG PRODUCT LIST

NIACIN

TABLET, EXTENDED RELEASE;ORAL

NIACIN

<u>AB</u>	EYWA	<u>500MG</u>	<u>A213090 001</u>	Aug 25, 2023
<u>AB</u>		<u>750MG</u>	<u>A213090 002</u>	Aug 25, 2023
<u>AB</u>		<u>1GM</u>	<u>A213090 003</u>	Aug 25, 2023
<u>AB</u>	LANNETT CO INC	<u>500MG</u>	<u>A203899 001</u>	Jun 16, 2017
<u>AB</u>		<u>1GM</u>	<u>A203899 002</u>	Jun 16, 2017
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A090860 001</u>	Mar 20, 2014
<u>AB</u>		<u>750MG</u>	<u>A090892 001</u>	Mar 20, 2014
<u>AB</u>		<u>1GM</u>	<u>A090446 001</u>	Mar 20, 2014
<u>AB</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A204934 001</u>	Mar 03, 2022
<u>AB</u>		<u>1GM</u>	<u>A204934 002</u>	Mar 03, 2022
<u>AB</u>	SUN PHARM	<u>500MG</u>	<u>A200484 001</u>	Apr 23, 2014
<u>AB</u>	!	<u>750MG</u>	<u>A201273 001</u>	Apr 23, 2014
<u>AB</u>	!	<u>1GM</u>	<u>A200484 002</u>	Apr 23, 2014

NICARDIPINE HYDROCHLORIDE

CAPSULE;ORAL

NICARDIPINE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>20MG</u>	<u>A074670 001</u>	Oct 28, 1996
<u>AB</u>		<u>30MG</u>	<u>A074670 002</u>	Oct 28, 1996
<u>AB</u>	BIONPHARMA	<u>20MG</u>	<u>A217555 001</u>	May 03, 2023
<u>AB</u>		<u>30MG</u>	<u>A217555 002</u>	May 03, 2023
<u>AB</u>	EPIC PHARMA LLC	<u>20MG</u>	<u>A074928 001</u>	Mar 19, 1998
<u>AB</u>	!	<u>30MG</u>	<u>A074928 002</u>	Mar 19, 1998
<u>AB</u>	GLENMARK PHARMS LTD	<u>20MG</u>	<u>A216357 001</u>	Dec 16, 2022
<u>AB</u>		<u>30MG</u>	<u>A216357 002</u>	Dec 16, 2022
<u>AB</u>	SENORES PHARMS	<u>20MG</u>	<u>A215377 001</u>	Jul 17, 2023
<u>AB</u>		<u>30MG</u>	<u>A215377 002</u>	Jul 17, 2023

INJECTABLE;INJECTION

NICARDIPINE HYDROCHLORIDE

<u>AP</u>	!	AM REGENT	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090534 001</u>	Nov 17, 2009
<u>AP</u>		EUGIA PHARMA	<u>25MG/10ML (2.5MG/ML)</u>	<u>A211121 001</u>	Apr 08, 2021
<u>AP</u>	+	HIKMA INTL PHARMS	<u>25MG/10ML (2.5MG/ML)</u>	<u>N022276 001</u>	Jul 24, 2008
<u>AP</u>		MICRO LABS	<u>25MG/10ML (2.5MG/ML)</u>	<u>A216420 001</u>	Jul 07, 2023

INJECTABLE;INTRAVENOUS

CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER

+! CHIESI 40MG/200ML (0.2MG/ML) N019734 004 Nov 07, 2008

CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER

+! CHIESI 20MG/200ML (0.1MG/ML) N019734 003 Jul 31, 2008

CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER

+! CHIESI 20MG/200ML (0.1MG/ML) N019734 002 Jul 31, 2008

NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE

+! HIKMA INTL PHARMS 20MG/200ML (0.1MG/ML) N022276 002 Apr 07, 2016

+! 40MG/200ML (0.2MG/ML) N022276 003 Apr 07, 2016

NICOTINE

INHALANT;ORAL

NICOTROL

+! PFIZER 4MG/CARTRIDGE N020714 001 May 02, 1997

SPRAY, METERED;NASAL

NICOTROL

+! PFIZER INC 0.5MG/SPRAY N020385 001 Mar 22, 1996

NIFEDIPINE

CAPSULE;ORAL

NIFEDIPINE

<u>AB</u>	ACELLA	<u>10MG</u>	<u>A072781 001</u>	Jul 30, 1993
<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072579 001</u>	Jan 08, 1991
<u>AB</u>		<u>20MG</u>	<u>A072556 001</u>	Sep 20, 1990
<u>AB</u>	HERITAGE PHARMA	<u>10MG</u>	<u>A202644 001</u>	Apr 25, 2013
<u>AB</u>		<u>20MG</u>	<u>A202644 002</u>	Apr 25, 2013
<u>AB</u>	LEADING	<u>10MG</u>	<u>A073250 001</u>	Oct 08, 1991
<u>AB</u>		<u>20MG</u>	<u>A074045 001</u>	Apr 30, 1992
	<u>PROCARDIA</u>			
<u>AB</u>	+	PFIZER	<u>10MG</u>	<u>N018482 001</u>

TABLET, EXTENDED RELEASE;ORAL

NIFEDIPINE

<u>AB1</u>	AUROBINDO PHARMA	<u>30MG</u>	<u>A213361 001</u>	Jul 19, 2021
<u>AB1</u>		<u>60MG</u>	<u>A213361 002</u>	Jul 19, 2021
<u>AB1</u>		<u>90MG</u>	<u>A213361 003</u>	Jul 19, 2021
<u>AB1</u>	NOVAST LABS	<u>30MG</u>	<u>A202987 001</u>	Aug 25, 2016
<u>AB1</u>	!	<u>60MG</u>	<u>A202987 002</u>	Aug 25, 2016

PRESCRIPTION DRUG PRODUCT LIST

NIFEDIPINE

TABLET, EXTENDED RELEASE;ORAL

NIFEDIPINE

<u>AB1</u>	!		<u>90MG</u>	<u>A202987</u>	<u>003</u>	Aug 25, 2016
<u>AB1</u>		VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075269</u>	<u>001</u>	Dec 04, 2000
<u>AB1</u>			<u>60MG</u>	<u>A075269</u>	<u>002</u>	Dec 04, 2000
<u>AB1</u>			<u>90MG</u>	<u>A076070</u>	<u>001</u>	Aug 16, 2002
<u>AB1</u>		ZYDUS PHARMS	<u>30MG</u>	<u>A210184</u>	<u>001</u>	Jun 29, 2018
<u>AB1</u>			<u>60MG</u>	<u>A210184</u>	<u>002</u>	Jun 29, 2018
<u>AB1</u>			<u>90MG</u>	<u>A210184</u>	<u>003</u>	Jun 29, 2018
<u>AB2</u>		ALEMBIC	<u>30MG</u>	<u>A216896</u>	<u>001</u>	Nov 18, 2022
<u>AB2</u>			<u>60MG</u>	<u>A216896</u>	<u>002</u>	Nov 18, 2022
<u>AB2</u>			<u>90MG</u>	<u>A216896</u>	<u>003</u>	Nov 18, 2022
<u>AB2</u>		ALKEM LABS LTD	<u>30MG</u>	<u>A216067</u>	<u>001</u>	Mar 29, 2022
<u>AB2</u>			<u>60MG</u>	<u>A216067</u>	<u>002</u>	Mar 29, 2022
<u>AB2</u>			<u>90MG</u>	<u>A216067</u>	<u>003</u>	Mar 29, 2022
<u>AB2</u>		ELITE PHARM SOLUTION	<u>90MG</u>	<u>A212016</u>	<u>001</u>	Nov 18, 2020
<u>AB2</u>		NOVAST LABS	<u>30MG</u>	<u>A210614</u>	<u>001</u>	Mar 12, 2019
<u>AB2</u>			<u>60MG</u>	<u>A210614</u>	<u>002</u>	Mar 12, 2019
<u>AB2</u>			<u>90MG</u>	<u>A210614</u>	<u>003</u>	Mar 12, 2019
<u>AB2</u>		OSMOTICA PHARM US	<u>30MG</u>	<u>A077127</u>	<u>001</u>	Nov 21, 2005
<u>AB2</u>			<u>60MG</u>	<u>A077127</u>	<u>002</u>	Nov 21, 2005
<u>AB2</u>			<u>90MG</u>	<u>A077127</u>	<u>003</u>	Oct 03, 2007
<u>AB2</u>		SPIL	<u>30MG</u>	<u>A210838</u>	<u>001</u>	Apr 16, 2019
<u>AB2</u>			<u>60MG</u>	<u>A210838</u>	<u>002</u>	Apr 16, 2019
<u>AB2</u>			<u>90MG</u>	<u>A210838</u>	<u>003</u>	Apr 16, 2019
<u>AB2</u>		TWI PHARMS	<u>30MG</u>	<u>A203126</u>	<u>001</u>	Apr 03, 2014
<u>AB2</u>			<u>60MG</u>	<u>A203126</u>	<u>002</u>	Apr 03, 2014
<u>AB2</u>			<u>90MG</u>	<u>A203126</u>	<u>003</u>	Apr 03, 2014
<u>AB2</u>		VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075289</u>	<u>002</u>	Feb 06, 2001
<u>AB2</u>			<u>60MG</u>	<u>A075289</u>	<u>001</u>	Sep 27, 2000
<u>AB2</u>		ZYDUS PHARMS	<u>30MG</u>	<u>A210012</u>	<u>001</u>	Dec 19, 2017
<u>AB2</u>			<u>60MG</u>	<u>A210012</u>	<u>002</u>	Dec 19, 2017
<u>AB2</u>			<u>90MG</u>	<u>A210012</u>	<u>003</u>	Dec 19, 2017
<u>PROCARDIA XL</u>						
<u>AB2</u>	+	PFIZER	<u>30MG</u>	<u>N019684</u>	<u>001</u>	Sep 06, 1989
<u>AB2</u>	+		<u>60MG</u>	<u>N019684</u>	<u>002</u>	Sep 06, 1989
<u>AB2</u>	+	!	<u>90MG</u>	<u>N019684</u>	<u>003</u>	Sep 06, 1989

NIFURTIMOX

TABLET;ORAL

LAMPIT

+	BAYER HEALTHCARE	30MG	N213464	001	Aug 06, 2020
+	!	120MG	N213464	002	Aug 06, 2020

NILOTINIB HYDROCHLORIDE

CAPSULE;ORAL

NILOTINIB HYDROCHLORIDE

<u>AB</u>		APOTEX	<u>EQ 50MG BASE</u>	<u>A203640</u>	<u>001</u>	Jan 05, 2024
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A203640</u>	<u>002</u>	Jan 05, 2024
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A203640</u>	<u>003</u>	Jan 05, 2024
<u>TASIGNA</u>						
<u>AB</u>	+	NOVARTIS	<u>EQ 50MG BASE</u>	<u>N022068</u>	<u>003</u>	Mar 22, 2018
<u>AB</u>	+		<u>EQ 150MG BASE</u>	<u>N022068</u>	<u>002</u>	Jun 17, 2010
<u>AB</u>	+	!	<u>EQ 200MG BASE</u>	<u>N022068</u>	<u>001</u>	Oct 29, 2007

NILUTAMIDE

TABLET;ORAL

NILANDRON

<u>AB</u>	+	CONCORDIA	<u>150MG</u>	<u>N020169</u>	<u>002</u>	Apr 30, 1999
<u>NILUTAMIDE</u>						
<u>AB</u>		ANI PHARMS	<u>150MG</u>	<u>A207631</u>	<u>001</u>	Jul 15, 2016

NIMODIPINE

CAPSULE;ORAL

NIMODIPINE

<u>AB</u>	!	BIONPHARMA	<u>30MG</u>	<u>A076740</u>	<u>001</u>	Jan 17, 2008
<u>AB</u>		HERITAGE PHARMS	<u>30MG</u>	<u>A077811</u>	<u>001</u>	May 02, 2007
<u>AB</u>		THEPHARMANETWORK LLC	<u>30MG</u>	<u>A090103</u>	<u>001</u>	Apr 07, 2014

PRESCRIPTION DRUG PRODUCT LIST

NIMODIPINE

SOLUTION;ORAL

NYMALIZE

+! AZURITY

6MG/ML

N203340 002 Apr 08, 2020

NINTEDANIB ESYLATE

CAPSULE;ORAL

OFEV

+ BOEHRINGER
INGELHEIM

EQ 100MG BASE

N205832 001 Oct 15, 2014

+!

EQ 150MG BASE

N205832 002 Oct 15, 2014

NIRAPARIB TOSYLATE

CAPSULE;ORAL

ZEJULA

+! GLAXOSMITHKLINE

EQ 100MG BASE

N208447 001 Mar 27, 2017

TABLET;ORAL

ZEJULA

+ GLAXOSMITHKLINE

EQ 100MG BASE

N214876 001 Apr 26, 2023

+

EQ 200MG BASE

N214876 002 Apr 26, 2023

+!

EQ 300MG BASE

N214876 003 Apr 26, 2023

NIRMATRELVIR; RITONAVIR

TABLET;ORAL

PAXLOVID (COPACKAGED)

+ PFIZER

150MG;100MG

N217188 001 May 25, 2023

+!

300MG;100MG

N217188 002 May 25, 2023

NIROGACESTAT HYDROBROMIDE

TABLET;ORAL

OGSIVEO

+! SPRINGWORKS

EQ 50MG BASE

N217677 001 Nov 27, 2023

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

NISOLDIPINEAB MYLAN8.5MGA091001 001 Jan 26, 2011AB17MGA091001 002 Jan 26, 2011AB34MGA091001 004 Jan 26, 2011SULARAB +! COVIS8.5MGN020356 008 Jan 02, 2008AB +!17MGN020356 007 Jan 02, 2008AB +!34MGN020356 005 Jan 02, 2008

NISOLDIPINE

! MYLAN

20MG

A079051 001 Jul 25, 2008

25.5MG

A091001 003 Jan 26, 2011

30MG

A079051 002 Jul 25, 2008

!

40MG

A079051 003 Jul 25, 2008

NITAZOXANIDE

FOR SUSPENSION;ORAL

ALINIA

+! ROMARK

100MG/5ML

N021498 001 Nov 22, 2002

TABLET;ORAL

ALINIAAB +! ROMARK500MGN021497 001 Jul 21, 2004NITAZOXANIDEAB RISING500MGA213820 001 Nov 27, 2020NITISINONE

CAPSULE;ORAL

NITISINONEAB ETON2MGA216201 001 May 25, 2023AB5MGA216201 002 May 25, 2023AB10MGA216201 003 May 25, 2023AB20MGA216201 004 May 25, 2023AB MEDUNIK2MGA212390 001 May 26, 2022AB5MGA212390 002 May 26, 2022AB10MGA212390 003 May 26, 2022AB20MGA212390 004 Apr 27, 2023AB NOVITIUM PHARMA2MGA211041 001 Aug 26, 2019AB5MGA211041 002 Aug 26, 2019AB10MGA211041 003 Aug 26, 2019AB TORRENT2MGA215908 001 Jan 09, 2023AB5MGA215908 002 Jan 09, 2023AB10MGA215908 003 Jan 09, 2023

PRESCRIPTION DRUG PRODUCT LIST

NITISINONE

CAPSULE; ORAL

NITISINONE

AB		20MG	A215908 004	Jan 09, 2023
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ORFADIN

AB	+	SWEDISH ORPHAN	2MG	N021232 001	Jan 18, 2002
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AB	+		5MG	N021232 002	Jan 18, 2002
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AB	+		10MG	N021232 003	Jan 18, 2002
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AB	+	!	20MG	N021232 004	Jun 13, 2016
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SUSPENSION; ORAL

ORFADIN

+	!	SWEDISH ORPHAN	4MG/ML	N206356 001	Apr 22, 2016
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TABLET; ORAL

NITYR

+		CYCLE	2MG	N209449 001	Jul 26, 2017
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+			5MG	N209449 002	Jul 26, 2017
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+	!		10MG	N209449 003	Jul 26, 2017
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NITRIC OXIDE

GAS; INHALATION

INOMAX

AA	+	!	MALLINCKRODT HOSP	800PPM	N020845 003	Dec 23, 1999
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NOXIVENT

AA			LINDE GAS EQUIP	800PPM	A207141 002	Oct 02, 2018
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ULSPIRA

AA			AIRGAS THERAP	800PPM	A203144 001	Jul 27, 2023
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GENOSYL

+	!	VERO BIOTECH INC	800PPM	N202860 001	Dec 20, 2019
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NOXIVENT

			LINDE GAS EQUIP	100PPM	A207141 001	Oct 02, 2018
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NITROFURANTOIN

SUSPENSION; ORAL

FURADANTIN

AB	+	CASPER PHARMA LLC	25MG/5ML	N009175 001	
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NITROFURANTOIN

AB		ACTAVIS MID ATLANTIC	25MG/5ML	A205180 001	May 03, 2016
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AB		AMNEAL PHARMS	25MG/5ML	A201679 001	May 11, 2011
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AB		AUROBINDO PHARMA	25MG/5ML	A212607 001	May 11, 2023
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AB		NOSTRUM LABS INC	25MG/5ML	A201355 001	Aug 14, 2013
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AB		NOVEL LABS INC	25MG/5ML	A201693 001	Sep 08, 2014
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AB		NOVITIUM PHARMA	25MG/5ML	A216385 001	Apr 14, 2023
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FURADANTIN

+	!	CASPER PHARMA LLC	50MG/5ML	N009175 002	Jun 09, 2023
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NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN

AB	+	ALMATICA	25MG	N016620 003	
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AB	+		50MG	N016620 001	
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AB	+	!	100MG	N016620 002	
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NITROFURANTOIN

AB		ACTAVIS LABS FL INC	25MG	A091095 001	Jun 18, 2015
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AB			50MG	A091095 002	Jun 18, 2015
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AB			100MG	A091095 003	Jun 18, 2015
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AB		ALEMBIC	25MG	A211935 001	Jun 25, 2021
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AB			50MG	A211935 002	Jun 25, 2021
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AB			100MG	A211935 003	Jun 25, 2021
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AB		AUROBINDO PHARMA USA	50MG	A074967 001	Jul 09, 1997
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AB			100MG	A074967 002	Jul 09, 1997
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AB			100MG	A077025 001	Aug 18, 2004
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AB		IMPAX LABS INC	50MG	A073671 001	Jan 28, 1993
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AB			100MG	A073652 001	Jan 28, 1993
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AB		MANKIND PHARMA	25MG	A217272 001	Mar 21, 2023
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AB			50MG	A217272 002	Mar 21, 2023
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AB			100MG	A217272 003	Mar 21, 2023
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AB		NOVEL LABS INC	50MG	A203233 001	Jul 09, 2018
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AB			100MG	A203233 002	Jul 09, 2018
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AB		SUN PHARM INDUSTRIES	25MG	A201722 001	Feb 16, 2016
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AB			50MG	A201722 002	Feb 16, 2016
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AB			100MG	A201722 003	Feb 16, 2016
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PRESCRIPTION DRUG PRODUCT LIST

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN

AB	ZYDUS PHARMS	50MG	A205005 001	Dec 12, 2017
AB		100MG	A205005 002	Dec 12, 2017

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID

AB	+ ! ALMATICA	75MG; 25MG	N020064 001	Dec 24, 1991
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NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)

AB	AMNEAL PHARMS	75MG; 25MG	A207372 001	May 15, 2017
AB	AUROBINDO PHARMA	75MG; 25MG	A209225 001	Mar 30, 2023
AB	INVENTIA	75MG; 25MG	A211013 001	Feb 18, 2022
AB	MANKIND PHARMA	75MG; 25MG	A217357 001	Jul 11, 2023
AB	OMSAV PHARMA	75MG; 25MG	A217073 001	Nov 03, 2023
AB	SANDOZ	75MG; 25MG	A077066 001	Apr 05, 2005
AB	SUNNY	75MG; 25MG	A208516 001	May 24, 2018
AB	WATSON LABS INC	75MG; 25MG	A202250 001	Jul 08, 2015

NITROGLYCERIN

AEROSOL, METERED; SUBLINGUAL

NITROMIST

+ !	EVUS	0.4MG/SPRAY	N021780 001	Nov 02, 2006
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FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN

AB2	! MYLAN TECHNOLOGIES	0.1MG/HR	A074559 004	Feb 06, 1998
AB2	!	0.2MG/HR	A074559 003	Aug 30, 1996
AB2	!	0.4MG/HR	A074559 002	Aug 30, 1996
AB2	!	0.6MG/HR	A074559 001	Aug 30, 1996
AB2	ZYDUS PHARMS	0.1MG/HR	A089885 002	Oct 30, 2017
AB2		0.2MG/HR	A089884 001	Oct 30, 1998
AB2		0.4MG/HR	A089885 001	Oct 30, 1998
AB2		0.6MG/HR	A089886 001	Oct 30, 1998

NITRO-DUR

+ !	USEPHARMA	0.1MG/HR	N020145 001	Apr 04, 1995
+ !		0.2MG/HR	N020145 002	Apr 04, 1995
+ !		0.3MG/HR	N020145 003	Apr 04, 1995
+ !		0.4MG/HR	N020145 004	Apr 04, 1995
+ !		0.6MG/HR	N020145 005	Apr 04, 1995
+ !		0.8MG/HR	N020145 006	Apr 04, 1995

INJECTABLE; INJECTION

NITROGLYCERIN

!	AM REGENT	5MG/ML	A072034 001	May 24, 1988
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NITROGLYCERIN IN DEXTROSE 5%

+ !	BAXTER HLTHCARE	10MG/100ML	N019970 001	Dec 29, 1989
+ !		20MG/100ML	N019970 002	Dec 29, 1989
+ !		40MG/100ML	N019970 003	Dec 29, 1989

OINTMENT; INTRA-ANAL

RECTIV

+ !	ABEVIE	0.4%	N021359 001	Jun 21, 2011
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OINTMENT; TRANSDERMAL

NITROGLYCERIN

+ !	FOUGERA PHARMS INC	2%	A087355 001	Jul 08, 1988
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SPRAY, METERED; SUBLINGUAL

NITROGLYCERIN

AB	PADAGIS ISRAEL	0.4MG/SPRAY	A091496 001	Sep 20, 2013
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NITROLINGUAL PUMPSPRAY

AB	+ ! POHL BOSKAMP	0.4MG/SPRAY	N018705 002	Jan 10, 1997
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TABLET; SUBLINGUAL

NITROGLYCERIN

AB	AUROBINDO PHARMA	0.3MG	A217879 001	Oct 26, 2023
AB		0.4MG	A217879 002	Oct 26, 2023
AB		0.6MG	A217879 003	Oct 26, 2023
AB	DR REDDYS	0.3MG	A208191 001	Aug 26, 2016
AB		0.4MG	A208191 002	Aug 26, 2016
AB		0.6MG	A208191 003	Aug 26, 2016
AB	GLENMARK PHARMS SA	0.3MG	A206391 001	Sep 19, 2017
AB		0.4MG	A206391 002	Sep 19, 2017
AB		0.6MG	A206391 003	Sep 19, 2017
AB	MANKIND PHARMA	0.3MG	A217970 001	Dec 05, 2023
AB		0.4MG	A217970 002	Dec 05, 2023
AB		0.6MG	A217970 003	Dec 05, 2023

PRESCRIPTION DRUG PRODUCT LIST

NITROGLYCERIN

TABLET;SUBLINGUAL

NITROGLYCERIN

<u>AB</u>	NATCO	<u>0.3MG</u>	<u>A211604 001</u>	Apr 30, 2019
<u>AB</u>		<u>0.4MG</u>	<u>A211604 002</u>	Apr 30, 2019
<u>AB</u>		<u>0.6MG</u>	<u>A211604 003</u>	Apr 30, 2019
<u>AB</u>	RUBICON	<u>0.3MG</u>	<u>A209779 001</u>	May 03, 2021
<u>AB</u>		<u>0.4MG</u>	<u>A209779 002</u>	May 03, 2021
<u>AB</u>		<u>0.6MG</u>	<u>A209779 003</u>	May 03, 2021
<u>NITROSTAT</u>				
<u>AB</u>	+ VIATRIS	<u>0.3MG</u>	<u>N021134 001</u>	May 01, 2000
<u>AB</u>	+	<u>0.4MG</u>	<u>N021134 002</u>	May 01, 2000
<u>AB</u>	+	<u>0.6MG</u>	<u>N021134 003</u>	May 01, 2000

NIZATIDINE

CAPSULE;ORAL

NIZATIDINE

<u>AB</u>	DR REDDYS LABS LTD	<u>150MG</u>	<u>A077314 001</u>	Sep 15, 2005
<u>AB</u>		<u>300MG</u>	<u>A077314 002</u>	Sep 15, 2005
<u>AB</u>	EPIC PHARMA LLC	<u>150MG</u>	<u>A076178 001</u>	Jul 05, 2002
<u>AB</u>		<u>300MG</u>	<u>A076178 002</u>	Jul 05, 2002
<u>AB</u>	GLENMARK GENERICS	<u>150MG</u>	<u>A090618 001</u>	Jul 15, 2011
<u>AB</u>		<u>300MG</u>	<u>A090618 002</u>	Jul 15, 2011
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A075616 001</u>	Jul 09, 2002
<u>AB</u>	!	<u>300MG</u>	<u>A075616 002</u>	Jul 09, 2002

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

LEVOPHED

<u>AP</u>	+	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N007513 001</u>
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NOREPINEPHRINE BITARTRATE

<u>AP</u>	AMNEAL	<u>EQ 1MG BASE/ML</u>	<u>A210839 001</u>	Dec 17, 2018
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 1MG BASE/ML</u>	<u>A040859 001</u>	Mar 27, 2012
<u>AP</u>	BRECKENRIDGE	<u>EQ 1MG BASE/ML</u>	<u>A214455 001</u>	Jan 22, 2021
<u>AP</u>	CAPLIN	<u>EQ 1MG BASE/ML</u>	<u>A217575 001</u>	Sep 15, 2023
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A211382 001</u>	Nov 03, 2020
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 1MG BASE/ML</u>	<u>A214323 001</u>	May 06, 2021
<u>AP</u>	HIKMA	<u>EQ 1MG BASE/ML</u>	<u>A040462 001</u>	Oct 31, 2003
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A203662 001</u>	Nov 07, 2018
<u>AP</u>	MEITHEAL	<u>EQ 1MG BASE/ML</u>	<u>A040455 001</u>	Mar 03, 2003
<u>AP</u>	MYLAN LABS LTD	<u>EQ 1MG BASE/ML</u>	<u>A211242 001</u>	Oct 04, 2018
<u>AP</u>	SANDOZ	<u>EQ 1MG BASE/ML</u>	<u>A211359 001</u>	Oct 18, 2018
<u>AP</u>	SUN PHARM	<u>EQ 1MG BASE/ML</u>	<u>A211980 001</u>	Jan 29, 2021

SOLUTION; INTRAVENOUS

NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE

+	INFORLIFE	<u>EQ 4MG BASE/250ML (EQ 16MCG BASE/ML)</u>	<u>N215700 001</u>	Sep 15, 2022
+		<u>EQ 8MG BASE/250ML (EQ 32MCG BASE/ML)</u>	<u>N215700 002</u>	Sep 15, 2022
+		<u>EQ 16MG BASE/250ML (EQ 64MCG BASE/ML)</u>	<u>N215700 003</u>	Sep 15, 2022

NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE

+	BAXTER HLTHCARE CORP	<u>EQ 4MG BASE/250ML (EQ 16MCG BASE/ML)</u>	<u>N214313 001</u>	Jan 15, 2021
+		<u>EQ 8MG BASE/250ML (EQ 32MCG BASE/ML)</u>	<u>N214313 002</u>	Jan 15, 2021
+		<u>EQ 16MG BASE/250ML (EQ 64MCG BASE/ML)</u>	<u>N214313 003</u>	Nov 21, 2023

NORETHINDRONE

TABLET;ORAL-28

CAMILA

<u>AB1</u>	DR REDDYS LABS SA	<u>0.35MG</u>	<u>A076177 001</u>	Oct 21, 2002
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HEATHER

<u>AB1</u>	GLENMARK GENERICS	<u>0.35MG</u>	<u>A090454 001</u>	Apr 23, 2010
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INCASSIA

<u>AB1</u>	AUROBINDO PHARMA	<u>0.35MG</u>	<u>A207304 001</u>	Sep 23, 2016
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NOR-QD

<u>AB1</u>	+	TEVA BRANDED PHARM	<u>0.35MG</u>	<u>N017060 001</u>
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NORETHINDRONE

<u>AB1</u>	LUPIN LTD	<u>0.35MG</u>	<u>A091325 001</u>	Sep 19, 2011
<u>AB1</u>	MYLAN LABS LTD	<u>0.35MG</u>	<u>A201483 001</u>	Jun 24, 2013
<u>AB1</u>	NAARI PTE LTD	<u>0.35MG</u>	<u>A206807 001</u>	Dec 13, 2016
<u>AB1</u>	NOVAST LABS	<u>0.35MG</u>	<u>A202014 001</u>	Sep 13, 2013

EMZAHH

<u>AB2</u>	AUROBINDO PHARMA	<u>0.35MG</u>	<u>A216796 001</u>	Jan 06, 2023
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ERRIN

<u>AB2</u>	DR REDDYS LABS SA	<u>0.35MG</u>	<u>A076225 001</u>	Oct 21, 2002
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PRESCRIPTION DRUG PRODUCT LIST

NORETHINDRONE

TABLET; ORAL-28

JENCYCLA

<u>AB2</u>	LUPIN LTD	<u>0.35MG</u>	<u>A091323</u>	<u>001</u>	Mar 28, 2013
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NORETHINDRONE

<u>AB2</u>	GLENMARK GENERICS	<u>0.35MG</u>	<u>A091209</u>	<u>001</u>	Jul 22, 2010
<u>AB2</u>	! MYLAN LABS LTD	<u>0.35MG</u>	<u>A200980</u>	<u>001</u>	Jun 12, 2013
<u>AB2</u>	NOVAST LABS	<u>0.35MG</u>	<u>A200961</u>	<u>001</u>	Sep 13, 2013

NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE

<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A200275</u>	<u>001</u>	Jul 30, 2012
<u>AB</u>	! BARR	<u>5MG</u>	<u>A075951</u>	<u>001</u>	May 25, 2001
<u>AB</u>	GLENMARK GENERICS	<u>5MG</u>	<u>A091090</u>	<u>001</u>	Jul 21, 2010
<u>AB</u>	MYLAN LABS LTD	<u>5MG</u>	<u>A205278</u>	<u>001</u>	Nov 10, 2016
<u>AB</u>	NOVAST LABS	<u>5MG</u>	<u>A206490</u>	<u>001</u>	Nov 05, 2018

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

NORTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS SA	<u>EQ 10MG BASE</u>	<u>A073556</u>	<u>002</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A073556</u>	<u>003</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A073556</u>	<u>004</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A073556</u>	<u>001</u>	Mar 30, 1992
<u>AB</u>	TARO	<u>EQ 10MG BASE</u>	<u>A075520</u>	<u>004</u>	May 08, 2000
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A075520</u>	<u>003</u>	May 08, 2000
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A075520</u>	<u>001</u>	May 08, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A075520</u>	<u>002</u>	May 08, 2000
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A074132</u>	<u>001</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A074132</u>	<u>002</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A074132</u>	<u>003</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A074132</u>	<u>004</u>	Mar 27, 1995

PAMELOR

<u>AB</u>	+ SPECGX LLC	<u>EQ 10MG BASE</u>	<u>N018013</u>	<u>001</u>	
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N018013</u>	<u>002</u>	
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N018013</u>	<u>004</u>	
<u>AB</u>	+!	<u>EQ 75MG BASE</u>	<u>N018013</u>	<u>003</u>	

SOLUTION; ORAL

NORTRIPTYLINE HYDROCHLORIDE

<u>AA</u>	! PHARM ASSOC	<u>EQ 10MG BASE/5ML</u>	<u>A075606</u>	<u>001</u>	Aug 23, 2000
<u>AA</u>	RUBICON	<u>EQ 10MG BASE/5ML</u>	<u>A217731</u>	<u>001</u>	Aug 15, 2023

NUSINERSEN SODIUM

SOLUTION; INTRATHECAL

SPINRAZA

+	! BIOGEN IDEC	<u>EQ 12MG BASE/5ML (EQ 2.4MG BASE/ML)</u>	<u>N209531</u>	<u>001</u>	Dec 23, 2016
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NYSTATIN

CREAM; TOPICAL

NYSTATIN

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062949</u>	<u>001</u>	Jun 13, 1988
<u>AT</u>	COSETTE	<u>100,000 UNITS/GM</u>	<u>A061966</u>	<u>001</u>	
<u>AT</u>	CROWN LABS INC	<u>100,000 UNITS/GM</u>	<u>A207733</u>	<u>001</u>	Sep 26, 2017
<u>AT</u>	FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<u>A062129</u>	<u>001</u>	
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM</u>	<u>A213566</u>	<u>001</u>	Aug 10, 2021
<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A062225</u>	<u>001</u>	
<u>AT</u>	! TARO	<u>100,000 UNITS/GM</u>	<u>A064022</u>	<u>001</u>	Jan 29, 1993
<u>AT</u>	TORRENT	<u>100,000 UNITS/GM</u>	<u>A212557</u>	<u>001</u>	Jul 24, 2019

OINTMENT; TOPICAL

NYSTATIN

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062840</u>	<u>001</u>	Nov 13, 1987
<u>AT</u>	CADILA	<u>100,000 UNITS/GM</u>	<u>A207767</u>	<u>001</u>	May 25, 2018
<u>AT</u>	COSETTE	<u>100,000 UNITS/GM</u>	<u>A209114</u>	<u>001</u>	Oct 06, 2017
<u>AT</u>	! FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<u>A062124</u>	<u>002</u>	Sep 23, 1982
<u>AT</u>	LYNE	<u>100,000 UNITS/GM</u>	<u>A209082</u>	<u>001</u>	May 21, 2018
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM</u>	<u>A213826</u>	<u>001</u>	Jan 14, 2021
<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A062472</u>	<u>001</u>	Feb 13, 1984

POWDER; TOPICAL

NYSTATIN

<u>AT</u>	BEOWULF ASSET	<u>100,000 UNITS/GM</u>	<u>A065175</u>	<u>001</u>	Dec 17, 2004
<u>AT</u>	! DR REDDYS LABS SA	<u>100,000 UNITS/GM</u>	<u>A065203</u>	<u>001</u>	Jul 15, 2004
<u>AT</u>	EPIC PHARMA LLC	<u>100,000 UNITS/GM</u>	<u>A210532</u>	<u>001</u>	Apr 30, 2018

PRESCRIPTION DRUG PRODUCT LIST

NYSTATIN

POWDER; TOPICAL

NYSTATIN

<u>AT</u>	LUPIN	<u>100,000 UNITS/GM</u>	<u>A065138 001</u>	Jul 23, 2004
<u>AT</u>	LYNE	<u>100,000 UNITS/GM</u>	<u>A208838 001</u>	May 30, 2017
<u>AT</u>	UPSHER SMITH LABS	<u>100,000 UNITS/GM</u>	<u>A065183 001</u>	May 03, 2005
<u>AT</u>	ZYDUS PHARMS	<u>100,000 UNITS/GM</u>	<u>A208581 001</u>	Jun 08, 2017

NYSTOP

<u>AT</u>	PADAGIS US	<u>100,000 UNITS/GM</u>	<u>A064118 001</u>	Aug 16, 1996
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SUSPENSION; ORAL

NYSTATIN

<u>AA</u>	FOUGERA PHARMS INC	<u>100,000 UNITS/ML</u>	<u>A062517 001</u>	Jun 07, 1984
<u>AA</u>	GENUS	<u>100,000 UNITS/ML</u>	<u>A065148 001</u>	Jun 28, 2005
<u>AA</u>	LEADING	<u>100,000 UNITS/ML</u>	<u>A214346 001</u>	Mar 10, 2022
<u>AA</u>	MLV	<u>100,000 UNITS/ML</u>	<u>A062832 001</u>	Dec 27, 1991
<u>AA</u>	PHARM ASSOC	<u>100,000 UNITS/ML</u>	<u>A203621 001</u>	Jan 07, 2016
<u>AA</u>	TARO	<u>100,000 UNITS/ML</u>	<u>A062876 001</u>	Feb 29, 1988
<u>AA</u>	! WOCKHARDT BIO AG	<u>100,000 UNITS/ML</u>	<u>A062512 001</u>	Oct 29, 1984

TABLET; ORAL

NYSTATIN

<u>AA</u>	HERITAGE PHARMS	<u>500,000 UNITS</u>	<u>A062474 001</u>	Dec 22, 1983
<u>AA</u>	SUN PHARM	<u>500,000 UNITS</u>	<u>A062838 001</u>	Dec 22, 1988
<u>AA</u>	! TEVA	<u>500,000 UNITS</u>	<u>A062506 001</u>	Jan 16, 1984

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYKACET

<u>AT</u>	COSETTE	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062367 001</u>	May 28, 1985
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	ALEMBIC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214090 001</u>	Mar 31, 2021
<u>AT</u>	AMNEAL	<u>100,000 UNITS/GM; 0.1%</u>	<u>A209990 001</u>	Feb 15, 2018
<u>AT</u>	CROWN LABS INC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207730 001</u>	Dec 26, 2017
<u>AT</u>	DR REDDYS	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208326 001</u>	Oct 26, 2016
<u>AT</u>	FOUGERA PHARMS INC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062599 001</u>	Oct 08, 1985
<u>AT</u>	GLENMARK PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208136 001</u>	Oct 24, 2016
<u>AT</u>	LUPIN LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208205 001</u>	May 31, 2018
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214181 001</u>	Jul 13, 2022
<u>AT</u>	PADAGIS ISRAEL	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208479 001</u>	Aug 14, 2017
<u>AT</u>	! TARO	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062364 001</u>	Dec 22, 1987

OINTMENT; TOPICAL

MYKACET

<u>AT</u>	COSETTE	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062733 001</u>	Mar 06, 1987
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	ALEMBIC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214186 001</u>	Mar 04, 2022
<u>AT</u>	CADILA	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207764 001</u>	Nov 08, 2018
<u>AT</u>	DR REDDYS	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207741 001</u>	Jan 31, 2017
<u>AT</u>	EPIC PHARMA LLC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207217 001</u>	Aug 04, 2017
<u>AT</u>	FOUGERA PHARMS INC	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062602 001</u>	Oct 09, 1985
<u>AT</u>	GLENMARK PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A208300 001</u>	Jun 23, 2016
<u>AT</u>	MACLEODS PHARMS LTD	<u>100,000 UNITS/GM; 0.1%</u>	<u>A214751 001</u>	Aug 17, 2021
<u>AT</u>	PADAGIS ISRAEL	<u>100,000 UNITS/GM; 0.1%</u>	<u>A207380 001</u>	Dec 20, 2016
<u>AT</u>	RISING	<u>100,000 UNITS/GM; 0.1%</u>	<u>A206785 001</u>	Dec 29, 2016
<u>AT</u>	STRIDES PHARMA	<u>100,000 UNITS/GM; 0.1%</u>	<u>A210077 001</u>	Jan 29, 2018
<u>AT</u>	! TARO	<u>100,000 UNITS/GM; 0.1%</u>	<u>A063305 001</u>	Mar 29, 1993

OBETICHOIC ACID

TABLET; ORAL

OCALIVA

+	INTERCEPT PHARMS	5MG	N207999 001	May 27, 2016
+	INC			
+	!	10MG	N207999 002	May 27, 2016

OCTREOTIDE ACETATE

CAPSULE, DELAYED RELEASE; ORAL

MYCAPSSA

+	!	CHIESI	EQ 20MG BASE	N208232 001	Jun 26, 2020
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INJECTABLE; INJECTION

OCTREOTIDE ACETATE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 0.2MG BASE/ML</u>	<u>A077450 001</u>	Feb 10, 2006
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A077450 002</u>	Feb 10, 2006
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 0.1MG BASE/ML</u>	<u>A216839 002</u>	Jun 22, 2023
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>A216807 001</u>	Jun 13, 2023
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A216839 001</u>	Jun 22, 2023

PRESCRIPTION DRUG PRODUCT LIST

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A216807 002</u>	Jun 13, 2023
<u>AP</u>	HERITAGE PHARMS	<u>EQ 0.05MG BASE/ML</u>	<u>A204669 001</u>	Dec 27, 2018
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A204669 002</u>	Dec 27, 2018
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>A203765 001</u>	Sep 07, 2018
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A204669 003</u>	Dec 27, 2018
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A203765 002</u>	Sep 07, 2018
<u>AP</u>	MEITHEAL	<u>EQ 0.05MG BASE/ML</u>	<u>A075957 001</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A075957 002</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>A075959 001</u>	Nov 21, 2005
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A075957 003</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075959 002</u>	Nov 21, 2005
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 0.2MG BASE/ML</u>	<u>A091041 001</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A091041 002</u>	Nov 12, 2013
<u>AP</u>	TEVA PHARMS USA INC	<u>EQ 10MG BASE/VIAL</u>	<u>A210317 001</u>	Dec 05, 2023
<u>AP</u>		<u>EQ 20MG BASE/VIAL</u>	<u>A210317 002</u>	Dec 05, 2023
<u>AP</u>		<u>EQ 30MG BASE/VIAL</u>	<u>A210317 003</u>	Dec 05, 2023
<u>AP</u>	! WEST-WARD PHARMS INT	<u>EQ 0.2MG BASE/ML</u>	<u>A076330 001</u>	Apr 08, 2005
<u>AP</u>	!	<u>EQ 1MG BASE/ML</u>	<u>A076330 002</u>	Apr 08, 2005

OCTREOTIDE ACETATE (PRESERVATIVE FREE)

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 0.05MG BASE/ML</u>	<u>A077457 001</u>	Feb 10, 2006
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A077457 002</u>	Feb 10, 2006
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A077457 003</u>	Feb 10, 2006
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 0.05MG BASE/ML</u>	<u>A079198 001</u>	Feb 10, 2011
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A079198 002</u>	Feb 10, 2011
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A079198 003</u>	Feb 10, 2011
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 0.05MG BASE/ML</u>	<u>A090834 001</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A090834 002</u>	Nov 12, 2013
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A090834 003</u>	Nov 12, 2013
<u>AP</u>	! WEST-WARD PHARMS INT	<u>EQ 0.05MG BASE/ML</u>	<u>A076313 001</u>	Mar 28, 2005
<u>AP</u>	!	<u>EQ 0.1MG BASE/ML</u>	<u>A076313 003</u>	Mar 28, 2005
<u>AP</u>	!	<u>EQ 0.5MG BASE/ML</u>	<u>A076313 002</u>	Mar 28, 2005

SANDOSTATIN

<u>AP</u>	+! NOVARTIS	<u>EQ 0.05MG BASE/ML</u>	<u>N019667 001</u>	Oct 21, 1988
<u>AP</u>	+!	<u>EQ 0.1MG BASE/ML</u>	<u>N019667 002</u>	Oct 21, 1988
<u>AP</u>	+!	<u>EQ 0.5MG BASE/ML</u>	<u>N019667 003</u>	Oct 21, 1988
<u>AP</u>	+ NOVARTIS	<u>EQ 10MG BASE/VIAL</u>	<u>N021008 001</u>	Nov 25, 1998
<u>AP</u>	+	<u>EQ 20MG BASE/VIAL</u>	<u>N021008 002</u>	Nov 25, 1998
<u>AP</u>	+!	<u>EQ 30MG BASE/VIAL</u>	<u>N021008 003</u>	Nov 25, 1998

ODEVIXIBAT

CAPSULE; ORAL

BYLVAY

+	ALBIREO	0.4MG	N215498 002	Jul 20, 2021
+	!	1.2MG	N215498 004	Jul 20, 2021

CAPSULE, PELLETS; ORAL

BYLVAY

+	ALBIREO	0.2MG	N215498 001	Jul 20, 2021
+	!	0.6MG	N215498 003	Jul 20, 2021

OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OCUFLOX

<u>AT</u>	+! ALLERGAN	<u>0.3%</u>	<u>N019921 001</u>	Jul 30, 1993
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OFLOXACIN

<u>AT</u>	ALTAIRE PHARMS INC	<u>0.3%</u>	<u>A202692 001</u>	Apr 29, 2013
<u>AT</u>	APOTEX INC	<u>0.3%</u>	<u>A076513 001</u>	May 14, 2004
<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>A076622 001</u>	May 14, 2004
<u>AT</u>	FDC LTD	<u>0.3%</u>	<u>A078559 001</u>	Feb 25, 2009
<u>AT</u>	MANKIND PHARMA	<u>0.3%</u>	<u>A215886 001</u>	Nov 23, 2022
<u>AT</u>	SENTISS	<u>0.3%</u>	<u>A076407 001</u>	Apr 15, 2008

SOLUTION/DROPS; OTIC

OFLOXACIN

<u>AT</u>	AMNEAL	<u>0.3%</u>	<u>A211525 001</u>	Aug 30, 2019
<u>AT</u>	APOTEX INC	<u>0.3%</u>	<u>A076527 001</u>	Sep 28, 2007
<u>AT</u>	! BAUSCH AND LOMB	<u>0.3%</u>	<u>A076128 001</u>	Mar 17, 2008
<u>AT</u>	CAPLIN	<u>0.3%</u>	<u>A217903 001</u>	Jan 05, 2024
<u>AT</u>	HIKMA	<u>0.3%</u>	<u>A076616 001</u>	Mar 17, 2008
<u>AT</u>	MANKIND PHARMA	<u>0.3%</u>	<u>A216130 001</u>	Jul 14, 2022

PRESCRIPTION DRUG PRODUCT LIST

OFLOXACIN

TABLET; ORAL

OFLOXACIN

<u>AB</u>	CADILA PHARMS LTD	<u>200MG</u>	<u>A091656 001</u>	Sep 18, 2014
<u>AB</u>		<u>300MG</u>	<u>A091656 002</u>	Sep 18, 2014
<u>AB</u>		<u>400MG</u>	<u>A091656 003</u>	Sep 18, 2014
<u>AB</u>	DR REDDYS LABS LTD	<u>200MG</u>	<u>A077098 001</u>	Feb 10, 2006
<u>AB</u>		<u>300MG</u>	<u>A077098 002</u>	Feb 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077098 003</u>	Feb 10, 2006
<u>AB</u>	LARKEN LABS	<u>400MG</u>	<u>A076093 003</u>	Sep 02, 2003
<u>AB</u>	TEVA	<u>200MG</u>	<u>A076182 001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>A076182 002</u>	Sep 02, 2003
<u>AB</u>	!	<u>400MG</u>	<u>A076182 003</u>	Sep 02, 2003

OLANZAPINE

INJECTABLE; INTRAMUSCULAR

OLANZAPINE

<u>AP</u>	AM REGENT	<u>10MG/VIAL</u>	<u>A201741 001</u>	Mar 20, 2012
<u>AP</u>	ASPIRO	<u>10MG/VIAL</u>	<u>A217466 001</u>	Mar 22, 2023
<u>AP</u>	EUGIA PHARMA	<u>10MG/VIAL</u>	<u>A210968 001</u>	Oct 22, 2020
<u>AP</u>	SANDOZ INC	<u>10MG/VIAL</u>	<u>A201588 001</u>	Oct 24, 2011

ZYPREXA

<u>AP</u>	+! CHEPLAPHARM	<u>10MG/VIAL</u>	<u>N021253 001</u>	Mar 29, 2004
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TABLET; ORAL

OLANZAPINE

<u>AB</u>	ALKEM LABS LTD	<u>2.5MG</u>	<u>A202295 001</u>	Oct 20, 2015
<u>AB</u>		<u>5MG</u>	<u>A202295 002</u>	Oct 20, 2015
<u>AB</u>		<u>7.5MG</u>	<u>A202295 003</u>	Oct 20, 2015
<u>AB</u>		<u>10MG</u>	<u>A202295 004</u>	Oct 20, 2015
<u>AB</u>		<u>15MG</u>	<u>A202295 005</u>	Oct 20, 2015
<u>AB</u>		<u>20MG</u>	<u>A202295 006</u>	Oct 20, 2015
<u>AB</u>	APOTEX INC	<u>2.5MG</u>	<u>A090798 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A090798 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A090798 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A090798 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A090798 005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A090798 006</u>	Apr 23, 2012
<u>AB</u>	AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A202050 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A202050 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A202050 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A202050 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A202050 005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A202050 006</u>	Apr 23, 2012
<u>AB</u>	CADILA PHARMS LTD	<u>2.5MG</u>	<u>A210022 001</u>	Feb 24, 2023
<u>AB</u>		<u>5MG</u>	<u>A210022 002</u>	Feb 24, 2023
<u>AB</u>		<u>7.5MG</u>	<u>A210022 003</u>	Feb 24, 2023
<u>AB</u>		<u>10MG</u>	<u>A210022 004</u>	Feb 24, 2023
<u>AB</u>		<u>15MG</u>	<u>A210022 005</u>	Feb 24, 2023
<u>AB</u>		<u>20MG</u>	<u>A210022 006</u>	Feb 24, 2023
<u>AB</u>	CHARTWELL MOLECULAR	<u>2.5MG</u>	<u>A203333 001</u>	Mar 15, 2016
<u>AB</u>		<u>5MG</u>	<u>A203333 002</u>	Mar 15, 2016
<u>AB</u>		<u>7.5MG</u>	<u>A203333 003</u>	Mar 15, 2016
<u>AB</u>		<u>10MG</u>	<u>A203333 004</u>	Mar 15, 2016
<u>AB</u>		<u>15MG</u>	<u>A203333 005</u>	Mar 15, 2016
<u>AB</u>		<u>20MG</u>	<u>A203333 006</u>	Mar 15, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>2.5MG</u>	<u>A076255 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A076255 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A076255 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A076255 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A076133 001</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A076133 002</u>	Oct 24, 2011
<u>AB</u>	INDOCO	<u>2.5MG</u>	<u>A206155 001</u>	Jul 31, 2020
<u>AB</u>		<u>5MG</u>	<u>A206155 002</u>	Jul 31, 2020
<u>AB</u>		<u>7.5MG</u>	<u>A206155 003</u>	Jul 31, 2020
<u>AB</u>		<u>10MG</u>	<u>A206155 004</u>	Jul 31, 2020
<u>AB</u>		<u>15MG</u>	<u>A206155 005</u>	Jul 31, 2020
<u>AB</u>		<u>20MG</u>	<u>A206155 006</u>	Jul 31, 2020
<u>AB</u>	MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A202862 001</u>	Aug 15, 2014
<u>AB</u>		<u>5MG</u>	<u>A202862 002</u>	Aug 15, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A202862 003</u>	Aug 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A202862 004</u>	Aug 15, 2014
<u>AB</u>		<u>15MG</u>	<u>A202862 005</u>	Aug 15, 2014

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET; ORAL

OLANZAPINE

<u>AB</u>		<u>20MG</u>	<u>A202862</u>	<u>006</u>	Aug 15, 2014
<u>AB</u>	ORBION PHARMS	<u>2.5MG</u>	<u>A202287</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A202287</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A202287</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A202287</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A202287</u>	<u>005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A202287</u>	<u>006</u>	Apr 23, 2012
<u>AB</u>	QILU	<u>2.5MG</u>	<u>A204319</u>	<u>001</u>	Jan 27, 2016
<u>AB</u>		<u>5MG</u>	<u>A204319</u>	<u>002</u>	Jan 27, 2016
<u>AB</u>		<u>7.5MG</u>	<u>A204319</u>	<u>003</u>	Jan 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A204319</u>	<u>004</u>	Jan 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A204319</u>	<u>005</u>	Jan 27, 2016
<u>AB</u>		<u>20MG</u>	<u>A204319</u>	<u>006</u>	Jan 27, 2016
<u>AB</u>	SUN PHARM INDS	<u>2.5MG</u>	<u>A091038</u>	<u>001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A091038</u>	<u>002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A091038</u>	<u>003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A091038</u>	<u>004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A091038</u>	<u>005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A091038</u>	<u>006</u>	Apr 23, 2012
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A090459</u>	<u>001</u>	Jul 16, 2018
<u>AB</u>		<u>5MG</u>	<u>A090459</u>	<u>002</u>	Jul 16, 2018
<u>AB</u>		<u>7.5MG</u>	<u>A090459</u>	<u>003</u>	Jul 16, 2018
<u>AB</u>		<u>10MG</u>	<u>A090459</u>	<u>004</u>	Jul 16, 2018
<u>AB</u>		<u>15MG</u>	<u>A090459</u>	<u>005</u>	Jul 16, 2018
<u>AB</u>		<u>20MG</u>	<u>A090459</u>	<u>006</u>	Jul 16, 2018

ZYPREXA

<u>AB</u>	+	CHEPLAPHARM	<u>2.5MG</u>	<u>N020592</u>	<u>001</u>	Sep 30, 1996
<u>AB</u>	+	!	<u>5MG</u>	<u>N020592</u>	<u>002</u>	Sep 30, 1996
<u>AB</u>	+		<u>7.5MG</u>	<u>N020592</u>	<u>003</u>	Sep 30, 1996
<u>AB</u>	+		<u>10MG</u>	<u>N020592</u>	<u>004</u>	Sep 30, 1996
<u>AB</u>	+		<u>15MG</u>	<u>N020592</u>	<u>005</u>	Sep 09, 1997
<u>AB</u>	+		<u>20MG</u>	<u>N020592</u>	<u>006</u>	Sep 09, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

OLANZAPINE

<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A091265</u>	<u>001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A091265</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A091265</u>	<u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A091265</u>	<u>004</u>	Oct 24, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A203708</u>	<u>001</u>	May 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A203708</u>	<u>002</u>	May 15, 2014
<u>AB</u>		<u>15MG</u>	<u>A203708</u>	<u>003</u>	May 15, 2014
<u>AB</u>		<u>20MG</u>	<u>A203708</u>	<u>004</u>	May 15, 2014
<u>AB</u>	BARR LABS INC	<u>5MG</u>	<u>A077243</u>	<u>001</u>	Jan 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077243</u>	<u>002</u>	Jan 30, 2012
<u>AB</u>		<u>15MG</u>	<u>A077243</u>	<u>003</u>	Jan 30, 2012
<u>AB</u>		<u>20MG</u>	<u>A077243</u>	<u>004</u>	Jan 30, 2012
<u>AB</u>	CHARTWELL MOLECULAR	<u>5MG</u>	<u>A203456</u>	<u>001</u>	Mar 16, 2016
<u>AB</u>		<u>10MG</u>	<u>A203456</u>	<u>002</u>	Mar 16, 2016
<u>AB</u>		<u>15MG</u>	<u>A203456</u>	<u>003</u>	Mar 16, 2016
<u>AB</u>		<u>20MG</u>	<u>A203456</u>	<u>004</u>	Mar 16, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076534</u>	<u>001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A076534</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A076534</u>	<u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A076534</u>	<u>004</u>	Oct 24, 2011
<u>AB</u>	HEC PHARM	<u>5MG</u>	<u>A208146</u>	<u>001</u>	Jul 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A208146</u>	<u>002</u>	Jul 02, 2018
<u>AB</u>		<u>15MG</u>	<u>A208146</u>	<u>003</u>	Jul 02, 2018
<u>AB</u>		<u>20MG</u>	<u>A208146</u>	<u>004</u>	Jul 02, 2018
<u>AB</u>	HISUN PHARM HANGZHOU	<u>5MG</u>	<u>A206892</u>	<u>001</u>	Dec 31, 2020
<u>AB</u>		<u>10MG</u>	<u>A206892</u>	<u>002</u>	Dec 31, 2020
<u>AB</u>		<u>15MG</u>	<u>A206892</u>	<u>003</u>	Dec 31, 2020
<u>AB</u>		<u>20MG</u>	<u>A206892</u>	<u>004</u>	Dec 31, 2020
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A200221</u>	<u>001</u>	Sep 12, 2012
<u>AB</u>		<u>10MG</u>	<u>A200221</u>	<u>002</u>	Sep 12, 2012
<u>AB</u>		<u>15MG</u>	<u>A200221</u>	<u>003</u>	Sep 12, 2012
<u>AB</u>		<u>20MG</u>	<u>A200221</u>	<u>004</u>	Sep 12, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A203044</u>	<u>001</u>	Feb 20, 2015
<u>AB</u>		<u>10MG</u>	<u>A203044</u>	<u>002</u>	Feb 20, 2015

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET, ORALLY DISINTEGRATING;ORAL

OLANZAPINE

<u>AB</u>		<u>15MG</u>	<u>A203044 003</u>	Feb 20, 2015
<u>AB</u>		<u>20MG</u>	<u>A203044 004</u>	Feb 20, 2015
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A202285 001</u>	May 12, 2014
<u>AB</u>		<u>10MG</u>	<u>A202285 002</u>	May 12, 2014
<u>AB</u>		<u>15MG</u>	<u>A202285 003</u>	May 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A202285 004</u>	May 12, 2014
<u>AB</u>	ORBION PHARMS	<u>5MG</u>	<u>A202937 001</u>	Mar 02, 2015
<u>AB</u>		<u>10MG</u>	<u>A202937 002</u>	Mar 02, 2015
<u>AB</u>		<u>15MG</u>	<u>A202937 003</u>	Mar 02, 2015
<u>AB</u>		<u>20MG</u>	<u>A202937 004</u>	Mar 02, 2015
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A078109 001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A078109 002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A078109 003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A078109 004</u>	Oct 24, 2011
<u>AB</u>	SUN PHARM INDS	<u>5MG</u>	<u>A090881 001</u>	Feb 28, 2012
<u>AB</u>		<u>10MG</u>	<u>A090881 002</u>	Feb 28, 2012
<u>AB</u>		<u>15MG</u>	<u>A090881 003</u>	Feb 28, 2012
<u>AB</u>		<u>20MG</u>	<u>A090881 004</u>	Feb 28, 2012
<u>AB</u>	TORRENT	<u>5MG</u>	<u>A091415 001</u>	Oct 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A091415 002</u>	Oct 25, 2011
<u>AB</u>		<u>15MG</u>	<u>A091415 003</u>	Oct 25, 2011
<u>AB</u>		<u>20MG</u>	<u>A091415 004</u>	Oct 25, 2011
<u>ZYPREXA ZYDIS</u>				
<u>AB</u>	+!	CHEPLAPHARM	<u>5MG</u>	<u>N021086 001</u> Apr 06, 2000
<u>AB</u>	+		<u>10MG</u>	<u>N021086 002</u> Apr 06, 2000
<u>AB</u>	+		<u>15MG</u>	<u>N021086 003</u> Apr 06, 2000
<u>AB</u>	+		<u>20MG</u>	<u>N021086 004</u> Apr 06, 2000

OLANZAPINE PAMOATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ZYPREXA RELPREVV

+	CHEPLAPHARM	EQ 210MG BASE/VIAL	N022173 001	Dec 11, 2009
+		EQ 300MG BASE/VIAL	N022173 002	Dec 11, 2009
+!		EQ 405MG BASE/VIAL	N022173 003	Dec 11, 2009

OLANZAPINE; SAMIDORPHAN L-MALATE

TABLET;ORAL

LYBALVI

+!	ALKERMES INC	5MG;EQ 10MG BASE	N213378 001	May 28, 2021
+		10MG;EQ 10MG BASE	N213378 002	May 28, 2021
+		15MG;EQ 10MG BASE	N213378 003	May 28, 2021
+		20MG;EQ 10MG BASE	N213378 004	May 28, 2021

OLAPARIB

TABLET;ORAL

LYNPARZA

+	ASTRAZENECA	100MG	N208558 001	Aug 17, 2017
+!		150MG	N208558 002	Aug 17, 2017

OLICERIDINE

SOLUTION;INTRAVENOUS

OLINVIK

+!	TREVENA	1MG/ML (1MG/ML)	N210730 001	Oct 30, 2020
+!		2MG/2ML (1MG/ML)	N210730 002	Oct 30, 2020
+!		30MG/30ML (1MG/ML)	N210730 003	Oct 30, 2020

OLIVE OIL; SOYBEAN OIL

EMULSION;INTRAVENOUS

CLINOLIPID 20%

+!	BAXTER HLTHCARE CORP	16% (160GM/1000ML); 4% (40GM/1000ML)	N204508 001	Oct 03, 2013
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OLMESARTAN MEDOXOMIL

TABLET;ORAL

BENICAR

<u>AB</u>	+	COSETTE	<u>5MG</u>	<u>N021286 001</u> Apr 25, 2002
<u>AB</u>	+		<u>20MG</u>	<u>N021286 003</u> Apr 25, 2002
<u>AB</u>	+!		<u>40MG</u>	<u>N021286 004</u> Apr 25, 2002

OLMESARTAN MEDOXOMIL

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A207662 001</u> Apr 24, 2017
<u>AB</u>			<u>20MG</u>	<u>A207662 002</u> Apr 24, 2017
<u>AB</u>			<u>40MG</u>	<u>A207662 003</u> Apr 24, 2017
<u>AB</u>		ALEMBIC	<u>5MG</u>	<u>A203012 001</u> Apr 24, 2017

PRESCRIPTION DRUG PRODUCT LIST

OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL

<u>AB</u>		<u>20MG</u>	<u>A203012</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A203012</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>	ALKEM LABS LTD	<u>5MG</u>	<u>A206763</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A206763</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A206763</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A204798</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A204798</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A204798</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>5MG</u>	<u>A203281</u>	<u>001</u>	May 25, 2017
<u>AB</u>		<u>20MG</u>	<u>A203281</u>	<u>002</u>	May 25, 2017
<u>AB</u>		<u>40MG</u>	<u>A203281</u>	<u>003</u>	May 25, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A204814</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A204814</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A204814</u>	<u>003</u>	Apr 24, 2017
<u>AB</u>	MICRO LABS	<u>5MG</u>	<u>A206372</u>	<u>001</u>	Sep 17, 2019
<u>AB</u>		<u>20MG</u>	<u>A206372</u>	<u>002</u>	Sep 17, 2019
<u>AB</u>		<u>40MG</u>	<u>A206372</u>	<u>003</u>	Sep 17, 2019
<u>AB</u>	MSN	<u>5MG</u>	<u>A217399</u>	<u>001</u>	Jan 18, 2023
<u>AB</u>		<u>20MG</u>	<u>A217399</u>	<u>002</u>	Jan 18, 2023
<u>AB</u>		<u>40MG</u>	<u>A217399</u>	<u>003</u>	Jan 18, 2023
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A206720</u>	<u>001</u>	Dec 02, 2022
<u>AB</u>		<u>20MG</u>	<u>A206720</u>	<u>002</u>	Dec 02, 2022
<u>AB</u>		<u>40MG</u>	<u>A206720</u>	<u>003</u>	Dec 02, 2022
<u>AB</u>	QILU	<u>5MG</u>	<u>A210552</u>	<u>001</u>	Jan 10, 2019
<u>AB</u>		<u>20MG</u>	<u>A210552</u>	<u>002</u>	Jan 10, 2019
<u>AB</u>		<u>40MG</u>	<u>A210552</u>	<u>003</u>	Jan 10, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A208130</u>	<u>001</u>	Jun 29, 2018
<u>AB</u>		<u>20MG</u>	<u>A208130</u>	<u>002</u>	Jun 29, 2018
<u>AB</u>		<u>40MG</u>	<u>A208130</u>	<u>003</u>	Jun 29, 2018
<u>AB</u>	SUNSHINE	<u>5MG</u>	<u>A211049</u>	<u>001</u>	Feb 22, 2019
<u>AB</u>		<u>20MG</u>	<u>A211049</u>	<u>002</u>	Feb 22, 2019
<u>AB</u>		<u>40MG</u>	<u>A211049</u>	<u>003</u>	Feb 22, 2019
<u>AB</u>	UMEDICA	<u>5MG</u>	<u>A207135</u>	<u>001</u>	Jul 18, 2019
<u>AB</u>		<u>20MG</u>	<u>A207135</u>	<u>002</u>	Jul 18, 2019
<u>AB</u>		<u>40MG</u>	<u>A207135</u>	<u>003</u>	Jul 18, 2019
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A205192</u>	<u>001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A205192</u>	<u>002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A205192</u>	<u>003</u>	Apr 24, 2017

OLODATEROL HYDROCHLORIDE

SPRAY, METERED; INHALATION

STRIVERDI RESPIMAT

+	BOEHRINGER INGELHEIM	EQ 0.0025MG BASE/INH	N203108	001	Jul 31, 2014
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OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE

SPRAY, METERED; INHALATION

STIOLTO RESPIMAT

+	BOEHRINGER INGELHEIM	EQ 0.0025MG BASE/INH; EQ 0.0025MG BASE/INH	N206756	001	May 21, 2015
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OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

	SOMERSET THERAPS LLC	EQ 0.1% BASE	A206306	001	Dec 07, 2015
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SPRAY, METERED; NASAL

OLOPATADINE HYDROCHLORIDE

<u>AB</u>	!	AMNEAL	<u>0.665MG/SPRAY</u>	<u>A210901</u>	<u>001</u>	Jan 28, 2020
<u>AB</u>		APOTEX INC	<u>0.665MG/SPRAY</u>	<u>A091572</u>	<u>001</u>	Oct 08, 2014
<u>AB</u>		HIKMA	<u>0.665MG/SPRAY</u>	<u>A213757</u>	<u>001</u>	Aug 19, 2020
<u>AB</u>		PADAGIS ISRAEL	<u>0.665MG/SPRAY</u>	<u>A202853</u>	<u>001</u>	Jan 31, 2017

OLSALAZINE SODIUM

CAPSULE; ORAL

DIPENTUM

+	MYLAN SPCLT VIATRIS	250MG	N019715	001	Jul 31, 1990
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PRESCRIPTION DRUG PRODUCT LIST

OLUTASIDENIB

CAPSULE; ORAL

REZLIDHIA

+! RIGEL PHARMS INC 150MG N215814 001 Dec 01, 2022

OMACETAXINE MEPESUCCINATE

POWDER; SUBCUTANEOUS

SYNRIBO

+! TEVA PHARMS INTL 3.5MG/VIAL N203585 001 Oct 26, 2012

OMADACYCLINE TOSYLATE

POWDER; INTRAVENOUS

NUZYRA

+! PARATEK PHARMS INC EQ 100MG BASE/VIAL N209817 001 Oct 02, 2018

TABLET; ORAL

NUZYRA

+! PARATEK PHARMS INC EQ 150MG BASE N209816 001 Oct 02, 2018

OMAVELOXOLONE

CAPSULE; ORAL

SKYCLARYS

+! REATA PHARMS 50MG N216718 001 Feb 28, 2023

OMEGA-3-ACID ETHYL ESTERS

CAPSULE; ORAL

LOVAZA**AB** +! WOODWARD 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **N021654 001** Nov 10, 2004OMEGA-3-ACID ETHYL ESTERS**AB** AMNEAL PHARMS 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A204940 001** Nov 27, 2015**AB** APOTEX 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A090973 001** Sep 30, 2014**AB** ASCENT PHARMS INC 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A207420 001** Feb 25, 2019**AB** BIONPHARMA 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A206455 001** Aug 07, 2019**AB** CSPEC-NBP PHARM 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A211979 001** May 12, 2020**AB** GLW 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A091028 001** Apr 07, 2014**AB** 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A212504 001** Aug 19, 2020**AB** MANKIND PHARMA 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A215458 001** Nov 15, 2021**AB** PURACAP PHARM LLC 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A210093 001** Jun 15, 2020**AB** SLAYBACK PHARMA LLC 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A211345 001** Dec 07, 2023**AB** SOFGEN PHARMS 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A211355 001** Jul 10, 2019**AB** STRIDES PHARMA 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A203893 001** Sep 19, 2017**AB** SUN PHARM 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS **A210834 001** Jan 09, 2020OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

OMEPRAZOLE**AB** ACTAVIS LABS FL INC 10MG **A075347 001** May 30, 2008**AB** 20MG **A075347 002** May 30, 2008**AB** 40MG **A075347 003** May 30, 2008**AB** APOTEX 10MG **A076048 001** Oct 22, 2007**AB** 20MG **A076048 002** Oct 22, 2007**AB** 40MG **A076048 003** Jan 21, 2009**AB** AUROBINDO PHARMA 10MG **A203270 001** Aug 19, 2015**AB** 20MG **A203270 002** Aug 19, 2015**AB** 40MG **A203270 003** Aug 19, 2015**AB** BRECKENRIDGE 10MG **A203481 001** Jul 03, 2017**AB** 20MG **A203481 002** Jul 03, 2017**AB** 40MG **A203481 003** Jul 03, 2017**AB** DR REDDYS LABS LTD 10MG **A075576 003** Oct 22, 2007**AB** 10MG **A078490 002** Mar 16, 2009**AB** 20MG **A075576 002** Oct 22, 2007**AB** 20MG **A078490 003** Mar 16, 2009**AB** 40MG **A075576 001** Jan 21, 2009**AB** 40MG **A078490 001** Apr 17, 2009**AB** GLENMARK GENERICS 10MG **A091672 001** Oct 31, 2014

PRESCRIPTION DRUG PRODUCT LIST

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

<u>AB</u>		<u>20MG</u>	<u>A091672 002</u>	Oct 31, 2014
<u>AB</u>		<u>40MG</u>	<u>A091672 003</u>	Oct 31, 2014
<u>AB</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A204012 001</u>	Sep 26, 2019
<u>AB</u>		<u>20MG</u>	<u>A204012 002</u>	Sep 26, 2019
<u>AB</u>	IMPAX LABS	<u>10MG</u>	<u>A075785 001</u>	Oct 22, 2007
<u>AB</u>		<u>20MG</u>	<u>A075785 002</u>	Oct 22, 2007
<u>AB</u>		<u>40MG</u>	<u>A075785 003</u>	Jan 21, 2009
<u>AB</u>	LANNETT CO INC	<u>10MG</u>	<u>A075410 001</u>	Nov 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075410 002</u>	Nov 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075410 003</u>	Jan 23, 2009
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A075757 001</u>	Jan 28, 2003
<u>AB</u>	!	<u>20MG</u>	<u>A075757 002</u>	Jan 28, 2003
<u>AB</u>	!	<u>40MG</u>	<u>A076515 001</u>	Jan 21, 2009
<u>AB</u>	TEVA PHARMS USA	<u>20MG</u>	<u>A204661 001</u>	Jun 13, 2017
<u>AB</u>	XIROMED	<u>10MG</u>	<u>A212977 001</u>	Dec 10, 2020
<u>AB</u>		<u>20MG</u>	<u>A212977 002</u>	Dec 10, 2020
<u>AB</u>	ZYDUS PHARMS USA INC	<u>10MG</u>	<u>A091352 001</u>	Nov 19, 2012
<u>AB</u>		<u>20MG</u>	<u>A091352 002</u>	Nov 19, 2012
<u>AB</u>		<u>40MG</u>	<u>A091352 003</u>	Nov 19, 2012

OMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE;ORAL

PRILOSEC

+	COVIS	EQ 2.5MG BASE/PACKET	N022056 001	Mar 20, 2008
+	!	EQ 10MG BASE/PACKET	N022056 002	Mar 20, 2008

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	AJANTA PHARMA LTD	<u>20MG;1.1GM</u>	<u>A204228 001</u>	Jul 15, 2016
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A204228 002</u>	Jul 15, 2016
<u>AB</u>	ANDA REPOSITORY	<u>20MG;1.1GM</u>	<u>A212587 001</u>	Apr 30, 2020
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A212587 002</u>	Apr 30, 2020
<u>AB</u>	AUROLIFE PHARMA LLC	<u>20MG;1.1GM</u>	<u>A204922 001</u>	Aug 19, 2016
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A204922 002</u>	Aug 19, 2016
<u>AB</u>	DR REDDYS	<u>20MG;1.1GM</u>	<u>A204068 001</u>	Jul 15, 2016
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A204068 002</u>	Jul 15, 2016
<u>AB</u>	SCIEGEN PHARMS INC	<u>20MG;1.1GM</u>	<u>A207476 001</u>	Dec 06, 2016
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A207476 002</u>	Dec 06, 2016
<u>AB</u>	ZYDUS PHARMS	<u>20MG;1.1GM</u>	<u>A203290 001</u>	May 25, 2018
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A203290 002</u>	May 25, 2018
	<u>ZEGERID</u>			
<u>AB</u>	+ SALIX	<u>20MG;1.1GM</u>	<u>N021849 001</u>	Feb 27, 2006
<u>AB</u>	+!	<u>40MG;1.1GM</u>	<u>N021849 002</u>	Feb 27, 2006

FOR SUSPENSION;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	AJANTA PHARMA LTD	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>A205545 001</u>	Jul 27, 2016
<u>AB</u>		<u>40MG/PACKET;1.68GM/PACKET</u>	<u>A205545 002</u>	Jul 27, 2016
<u>AB</u>	STRIDES PHARMA	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>A079182 001</u>	Apr 19, 2013
<u>AB</u>		<u>40MG/PACKET;1.68GM/PACKET</u>	<u>A079182 002</u>	Apr 19, 2013
	<u>ZEGERID</u>			
<u>AB</u>	+ SALIX	<u>20MG/PACKET;1.68GM/PACKET</u>	<u>N021636 001</u>	Jun 15, 2004
<u>AB</u>	+!	<u>40MG/PACKET;1.68GM/PACKET</u>	<u>N021636 002</u>	Dec 21, 2004
	KONVOMEF			
	+! AZURITY	2MG/ML; 84MG/ML	N213593 001	Aug 30, 2022

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

<u>AB</u>	AUROBINDO PHARMA	<u>4MG</u>	<u>A090469 001</u>	Apr 12, 2010
<u>AB</u>	!	<u>8MG</u>	<u>A090469 002</u>	Apr 12, 2010
<u>AB</u>	CHARTWELL MOLECULES	<u>4MG</u>	<u>A077406 003</u>	Dec 26, 2006
<u>AB</u>		<u>8MG</u>	<u>A077406 004</u>	Dec 26, 2006
<u>AB</u>	GLENMARK GENERICS	<u>4MG</u>	<u>A078152 001</u>	Jun 27, 2007
<u>AB</u>		<u>8MG</u>	<u>A078152 002</u>	Jun 27, 2007
<u>AB</u>	IPCA LABS LTD	<u>4MG</u>	<u>A209389 001</u>	Oct 30, 2023
<u>AB</u>		<u>8MG</u>	<u>A209389 002</u>	Oct 30, 2023
<u>AB</u>	MYLAN	<u>4MG</u>	<u>A078139 001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A078139 002</u>	Jun 25, 2007
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A078050 001</u>	Aug 13, 2007

PRESCRIPTION DRUG PRODUCT LIST

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

<u>AB</u>		<u>8MG</u>	<u>A078050</u>	<u>002</u>	Aug 13, 2007
<u>AB</u>	SUN PHARM INDS	<u>4MG</u>	<u>A077557</u>	<u>001</u>	Aug 02, 2007
<u>AB</u>		<u>8MG</u>	<u>A077557</u>	<u>002</u>	Aug 02, 2007
<u>AB</u>	SUN PHARM INDS LTD	<u>4MG</u>	<u>A078602</u>	<u>001</u>	Feb 24, 2011
<u>AB</u>		<u>8MG</u>	<u>A078602</u>	<u>002</u>	Feb 24, 2011
!	CHARTWELL MOLECULES	16MG	A077406	001	Dec 26, 2006
!		24MG	A077406	002	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A206846</u>	<u>001</u>	Jul 13, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A076974</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 2MG BASE/ML</u>	<u>A079224</u>	<u>001</u>	Sep 25, 2009
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A090648</u>	<u>001</u>	Jun 15, 2012
<u>AP</u>	HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A076967</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077365</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 2MG BASE/ML</u>	<u>A076781</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077473</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A203711</u>	<u>001</u>	Sep 08, 2014
<u>AP</u>	SANDOZ	<u>EQ 2MG BASE/ML</u>	<u>A077430</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077577</u>	<u>001</u>	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A206845</u>	<u>001</u>	Mar 10, 2016
<u>AP</u>	!	<u>EQ 2MG BASE/ML</u>	<u>A078287</u>	<u>001</u>	Feb 22, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A076972</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A202253</u>	<u>001</u>	Jul 19, 2013
<u>AP</u>	HIKMA	<u>EQ 2MG BASE/ML</u>	<u>A077011</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077541</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077548</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	SANDOZ	<u>EQ 2MG BASE/ML</u>	<u>A077551</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077716</u>	<u>001</u>	Dec 26, 2006

SOLUTION;ORAL

ONDANSETRON HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>EQ 4MG BASE/5ML</u>	<u>A091483</u>	<u>001</u>	Jan 31, 2011
<u>AA</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE/5ML</u>	<u>A078776</u>	<u>001</u>	Nov 28, 2007
<u>AA</u>	CHARTWELL MOLECULAR	<u>EQ 4MG BASE/5ML</u>	<u>A091342</u>	<u>001</u>	Jan 27, 2011
<u>AA</u>	!	<u>EQ 4MG BASE/5ML</u>	<u>A076960</u>	<u>001</u>	Dec 26, 2006
<u>AA</u>	PHARM ASSOC	<u>EQ 4MG BASE/5ML</u>	<u>A078127</u>	<u>001</u>	Jun 25, 2007
<u>AA</u>	TARO	<u>EQ 4MG BASE/5ML</u>	<u>A077009</u>	<u>001</u>	Nov 30, 2007

TABLET;ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 4MG BASE</u>	<u>A077306</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077306</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE</u>	<u>A078539</u>	<u>001</u>	Jul 31, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A078539</u>	<u>002</u>	Jul 31, 2007
<u>AB</u>	CHARTWELL MOLECULES	<u>EQ 4MG BASE</u>	<u>A077303</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077303</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A076183</u>	<u>003</u>	Dec 26, 2006
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076183</u>	<u>002</u>	Dec 26, 2006
<u>AB</u>	GLENMARK GENERICS	<u>EQ 4MG BASE</u>	<u>A077535</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077535</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	IPCA LABS LTD	<u>EQ 4MG BASE</u>	<u>A203761</u>	<u>001</u>	Jan 23, 2014
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A203761</u>	<u>002</u>	Jan 23, 2014
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A077851</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077851</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	SUN PHARM INDS (IN)	<u>EQ 4MG BASE</u>	<u>A077050</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077050</u>	<u>002</u>	Jun 25, 2007
!	CHARTWELL MOLECULES	EQ 24MG BASE	A077303	004	Jun 25, 2007

OPICAPONE

CAPSULE;ORAL

ONGENTYS

+	NEUROCRINE	25MG	N212489	001	Apr 24, 2020
+	!	50MG	N212489	002	Apr 24, 2020

PRESCRIPTION DRUG PRODUCT LIST

ORITAVANCIN DIPHOSPHATE

POWDER; INTRAVENOUS

KIMYRSA

+! MELINTA THERAP EQ 1.2GM BASE/VIAL N214155 001 Mar 12, 2021

ORBACTIV

+! MELINTA THERAP EQ 400MG BASE/VIAL N206334 001 Aug 06, 2014

ORLISTAT

CAPSULE; ORAL

XENICAL

+! CHEPLAPHARM 120MG N020766 001 Apr 23, 1999

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

ORPHENADRINE CITRATEAP HIKMA 30MG/ML A040463 001 Mar 04, 2003AP ! RISING 30MG/ML A040484 001 May 24, 2006AP SAGENT PHARMS 30MG/ML A090585 001 Aug 30, 2011AP WATSON LABS 30MG/ML A084779 001 Mar 15, 1982

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATEAB BAYSHORE PHARMS LLC 100MG A091158 001 Jul 27, 2012AB ! LUPIN 100MG A040284 001 Jun 19, 1998AB RISING 100MG A040249 001 Jan 29, 1999AB SANDOZ 100MG A040327 001 Feb 15, 2000OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATEAB ALEMBIC EQ 30MG BASE A211823 001 Jun 24, 2019AB EQ 45MG BASE A211823 002 Jun 24, 2019AB EQ 75MG BASE A211823 003 Jun 24, 2019AB AMNEAL PHARMS EQ 30MG BASE A209093 001 May 17, 2017AB EQ 45MG BASE A209093 002 May 17, 2017AB EQ 75MG BASE A209093 003 May 17, 2017AB EPIC PHARMA LLC EQ 30MG BASE A215208 001 Oct 01, 2021AB EQ 45MG BASE A215208 002 Oct 01, 2021AB EQ 75MG BASE A215208 003 Oct 01, 2021AB HAINAN POLY EQ 25MG BASE A218009 001 Nov 24, 2023AB HETERO LABS LTD III EQ 30MG BASE A209438 001 Feb 23, 2018AB EQ 45MG BASE A209438 002 Feb 23, 2018AB EQ 75MG BASE A209438 003 Feb 23, 2018AB INVAGEN PHARMS EQ 30MG BASE A217467 001 Oct 03, 2023AB EQ 45MG BASE A217467 002 Oct 03, 2023AB EQ 75MG BASE A217467 003 Oct 03, 2023AB LUPIN EQ 30MG BASE A208348 001 Jan 09, 2018AB EQ 45MG BASE A208348 002 Jan 09, 2018AB EQ 75MG BASE A208348 003 Jan 09, 2018AB MACLEODS PHARMS LTD EQ 30MG BASE A207211 001 Sep 14, 2017AB EQ 45MG BASE A207211 002 Sep 14, 2017AB EQ 75MG BASE A207211 003 Sep 14, 2017AB MSN EQ 30MG BASE A212544 001 May 20, 2020AB EQ 45MG BASE A212544 002 May 20, 2020AB EQ 75MG BASE A212544 003 May 20, 2020AB NATCO EQ 30MG BASE A202595 001 Aug 03, 2016AB EQ 45MG BASE A202595 002 Aug 03, 2016AB EQ 75MG BASE A202595 003 Aug 03, 2016AB STRIDES PHARMA EQ 30MG BASE A209421 001 Jun 08, 2018AB EQ 45MG BASE A209421 002 Jun 08, 2018AB EQ 75MG BASE A209421 003 Jun 08, 2018AB SUNSHINE EQ 30MG BASE A212739 001 Mar 04, 2020AB EQ 45MG BASE A212739 002 Mar 04, 2020AB EQ 75MG BASE A212739 003 Mar 04, 2020AB ZYDUS PHARMS EQ 30MG BASE A208578 001 Feb 24, 2017AB EQ 45MG BASE A208578 002 Feb 24, 2017AB EQ 75MG BASE A208578 003 Feb 24, 2017TAMIFLUAB + ROCHE EQ 30MG BASE N021087 003 Jul 02, 2007AB + EQ 45MG BASE N021087 002 Jul 02, 2007AB +! EQ 75MG BASE N021087 001 Oct 27, 1999

FOR SUSPENSION; ORAL

OSELTAMIVIR PHOSPHATEAB AJANTA PHARMA LTD EQ 6MG BASE/ML A212784 001 May 27, 2020AB ALVOGEN EQ 6MG BASE/ML A208823 001 Oct 31, 2017

PRESCRIPTION DRUG PRODUCT LIST

OSELTAMIVIR PHOSPHATE

FOR SUSPENSION;ORAL

OSELTAMIVIR PHOSPHATE

AB	AMNEAL PHARMS NY	<u>EQ 6MG BASE/ML</u>	<u>A210186 001</u>	Feb 27, 2018
AB	APTAPHARMA INC	<u>EQ 6MG BASE/ML</u>	<u>A212858 001</u>	Aug 30, 2021
AB	EPIC PHARMA LLC	<u>EQ 6MG BASE/ML</u>	<u>A215538 001</u>	Feb 14, 2023
AB	HETERO LABS LTD V	<u>EQ 6MG BASE/ML</u>	<u>A209590 001</u>	Mar 28, 2022
AB	LUPIN	<u>EQ 6MG BASE/ML</u>	<u>A208347 001</u>	Feb 20, 2018
AB	MSN	<u>EQ 6MG BASE/ML</u>	<u>A215313 001</u>	May 02, 2022
AB	STRIDES PHARMA	<u>EQ 6MG BASE/ML</u>	<u>A211894 001</u>	Jan 13, 2022
AB	SUNSHINE	<u>EQ 6MG BASE/ML</u>	<u>A213594 001</u>	Jan 05, 2022
AB	TEVA PHARMS USA	<u>EQ 6MG BASE/ML</u>	<u>A211125 001</u>	Feb 27, 2019
AB	ZYDUS PHARMS	<u>EQ 6MG BASE/ML</u>	<u>A209113 001</u>	Sep 14, 2017

TAMIFLU

AB	+ ! ROCHE	<u>EQ 6MG BASE/ML</u>	<u>N021246 002</u>	Mar 21, 2011
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OSILODROSTAT PHOSPHATE

TABLET;ORAL

ISTURISA

+	RECORDATI RARE	EQ 1MG BASE	N212801 001	Mar 06, 2020
+		EQ 5MG BASE	N212801 002	Mar 06, 2020

OSIMERTINIB MESYLATE

TABLET;ORAL

TAGRISSO

+	ASTRAZENECA	EQ 40MG BASE	N208065 001	Nov 13, 2015
+ !		EQ 80MG BASE	N208065 002	Nov 13, 2015

OSPEMIFENE

TABLET;ORAL

OSPHENA

+ !	DUCHESNAY	60MG	N203505 001	Feb 26, 2013
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OTESECONAZOLE

CAPSULE;ORAL

VIVJOA

+ !	MYCOVIA PHARMS	150MG	N215888 001	Apr 26, 2022
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OXACILLIN SODIUM

INJECTABLE;INJECTION

OXACILLIN SODIUM

AP	EUGIA PHARMA SPECLTS	<u>EQ 1GM BASE/VIAL</u>	<u>A201539 001</u>	Jan 18, 2013
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A201539 002</u>	Jan 18, 2013
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A201538 001</u>	Jan 18, 2013
AP	FRESENIUS KABI USA	<u>EQ 1GM BASE/VIAL</u>	<u>A206198 001</u>	Jul 20, 2020
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A206198 002</u>	Jul 20, 2020
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A206199 001</u>	Jul 27, 2020
AP	! SAGENT PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A091246 001</u>	Mar 30, 2012
AP	!	<u>EQ 2GM BASE/VIAL</u>	<u>A091246 002</u>	Mar 30, 2012
AP	!	<u>EQ 10GM BASE/VIAL</u>	<u>A091245 001</u>	Mar 30, 2012
AP	STERISCIENCE	<u>EQ 2GM BASE/VIAL</u>	<u>A091486 002</u>	Aug 25, 2014
AP	WOCKHARDT BIO AG	<u>EQ 1GM BASE/VIAL</u>	<u>A207147 001</u>	Jul 31, 2017
AP		<u>EQ 2GM BASE/VIAL</u>	<u>A207147 002</u>	Jul 31, 2017
AP		<u>EQ 10GM BASE/VIAL</u>	<u>A207148 001</u>	Nov 24, 2017
	BACTOCILL IN PLASTIC CONTAINER			
+ !	BAXTER HLTHCARE	EQ 20MG BASE/ML	N050640 001	Oct 26, 1989
+ !		EQ 40MG BASE/ML	N050640 002	Oct 26, 1989

OXALIPLATIN

INJECTABLE;INTRAVENOUS

ELOXATIN

AP	+ ! SANOFI AVENTIS US	<u>50MG/10ML (5MG/ML)</u>	<u>N021759 001</u>	Jan 31, 2005
AP	+ !	<u>100MG/20ML (5MG/ML)</u>	<u>N021759 002</u>	Jan 31, 2005

OXALIPLATIN

AP	ACCORD HLTHCARE	<u>50MG/10ML (5MG/ML)</u>	<u>A207474 001</u>	Mar 21, 2017
AP		<u>100MG/20ML (5MG/ML)</u>	<u>A207474 002</u>	Mar 21, 2017
AP	ACTAVIS	<u>50MG/10ML (5MG/ML)</u>	<u>A204880 001</u>	Mar 05, 2018
AP		<u>100MG/20ML (5MG/ML)</u>	<u>A204880 002</u>	Mar 05, 2018
AP	! ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078803 001</u>	Aug 08, 2012
AP	!	<u>100MG/VIAL</u>	<u>A078803 002</u>	Aug 08, 2012
AP	EUGIA PHARMA	<u>50MG/10ML (5MG/ML)</u>	<u>A205529 001</u>	Sep 06, 2017
AP		<u>100MG/20ML (5MG/ML)</u>	<u>A205529 002</u>	Sep 06, 2017
AP	FRESENIUS KABI USA	<u>50MG/10ML (5MG/ML)</u>	<u>A078811 001</u>	Jun 10, 2010
AP		<u>100MG/20ML (5MG/ML)</u>	<u>A078811 002</u>	Jun 10, 2010
AP		<u>50MG/VIAL</u>	<u>A078819 001</u>	Jun 02, 2010

PRESCRIPTION DRUG PRODUCT LIST

OXALIPLATIN

INJECTABLE; INTRAVENOUS

OXALIPLATIN

<u>AP</u>		<u>50MG/10ML (5MG/ML)</u>	<u>A090030 001</u>	Jan 31, 2017
<u>AP</u>		<u>100MG/VIAL</u>	<u>A078819 002</u>	Jun 02, 2010
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A090030 002</u>	Jan 31, 2017
<u>AP</u>	GLAND	<u>50MG/10ML (5MG/ML)</u>	<u>A207325 001</u>	Feb 10, 2017
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A207325 002</u>	Feb 10, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/VIAL</u>	<u>A207385 001</u>	May 23, 2017
<u>AP</u>		<u>100MG/VIAL</u>	<u>A207385 002</u>	May 23, 2017
<u>AP</u>	HENGRUI PHARMA	<u>50MG/10ML (5MG/ML)</u>	<u>A203869 001</u>	Jun 18, 2014
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A203869 002</u>	Jun 18, 2014
<u>AP</u>	HOSPIRA WORLDWIDE	<u>50MG/10ML (5MG/ML)</u>	<u>A078813 001</u>	Aug 07, 2009
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A078813 002</u>	Aug 07, 2009
<u>AP</u>	MYLAN LABS LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A091358 001</u>	Aug 07, 2012
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A091358 002</u>	Aug 07, 2012
<u>AP</u>	NOVAST LABS	<u>50MG/10ML (5MG/ML)</u>	<u>A207562 001</u>	Oct 16, 2018
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A207562 002</u>	Oct 16, 2018
<u>AP</u>	QILU PHARM HAINAN	<u>50MG/10ML (5MG/ML)</u>	<u>A204368 001</u>	Jun 07, 2016
<u>AP</u>		<u>50MG/VIAL</u>	<u>A204616 001</u>	May 11, 2016
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A204368 002</u>	Jun 07, 2016
<u>AP</u>		<u>100MG/VIAL</u>	<u>A204616 002</u>	May 11, 2016
<u>AP</u>	SANDOZ	<u>50MG/10ML (5MG/ML)</u>	<u>A078817 001</u>	Jan 24, 2011
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A078817 002</u>	Jan 24, 2011
<u>AP</u>	+! TEVA PHARMS	<u>50MG/10ML (5MG/ML)</u>	<u>N022160 001</u>	Aug 07, 2009
<u>AP</u>	+!	<u>100MG/20ML (5MG/ML)</u>	<u>N022160 002</u>	Aug 07, 2009
	! QILU PHARM HAINAN	<u>200MG/40ML (5MG/ML)</u>	<u>A204368 003</u>	Jun 07, 2016

OXAPROZIN

CAPSULE; ORAL

COXANTO

+! SOLUBIOMIX

300MG

N217927 001 Oct 20, 2023

TABLET; ORAL

DAYPRO

<u>AB</u>	+! PFIZER	<u>600MG</u>	<u>N018841 004</u>	Oct 29, 1992
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OXAPROZIN

<u>AB</u>	AMNEAL PHARMS CO	<u>600MG</u>	<u>A208633 001</u>	May 04, 2017
<u>AB</u>	CHARTWELL	<u>600MG</u>	<u>A075987 001</u>	Sep 02, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>600MG</u>	<u>A075855 001</u>	Jan 31, 2001
<u>AB</u>	SANDOZ	<u>600MG</u>	<u>A075845 001</u>	Jan 31, 2001
<u>AB</u>	TEVA	<u>600MG</u>	<u>A075849 001</u>	Jul 03, 2002

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072253 002</u>	Apr 14, 1988
<u>AB</u>		<u>15MG</u>	<u>A072253 003</u>	Apr 14, 1988
<u>AB</u>	!	<u>30MG</u>	<u>A072253 001</u>	Apr 14, 1988
<u>AB</u>	EPIC PHARMA LLC	<u>10MG</u>	<u>A071813 001</u>	Apr 19, 1988
<u>AB</u>		<u>15MG</u>	<u>A071756 001</u>	Apr 19, 1988
<u>AB</u>		<u>30MG</u>	<u>A071814 001</u>	Apr 19, 1988
<u>AB</u>	UNITED RES LABS	<u>10MG</u>	<u>A071026 002</u>	Aug 10, 1987
<u>AB</u>		<u>15MG</u>	<u>A071026 003</u>	Aug 10, 1987
<u>AB</u>		<u>30MG</u>	<u>A071026 001</u>	Aug 10, 1987

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

<u>AB</u>	AMNEAL PHARMS	<u>300MG/5ML</u>	<u>A202961 001</u>	Sep 17, 2012
<u>AB</u>	AUCTA	<u>300MG/5ML</u>	<u>A215332 001</u>	Nov 30, 2022
<u>AB</u>	BIONPHARMA	<u>300MG/5ML</u>	<u>A209652 001</u>	Nov 04, 2022
<u>AB</u>	CHARTWELL RX	<u>300MG/5ML</u>	<u>A212428 001</u>	Jun 21, 2021
<u>AB</u>	HETERO LABS LTD III	<u>300MG/5ML</u>	<u>A216749 001</u>	Oct 20, 2023
<u>AB</u>	RUBICON	<u>300MG/5ML</u>	<u>A215726 001</u>	Aug 30, 2022
<u>AB</u>	SUN PHARM INDS LTD	<u>300MG/5ML</u>	<u>A078734 001</u>	Jun 26, 2009

TRILEPTAL

<u>AB</u>	+! NOVARTIS	<u>300MG/5ML</u>	<u>N021285 001</u>	May 25, 2001
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TABLET; ORAL

OXCARBAZEPINE

<u>AB</u>	ANNORA PHARMA	<u>150MG</u>	<u>A215939 001</u>	Jan 11, 2022
<u>AB</u>		<u>300MG</u>	<u>A215939 002</u>	Jan 11, 2022
<u>AB</u>		<u>600MG</u>	<u>A215939 003</u>	Jan 11, 2022
<u>AB</u>	BRECKENRIDGE PHARM	<u>150MG</u>	<u>A078069 001</u>	Jan 11, 2008
<u>AB</u>		<u>300MG</u>	<u>A078069 002</u>	Jan 11, 2008

PRESCRIPTION DRUG PRODUCT LIST

OXCARBAZEPINE

TABLET; ORAL

OXCARBAZEPINE

AB		600MG	A078069 003	Jan 11, 2008
AB	GLENMARK PHARMS LTD	150MG	A077802 001	Oct 09, 2007
AB		300MG	A077802 002	Oct 09, 2007
AB		600MG	A077802 003	Oct 09, 2007
AB	RUBICON	150MG	A077747 001	Apr 09, 2008
AB		300MG	A077747 002	Apr 09, 2008
AB		600MG	A077747 003	Apr 09, 2008
AB	SUN PHARM INDS	150MG	A077794 001	Oct 09, 2007
AB		300MG	A077794 002	Oct 09, 2007
AB		600MG	A077794 003	Oct 09, 2007
AB	TARO	150MG	A077801 001	Nov 15, 2007
AB		300MG	A077801 002	Nov 15, 2007
AB		600MG	A077801 003	Nov 15, 2007
AB	ZYDUS	150MG	A211747 001	Jul 03, 2023
AB		300MG	A211747 002	Jul 03, 2023
AB		600MG	A211747 003	Jul 03, 2023

TRILEPTAL

AB	+	NOVARTIS	150MG	N021014 001	Jan 14, 2000
AB	+		300MG	N021014 002	Jan 14, 2000
AB	+	!	600MG	N021014 003	Jan 14, 2000

TABLET, EXTENDED RELEASE; ORAL

OXTELLAR XR

	+	SUPERNUS PHARMS	150MG	N202810 001	Oct 19, 2012
	+		300MG	N202810 002	Oct 19, 2012
	+	!	600MG	N202810 003	Oct 19, 2012

OXICONAZOLE NITRATE

CREAM; TOPICAL

OXICONAZOLE NITRATE

AB		TARO	EQ 1% BASE	A205076 001	Mar 07, 2016			
AB	+	!	FOUGERA PHARMS	EQ 1% BASE	N019828 001	Dec 30, 1988		
			LOTION; TOPICAL					
			OXISTAT					
			+	!	ANI PHARMS	EQ 1% BASE	N020209 001	Sep 30, 1992

OXYBUTYNYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYTROL

	+	!	ALLERGAN	3.9MG/24HR	N021351 002	Feb 26, 2003
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OXYBUTYNYNIN CHLORIDE

SYRUP; ORAL

OXYBUTYNYNIN CHLORIDE

AA		CHARTWELL RX	5MG/5ML	A075039 001	Jan 29, 1999
AA	!	LANNETT CO INC	5MG/5ML	A074520 001	Mar 29, 1996
AA		PHARM ASSOC	5MG/5ML	A075137 001	Dec 18, 1998

TABLET; ORAL

OXYBUTYNYNIN CHLORIDE

AB		ABHAI LLC	5MG	A209335 001	Dec 22, 2017
AB		BEXIMCO PHARMS USA	5MG	A213550 001	Jul 14, 2022
AB		LEADING	5MG	A212798 001	Apr 06, 2020
AB		NOVAST LABS	5MG	A210611 001	Oct 30, 2019
AB		NOVITIUM PHARMA	5MG	A209823 001	Oct 23, 2017
AB		RISING	5MG	A209025 001	Dec 21, 2017
AB	!	STRIDES PHARMA	5MG	A075079 001	Oct 31, 1997
AB			5MG	A208165 001	Dec 17, 2020
AB		TEVA PHARMS USA	5MG	A071655 001	Nov 14, 1988
AB		TULEX PHARMS INC	5MG	A210125 001	Sep 06, 2018
AB		UPSHER SMITH LABS	5MG	A074625 001	Jul 31, 1996
BX		EYWA	5MG	A211062 001	Feb 06, 2019
	!	RISING	2.5MG	A209025 002	Feb 07, 2023

TABLET, EXTENDED RELEASE; ORAL

OXYBUTYNYNIN CHLORIDE

AB		ACCORD HLTHCARE	5MG	A207138 001	Feb 29, 2016
AB			10MG	A207138 002	Feb 29, 2016
AB	!		15MG	A207138 003	Feb 29, 2016
AB		AJANTA PHARMA LTD	5MG	A211655 001	Feb 28, 2019
AB			10MG	A211655 002	Feb 28, 2019
AB			15MG	A211655 003	Feb 28, 2019
AB		AMNEAL PHARMS	5MG	A204010 001	Nov 23, 2015

PRESCRIPTION DRUG PRODUCT LIST

OXYBUTYNIN CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

OXYBUTYNIN CHLORIDE

<u>AB</u>		<u>10MG</u>	<u>A204010</u>	<u>002</u>	Nov 23, 2015
<u>AB</u>		<u>15MG</u>	<u>A204010</u>	<u>003</u>	Nov 23, 2015
<u>AB</u>	BIONPHARMA	<u>5MG</u>	<u>A210717</u>	<u>001</u>	Dec 17, 2019
<u>AB</u>		<u>10MG</u>	<u>A210717</u>	<u>002</u>	Dec 17, 2019
<u>AB</u>		<u>15MG</u>	<u>A210717</u>	<u>003</u>	Dec 17, 2019
<u>AB</u>	OSMOTICA PHARM US	<u>5MG</u>	<u>A078503</u>	<u>001</u>	Feb 04, 2009
<u>AB</u>		<u>10MG</u>	<u>A078503</u>	<u>002</u>	Feb 04, 2009
<u>AB</u>		<u>15MG</u>	<u>A078503</u>	<u>003</u>	Feb 04, 2009
<u>AB</u>	RUBICON	<u>5MG</u>	<u>A214415</u>	<u>001</u>	Oct 27, 2020
<u>AB</u>		<u>10MG</u>	<u>A214415</u>	<u>002</u>	Oct 27, 2020
<u>AB</u>		<u>15MG</u>	<u>A214415</u>	<u>003</u>	Oct 27, 2020
<u>AB</u>	UNIQUE	<u>5MG</u>	<u>A206121</u>	<u>001</u>	May 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A206121</u>	<u>002</u>	May 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A206121</u>	<u>003</u>	May 27, 2016
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A202332</u>	<u>001</u>	Jun 26, 2017
<u>AB</u>		<u>10MG</u>	<u>A202332</u>	<u>002</u>	Jun 26, 2017
<u>AB</u>		<u>15MG</u>	<u>A202332</u>	<u>003</u>	Jun 26, 2017

OXYCODONE

CAPSULE, EXTENDED RELEASE;ORAL

XTAMPZA ER

+	COLLEGIUM PHARM INC	9MG	N208090	001	Apr 26, 2016
+		13.5MG	N208090	002	Apr 26, 2016
+		18MG	N208090	003	Apr 26, 2016
+		27MG	N208090	004	Apr 26, 2016
+		36MG	N208090	005	Apr 26, 2016

OXYCODONE HYDROCHLORIDE

CAPSULE;ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>5MG</u>	<u>A205177</u>	<u>001</u>	Mar 31, 2016
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A202773</u>	<u>001</u>	Aug 17, 2015
<u>AB</u>	+!	GENUS LIFESCIENCES	<u>N200534</u>	<u>001</u>	Oct 20, 2010
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204752</u>	<u>001</u>	Aug 24, 2015

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

<u>AA</u>	ABHAI LLC	<u>5MG/5ML</u>	<u>A208593</u>	<u>001</u>	Jul 21, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A208593</u>	<u>002</u>	Jul 21, 2017
<u>AA</u>	ALKEM LABS LTD	<u>5MG/5ML</u>	<u>A211748</u>	<u>001</u>	Feb 07, 2019
<u>AA</u>		<u>100MG/5ML</u>	<u>A211749</u>	<u>001</u>	Feb 04, 2019
<u>AA</u>	ANI PHARMS	<u>5MG/5ML</u>	<u>A204979</u>	<u>001</u>	Jun 01, 2015
<u>AA</u>	ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A209021</u>	<u>001</u>	Nov 09, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A209021</u>	<u>002</u>	Nov 09, 2017
<u>AA</u>	+	GENUS LIFESCIENCES	<u>N200535</u>	<u>002</u>	Aug 22, 2013
<u>AA</u>	+		<u>N200535</u>	<u>001</u>	Oct 20, 2010
<u>AA</u>	!	HIKMA	<u>A204037</u>	<u>001</u>	Jul 15, 2013
<u>AA</u>		<u>5MG/5ML</u>	<u>A208817</u>	<u>001</u>	Aug 10, 2017
<u>AA</u>		<u>100MG/5ML</u>	<u>A208795</u>	<u>001</u>	Aug 07, 2017
<u>AA</u>	PHARM ASSOC	<u>5MG/5ML</u>	<u>A206914</u>	<u>001</u>	Feb 01, 2019
<u>AA</u>		<u>100MG/5ML</u>	<u>A206822</u>	<u>001</u>	Aug 15, 2017
<u>AA</u>	QUAGEN	<u>5MG/5ML</u>	<u>A213761</u>	<u>001</u>	Jun 02, 2021
<u>AA</u>		<u>100MG/5ML</u>	<u>A213761</u>	<u>002</u>	Jun 02, 2021
<u>AA</u>	SPECGX LLC	<u>5MG/5ML</u>	<u>A210758</u>	<u>001</u>	Apr 30, 2018
<u>AA</u>		<u>100MG/5ML</u>	<u>A210758</u>	<u>002</u>	Apr 30, 2018
<u>AA</u>	WES PHARMA INC	<u>5MG/5ML</u>	<u>A207511</u>	<u>001</u>	Nov 23, 2016
<u>AA</u>		<u>100MG/5ML</u>	<u>A209897</u>	<u>001</u>	Sep 06, 2017

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ALVOGEN	<u>5MG</u>	<u>A202116</u>	<u>001</u>	Dec 30, 2011
<u>AB</u>		<u>15MG</u>	<u>A202116</u>	<u>002</u>	Dec 30, 2011
<u>AB</u>		<u>30MG</u>	<u>A202116</u>	<u>003</u>	Dec 30, 2011
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A203638</u>	<u>001</u>	Jun 03, 2014
<u>AB</u>		<u>10MG</u>	<u>A203638</u>	<u>002</u>	Jun 03, 2014
<u>AB</u>		<u>15MG</u>	<u>A203638</u>	<u>003</u>	Jun 03, 2014
<u>AB</u>		<u>20MG</u>	<u>A203638</u>	<u>004</u>	Jun 03, 2014
<u>AB</u>		<u>30MG</u>	<u>A203638</u>	<u>005</u>	Jun 03, 2014
<u>AB</u>	ASCENT PHARMS INC	<u>15MG</u>	<u>A207418</u>	<u>001</u>	Aug 07, 2017
<u>AB</u>		<u>30MG</u>	<u>A207418</u>	<u>002</u>	Aug 07, 2017
<u>AB</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A202160</u>	<u>001</u>	Nov 19, 2012
<u>AB</u>		<u>15MG</u>	<u>A202160</u>	<u>002</u>	Nov 19, 2012

PRESCRIPTION DRUG PRODUCT LIST

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>		<u>30MG</u>	<u>A202160</u>	<u>003</u>	Nov 19, 2012
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A091393</u>	<u>001</u>	Aug 31, 2009
<u>AB</u>	!	<u>10MG</u>	<u>A091393</u>	<u>002</u>	Aug 31, 2009
<u>AB</u>		<u>15MG</u>	<u>A091393</u>	<u>003</u>	Aug 31, 2009
<u>AB</u>		<u>20MG</u>	<u>A091393</u>	<u>004</u>	Aug 31, 2009
<u>AB</u>		<u>30MG</u>	<u>A091393</u>	<u>005</u>	Aug 31, 2009
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A090895</u>	<u>001</u>	Aug 24, 2009
<u>AB</u>		<u>5MG</u>	<u>A202662</u>	<u>001</u>	Sep 22, 2015
<u>AB</u>		<u>10MG</u>	<u>A202662</u>	<u>002</u>	Sep 22, 2015
<u>AB</u>		<u>15MG</u>	<u>A090895</u>	<u>002</u>	Aug 24, 2009
<u>AB</u>		<u>15MG</u>	<u>A202662</u>	<u>003</u>	Sep 22, 2015
<u>AB</u>		<u>20MG</u>	<u>A202662</u>	<u>005</u>	Apr 27, 2017
<u>AB</u>		<u>30MG</u>	<u>A090895</u>	<u>003</u>	Aug 24, 2009
<u>AB</u>		<u>30MG</u>	<u>A202662</u>	<u>004</u>	Sep 22, 2015
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204021</u>	<u>001</u>	Jun 12, 2017
<u>AB</u>		<u>10MG</u>	<u>A204021</u>	<u>002</u>	Jun 12, 2017
<u>AB</u>		<u>15MG</u>	<u>A204021</u>	<u>003</u>	Jun 12, 2017
<u>AB</u>		<u>20MG</u>	<u>A204021</u>	<u>004</u>	Jun 12, 2017
<u>AB</u>		<u>30MG</u>	<u>A204021</u>	<u>005</u>	Jun 12, 2017
<u>AB</u>	NUVO PHARM	<u>5MG</u>	<u>A207119</u>	<u>001</u>	Apr 12, 2016
<u>AB</u>		<u>10MG</u>	<u>A207119</u>	<u>002</u>	Apr 12, 2016
<u>AB</u>		<u>15MG</u>	<u>A207119</u>	<u>003</u>	Apr 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A207119</u>	<u>004</u>	Apr 12, 2016
<u>AB</u>		<u>30MG</u>	<u>A207119</u>	<u>005</u>	Apr 12, 2016
<u>AB</u>	RHODES PHARMS	<u>5MG</u>	<u>A091490</u>	<u>001</u>	Mar 09, 2011
<u>AB</u>		<u>10MG</u>	<u>A091490</u>	<u>002</u>	Mar 09, 2011
<u>AB</u>		<u>15MG</u>	<u>A091490</u>	<u>003</u>	Mar 09, 2011
<u>AB</u>		<u>20MG</u>	<u>A091490</u>	<u>004</u>	Mar 09, 2011
<u>AB</u>		<u>30MG</u>	<u>A091490</u>	<u>005</u>	Mar 09, 2011
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A076758</u>	<u>003</u>	Mar 19, 2007
<u>AB</u>		<u>10MG</u>	<u>A076758</u>	<u>004</u>	Oct 12, 2021
<u>AB</u>		<u>15MG</u>	<u>A076758</u>	<u>001</u>	Jun 30, 2004
<u>AB</u>		<u>20MG</u>	<u>A076758</u>	<u>005</u>	Oct 12, 2021
<u>AB</u>		<u>30MG</u>	<u>A076758</u>	<u>002</u>	Jun 30, 2004
<u>AB</u>	SUN PHARM INDS INC	<u>5MG</u>	<u>A090659</u>	<u>001</u>	Apr 10, 2009
<u>AB</u>		<u>10MG</u>	<u>A090659</u>	<u>005</u>	Nov 06, 2012
<u>AB</u>		<u>15MG</u>	<u>A090659</u>	<u>002</u>	Apr 10, 2009
<u>AB</u>		<u>20MG</u>	<u>A090659</u>	<u>004</u>	Nov 06, 2012
<u>AB</u>		<u>30MG</u>	<u>A090659</u>	<u>003</u>	Apr 10, 2009
<u>ROXICODONE</u>					
<u>AB</u>	+ SPECGX LLC	<u>5MG</u>	<u>N021011</u>	<u>003</u>	May 15, 2009
<u>AB</u>	+!	<u>15MG</u>	<u>N021011</u>	<u>001</u>	Aug 31, 2000
<u>AB</u>	+	<u>30MG</u>	<u>N021011</u>	<u>002</u>	Aug 31, 2000

ROXYBOND

PROTEGA PHARMS	5MG	N209777	001	Apr 20, 2017
	15MG	N209777	002	Apr 20, 2017
	30MG	N209777	003	Apr 20, 2017

TABLET, EXTENDED RELEASE; ORAL

OXYCONTIN

+	PURDUE PHARMA LP	10MG	N022272	001	Apr 05, 2010
+		15MG	N022272	002	Apr 05, 2010
+		20MG	N022272	003	Apr 05, 2010
+		30MG	N022272	004	Apr 05, 2010
+	!	40MG	N022272	005	Apr 05, 2010
+		60MG	N022272	006	Apr 05, 2010
+		80MG	N022272	007	Apr 05, 2010

OXYMETAZOLINE HYDROCHLORIDE

CREAM; TOPICAL

RHOFADE

+	MAYNE PHARMA	1%	N208552	001	Jan 18, 2017
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SOLUTION/DROPS; OPHTHALMIC

UPNEEQ

+	RVL PHARMS	0.1%	N212520	001	Jul 08, 2020
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PRESCRIPTION DRUG PRODUCT LIST

OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE

SPRAY, METERED;NASAL

KOVANAZE

+! ST RENATUS

0.1MG/SPRAY;6MG/SPRAY

N208032 001 Jun 29, 2016

OXYMORPHONE HYDROCHLORIDE

TABLET;ORAL

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>	ASCENT PHARMS INC	<u>5MG</u>	<u>A210175 001</u>	Feb 02, 2018
<u>AB</u>		<u>10MG</u>	<u>A210175 002</u>	Feb 02, 2018
<u>AB</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A204459 001</u>	Apr 26, 2016
<u>AB</u>		<u>10MG</u>	<u>A204459 002</u>	Apr 26, 2016
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A203601 001</u>	Jan 30, 2013
<u>AB</u>	!	<u>10MG</u>	<u>A203601 002</u>	Jan 30, 2013
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A201187 001</u>	Dec 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A201187 002</u>	Dec 15, 2014
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A090964 001</u>	Sep 27, 2010
<u>AB</u>		<u>10MG</u>	<u>A090964 002</u>	Sep 27, 2010
<u>AB</u>	TEVA	<u>5MG</u>	<u>A091443 002</u>	Feb 15, 2011
<u>AB</u>		<u>10MG</u>	<u>A091443 001</u>	Feb 15, 2011

TABLET, EXTENDED RELEASE;ORAL

OXYMORPHONE HYDROCHLORIDE

IMPAX LABS

5MG

A079087 001 Jun 14, 2010

7.5MG

A079087 002 Dec 21, 2010

10MG

A079087 003 Jun 14, 2010

15MG

A079087 004 Dec 21, 2010

20MG

A079087 005 Jun 14, 2010

30MG

A079087 006 Jul 22, 2010

!

40MG

A079087 007 Jun 14, 2010

OXYTOCIN

INJECTABLE;INJECTION

OXYTOCIN

<u>AP</u>	+!	FRESENIUS KABI USA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018248 001</u>	
<u>AP</u>	+!		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018248 002</u>	
<u>AP</u>	+!	HIKMA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018243 001</u>	
<u>AP</u>	+!		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018243 002</u>	Jan 10, 2007
<u>AP</u>		HIKMA FARMACEUTICA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A200219 001</u>	Feb 13, 2013
<u>AP</u>		SAGENT PHARMS INC	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A091676 001</u>	Jul 13, 2018
<u>AP</u>			<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>A091676 002</u>	Jul 13, 2018

PITOCIN

<u>AP</u>	+!	PAR STERILE PRODUCTS	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018261 001</u>	
<u>AP</u>	+		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018261 002</u>	Jul 27, 2007
		OXYTOCIN			
	+!	FRESENIUS KABI USA	<u>300USP UNITS/30ML (10USP UNITS/ML)</u>	<u>N018248 003</u>	Jul 27, 2007
		PITOCIN			
	+	PAR STERILE PRODUCTS	<u>500USP UNITS/50ML (10USP UNITS/ML)</u>	<u>N018261 003</u>	Sep 05, 2012

OZANIMOD HYDROCHLORIDE

CAPSULE;ORAL

ZEPOSIA

+ CELGENE INTL

EQ 0.23MG BASE

N209899 001 Mar 25, 2020

+

EQ 0.46MG BASE

N209899 002 Mar 25, 2020

+!

EQ 0.92MG BASE

N209899 003 Mar 25, 2020

OZENOXACIN

CREAM;TOPICAL

XEPI

+! FERRER

1%

N208945 001 Dec 11, 2017

INTERNACIONAL

PACLITAXEL

INJECTABLE;INJECTION

PACLITAXEL

<u>AP</u>		ACCORD HLTHCARE	<u>6MG/ML</u>	<u>A205720 001</u>	Aug 17, 2018
<u>AP</u>		ACTAVIS TOTOWA	<u>6MG/ML</u>	<u>A090130 001</u>	Dec 09, 2009
<u>AP</u>		ALEMBIC	<u>6MG/ML</u>	<u>A216874 001</u>	Oct 20, 2022
<u>AP</u>		FRESENIUS KABI USA	<u>6MG/ML</u>	<u>A077574 001</u>	Nov 27, 2006
<u>AP</u>		GLAND PHARMA LTD	<u>6MG/ML</u>	<u>A207326 001</u>	Aug 23, 2016
<u>AP</u>		HIKMA	<u>6MG/ML</u>	<u>A075190 001</u>	Jan 28, 2002
<u>AP</u>	!	HOSPIRA	<u>6MG/ML</u>	<u>A076131 001</u>	May 08, 2002
<u>AP</u>		MSN	<u>6MG/ML</u>	<u>A213434 001</u>	Aug 24, 2020
<u>AP</u>		TEVA PHARMS	<u>6MG/ML</u>	<u>A075184 001</u>	Jan 25, 2002

PRESCRIPTION DRUG PRODUCT LISTPACLITAXEL

POWDER; INTRAVENOUS

ABRAXANE

+! BRISTOL-MYERS 100MG/VIAL N021660 001 Jan 07, 2005

PACLITAXEL

+! AM REGENT 100MG/VIAL N211875 001 Jul 27, 2022

PACRITINIB CITRATE

CAPSULE; ORAL

VONJO

+! CTI BIOPHARMA CORP EQ 100MG BASE N208712 001 Feb 28, 2022

PAFOLACIANINE SODIUM

SOLUTION; INTRAVENOUS

CYTALUX

+! ON TARGET LABS EQ 3.2MG BASE/1.6ML (EQ 2MG BASE/ML) N214907 001 Nov 29, 2021

PALBOCICLIB

CAPSULE; ORAL

IBRANCE

+ PFIZER 75MG N207103 001 Feb 03, 2015

+ 100MG N207103 002 Feb 03, 2015

+! 125MG N207103 003 Feb 03, 2015

TABLET; ORAL

IBRANCEAB + PFIZER 75MG N212436 001 Nov 01, 2019AB + 100MG N212436 002 Nov 01, 2019AB +! 125MG N212436 003 Nov 01, 2019PALBOCICLIBAB SYNTHON PHARMS INC 75MG A215570 001 Aug 28, 2023AB 100MG A215570 002 Aug 28, 2023AB 125MG A215570 003 Aug 28, 2023PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGAAB + JANSSEN PHARMS 1.5MG N021999 006 Aug 26, 2008AB + 3MG N021999 001 Dec 19, 2006AB +! 6MG N021999 002 Dec 19, 2006AB + 9MG N021999 003 Dec 19, 2006PALIPERIDONEAB ACTAVIS LABS FL INC 1.5MG A202645 001 Aug 03, 2015AB 3MG A202645 002 Aug 03, 2015AB 6MG A202645 003 Aug 03, 2015AB 9MG A202645 004 Aug 03, 2015AB AMNEAL PHARMS 1.5MG A204707 001 Sep 23, 2019AB 3MG A204707 002 Sep 23, 2019AB 6MG A204707 003 Sep 23, 2019AB 9MG A204707 004 Sep 23, 2019AB ASCENT PHARMS INC 1.5MG A216174 001 Aug 23, 2023AB 3MG A216174 002 Aug 23, 2023AB 6MG A216174 003 Aug 23, 2023AB 9MG A216174 004 Aug 23, 2023AB CSPC OUYI 1.5MG A212807 001 Oct 29, 2020AB 3MG A212807 002 Oct 29, 2020AB 6MG A212807 003 Oct 29, 2020AB 9MG A212807 004 Oct 29, 2020AB INVENTIA 1.5MG A204452 001 Jun 12, 2019AB 3MG A204452 002 Jun 12, 2019AB 6MG A204452 003 Jun 12, 2019AB 9MG A204452 004 Jun 12, 2019AB LUPIN LTD 1.5MG A208643 001 Jun 29, 2022AB 3MG A208643 002 Jun 29, 2022AB 6MG A208643 003 Jun 29, 2022AB 9MG A208643 004 Jun 29, 2022AB RK PHARMA 1.5MG A203802 001 Sep 24, 2015AB 3MG A203802 002 Sep 24, 2015AB 6MG A203802 003 Sep 24, 2015AB 9MG A203802 004 Sep 24, 2015AB SUN PHARM 1.5MG A205618 001 Apr 06, 2018AB 3MG A205618 002 Apr 06, 2018AB 6MG A205618 003 Apr 06, 2018AB 9MG A205618 004 Apr 06, 2018

PRESCRIPTION DRUG PRODUCT LIST

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

INVEGA HAFYERA

+	JANSSEN PHARMS	1.092GM/3.5ML (312MG/ML)	N207946	005	Aug 30, 2021
+		1.560GM/5ML (312MG/ML)	N207946	006	Aug 30, 2021

INVEGA SUSTENNA

+	JANSSEN PHARMS	39MG/0.25ML (39MG/0.25ML)	N022264	001	Jul 31, 2009
+		78MG/0.5ML (78MG/0.5ML)	N022264	002	Jul 31, 2009
+		117MG/0.75ML (117MG/0.75ML)	N022264	003	Jul 31, 2009
+		156MG/ML (156MG/ML)	N022264	004	Jul 31, 2009
+		234MG/1.5ML (156MG/ML)	N022264	005	Jul 31, 2009

INVEGA TRINZA

+	JANSSEN PHARMS	273MG/0.875ML (273MG/0.875ML)	N207946	001	May 18, 2015
+		410MG/1.315ML (311.79MG/ML)	N207946	002	May 18, 2015
+		546MG/1.75ML (312MG/ML)	N207946	003	May 18, 2015
+		819MG/2.625ML (312MG/ML)	N207946	004	May 18, 2015

PALONOSETRON HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

AP	AVET LIFESCIENCES	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A202951	002	Jun 29, 2021
AP	BAXTER HLTHCARE CORP	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A206916	001	Nov 12, 2021
AP	CHARTWELL RX	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A209287	001	Sep 19, 2018
AP	! DR REDDYS	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A201533	002	Apr 21, 2016
AP	EUGIA PHARMA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A204702	001	Nov 06, 2018
AP	FRESENIUS KABI USA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A206801	001	Sep 19, 2018
AP		EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A206802	001	Sep 19, 2018
AP	HOSPIRA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A207005	001	Sep 19, 2018
AP	MYLAN INSTITUTIONAL	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A206416	001	Sep 19, 2018
AP	NANJING KING-FRIEND	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A215861	001	Aug 14, 2023
AP	QILU PHARM HAINAN	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A205648	001	Sep 19, 2018
AP	SAGENT PHARMS INC	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A204289	001	Sep 19, 2018
AP		EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A205870	001	Sep 19, 2018
AP	SANDOZ	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A202521	001	Oct 13, 2015
AP	TEVA PHARMS USA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	A090713	001	Mar 23, 2018
	! AVET LIFESCIENCES	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	A202951	001	Jun 29, 2021

PALOVAROTENE

CAPSULE; ORAL

SOHONOS

+	IPSEN	1MG	N215559	001	Aug 16, 2023
+		1.5MG	N215559	002	Aug 16, 2023
+		2.5MG	N215559	003	Aug 16, 2023
+		5MG	N215559	004	Aug 16, 2023
+		10MG	N215559	005	Aug 16, 2023

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

AP	AREVA PHARMS	30MG/VIAL	A077433	001	Nov 26, 2008
AP		90MG/VIAL	A077433	003	Nov 26, 2008
AP	DR REDDYS	60MG/10ML (6MG/ML)	A078156	002	Aug 19, 2008
AP	HIKMA	30MG/VIAL	A075290	001	Apr 30, 2001
AP		90MG/VIAL	A075290	003	Apr 30, 2001
AP	! HOSPIRA	60MG/10ML (6MG/ML)	A075841	002	Jun 27, 2002
AP1	! HIKMA	30MG/10ML (3MG/ML)	N021113	001	Mar 04, 2002
AP1	! HIKMA	90MG/10ML (9MG/ML)	N021113	002	Mar 04, 2002
AP1	SAGENT PHARMS INC	30MG/10ML (3MG/ML)	A078373	001	Dec 23, 2008
AP1		90MG/10ML (9MG/ML)	A078373	002	Dec 23, 2008
AP2	DR REDDYS	30MG/10ML (3MG/ML)	A078156	001	Aug 19, 2008
AP2		90MG/10ML (9MG/ML)	A078156	003	Aug 19, 2008
AP2	HOSPIRA	30MG/10ML (3MG/ML)	A075841	001	Jun 27, 2002
AP2		90MG/10ML (9MG/ML)	A075841	003	Jun 27, 2002
AP2	MYLAN LABS LTD	30MG/10ML (3MG/ML)	A078520	001	Oct 31, 2008
AP2		90MG/10ML (9MG/ML)	A078520	002	Oct 31, 2008
	AREVA PHARMS	60MG/VIAL	A077433	002	Nov 26, 2008

PRESCRIPTION DRUG PRODUCT LIST

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

AP	!	DR REDDYS	<u>1MG/ML</u>	<u>A072759</u>	<u>001</u>	Jul 31, 1990
AP		HOSPIRA	<u>1MG/ML</u>	<u>A072320</u>	<u>001</u>	Jan 19, 1989
	!	DR REDDYS	2MG/ML	A072760	001	Jul 31, 1990

PANTOPRAZOLE SODIUM

FOR SUSPENSION, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

AB		AJANTA PHARMA LTD	<u>EQ 40MG BASE</u>	<u>A217416</u>	<u>001</u>	Feb 09, 2023
AB		ANNORA PHARMA	<u>EQ 40MG BASE</u>	<u>A216139</u>	<u>001</u>	Oct 27, 2023
AB		DEXCEL	<u>EQ 40MG BASE</u>	<u>A216247</u>	<u>001</u>	Jun 16, 2023
AB		SUN PHARM	<u>EQ 40MG BASE</u>	<u>A213725</u>	<u>001</u>	Jun 30, 2020

PROTONIX

AB	+	WYETH PHARMS	<u>EQ 40MG BASE</u>	<u>N022020</u>	<u>001</u>	Nov 14, 2007
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INJECTABLE; INTRAVENOUS

PANTOPRAZOLE SODIUM

AP		ACIC PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>A209524</u>	<u>001</u>	Aug 30, 2021
AP		AKORN	<u>EQ 40MG BASE/VIAL</u>	<u>A079197</u>	<u>001</u>	Nov 08, 2012
AP		ASPIRO	<u>EQ 40MG BASE/VIAL</u>	<u>A213778</u>	<u>001</u>	May 18, 2022
AP		BE PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>A216171</u>	<u>001</u>	May 18, 2022
AP		EUGIA PHARMA	<u>EQ 40MG BASE/VIAL</u>	<u>A205675</u>	<u>001</u>	Mar 30, 2016
AP		GLAND PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A204400</u>	<u>001</u>	May 18, 2022
AP		KNACK	<u>EQ 40MG BASE/VIAL</u>	<u>A214680</u>	<u>001</u>	May 19, 2022
AP		MEITHEAL	<u>EQ 40MG BASE/VIAL</u>	<u>A215860</u>	<u>001</u>	Aug 29, 2022
AP		SANDOZ	<u>EQ 40MG BASE/VIAL</u>	<u>A090296</u>	<u>001</u>	Jul 14, 2015
AP		SUN PHARM	<u>EQ 40MG BASE/VIAL</u>	<u>A077674</u>	<u>001</u>	Aug 19, 2019

PROTONIX IV

AP	+	WYETH PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>N020988</u>	<u>001</u>	Mar 22, 2001
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POWDER; INTRAVENOUS

PANTOPRAZOLE SODIUM

	+	HIKMA	EQ 40MG BASE/VIAL	N209463	001	Jun 30, 2017
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TABLET, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

AB		ACTAVIS TOTOWA	<u>EQ 20MG BASE</u>	<u>A090797</u>	<u>001</u>	Feb 07, 2011
AB			<u>EQ 40MG BASE</u>	<u>A090797</u>	<u>002</u>	Feb 07, 2011
AB		AMNEAL PHARMS	<u>EQ 20MG BASE</u>	<u>A205119</u>	<u>001</u>	Jan 26, 2016
AB			<u>EQ 40MG BASE</u>	<u>A205119</u>	<u>002</u>	Jan 26, 2016
AB		AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A202038</u>	<u>001</u>	Sep 28, 2012
AB			<u>EQ 40MG BASE</u>	<u>A202038</u>	<u>002</u>	Sep 28, 2012
AB		DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A077619</u>	<u>001</u>	Jan 19, 2011
AB			<u>EQ 40MG BASE</u>	<u>A077619</u>	<u>002</u>	Jan 19, 2011
AB		GRANULES	<u>EQ 20MG BASE</u>	<u>A217282</u>	<u>001</u>	Dec 11, 2023
AB			<u>EQ 40MG BASE</u>	<u>A217282</u>	<u>002</u>	Dec 11, 2023
AB		HETERO LABS LTD V	<u>EQ 20MG BASE</u>	<u>A202882</u>	<u>001</u>	Sep 10, 2014
AB			<u>EQ 40MG BASE</u>	<u>A202882</u>	<u>002</u>	Sep 10, 2014
AB		INGENUS PHARMS LLC	<u>EQ 40MG BASE</u>	<u>A211368</u>	<u>001</u>	Mar 01, 2019
AB		LANNETT CO INC	<u>EQ 20MG BASE</u>	<u>A078281</u>	<u>001</u>	Jan 20, 2011
AB			<u>EQ 40MG BASE</u>	<u>A078281</u>	<u>002</u>	Jan 20, 2011
AB		MANKIND PHARMA	<u>EQ 40MG BASE</u>	<u>A215880</u>	<u>001</u>	Jul 26, 2022
AB		MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A090970</u>	<u>001</u>	Jan 19, 2011
AB			<u>EQ 40MG BASE</u>	<u>A090970</u>	<u>002</u>	Jan 19, 2011
AB		ORBION PHARMS	<u>EQ 20MG BASE</u>	<u>A202052</u>	<u>001</u>	Dec 02, 2014
AB			<u>EQ 40MG BASE</u>	<u>A202052</u>	<u>002</u>	Dec 02, 2014
AB		RUBICON	<u>EQ 20MG BASE</u>	<u>A090807</u>	<u>001</u>	May 02, 2012
AB			<u>EQ 40MG BASE</u>	<u>A090807</u>	<u>002</u>	May 02, 2012
AB		TORRENT PHARMS	<u>EQ 20MG BASE</u>	<u>A090074</u>	<u>001</u>	Jan 19, 2011
AB			<u>EQ 40MG BASE</u>	<u>A090074</u>	<u>002</u>	Jan 19, 2011
AB		WOCKHARDT BIO AG	<u>EQ 20MG BASE</u>	<u>A091231</u>	<u>001</u>	Jan 19, 2011
AB			<u>EQ 40MG BASE</u>	<u>A091231</u>	<u>002</u>	Jan 19, 2011

PROTONIX

AB	+	WYETH PHARMS	<u>EQ 20MG BASE</u>	<u>N020987</u>	<u>002</u>	Jun 12, 2001
AB	+		<u>EQ 40MG BASE</u>	<u>N020987</u>	<u>001</u>	Feb 02, 2000

PARICALCITOL

CAPSULE; ORAL

PARICALCITOL

AB		AMNEAL PHARMS	<u>1MCG</u>	<u>A204327</u>	<u>001</u>	Jan 13, 2016
AB			<u>2MCG</u>	<u>A204327</u>	<u>002</u>	Jan 13, 2016
AB			<u>4MCG</u>	<u>A204327</u>	<u>003</u>	Jan 13, 2016
AB		AUROBINDO PHARMA LTD	<u>1MCG</u>	<u>A207672</u>	<u>001</u>	Jan 14, 2016

PRESCRIPTION DRUG PRODUCT LIST

PARICALCITOL

CAPSULE; ORAL

PARICALCITOL

<u>AB</u>		<u>2MCG</u>	<u>A207672 002</u>	Jan 14, 2016
<u>AB</u>		<u>4MCG</u>	<u>A207672 003</u>	Jan 14, 2016
<u>AB</u>	BIONPHARMA	<u>1MCG</u>	<u>A202539 001</u>	Mar 27, 2014
<u>AB</u>		<u>2MCG</u>	<u>A202539 002</u>	Mar 27, 2014
<u>AB</u>		<u>4MCG</u>	<u>A202539 003</u>	Mar 27, 2014
<u>AB</u>	DR REDDYS	<u>1MCG</u>	<u>A091412 001</u>	Jun 24, 2014
<u>AB</u>		<u>2MCG</u>	<u>A091412 002</u>	Jun 24, 2014
<u>AB</u>		<u>4MCG</u>	<u>A091412 003</u>	Jun 24, 2014
<u>AB</u>	MARKSANS PHARMA	<u>1MCG</u>	<u>A204948 001</u>	Oct 07, 2016
<u>AB</u>		<u>2MCG</u>	<u>A204948 002</u>	Oct 07, 2016
<u>AB</u>		<u>4MCG</u>	<u>A204948 003</u>	Oct 07, 2016
<u>AB</u>	RISING	<u>1MCG</u>	<u>A202124 001</u>	Jun 24, 2014
<u>AB</u>		<u>2MCG</u>	<u>A202124 002</u>	Jun 24, 2014
<u>AB</u>		<u>4MCG</u>	<u>A202124 003</u>	Jun 24, 2014
<u>AB</u>	TEVA PHARMS USA	<u>1MCG</u>	<u>A090829 001</u>	Sep 27, 2013
<u>AB</u>		<u>2MCG</u>	<u>A090829 002</u>	Sep 27, 2013
<u>AB</u>	!	<u>4MCG</u>	<u>A090829 003</u>	Sep 27, 2013

ZEMPLAR

<u>AB</u>	+ ABBVIE	<u>1MCG</u>	<u>N021606 001</u>	May 26, 2005
<u>AB</u>	+	<u>2MCG</u>	<u>N021606 002</u>	May 26, 2005

SOLUTION; INTRAVENOUS

PARICALCITOL

<u>AP</u>	ACCORD HLTHCARE	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N207174 001</u>	Feb 04, 2016
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N207174 002</u>	Feb 04, 2016
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N207174 003</u>	Feb 04, 2016
<u>AP</u>	AMNEAL PHARMS CO	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A206699 001</u>	Mar 09, 2017
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A206699 002</u>	Mar 09, 2017
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A206699 003</u>	Mar 09, 2017
<u>AP</u>	DR REDDYS	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A204910 001</u>	Aug 17, 2016
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A204910 002</u>	Aug 17, 2016
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A204910 003</u>	Aug 17, 2016
<u>AP</u>	EUGIA PHARMA	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A205982 001</u>	Oct 09, 2018
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A205982 002</u>	Oct 09, 2018
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A205982 003</u>	Oct 09, 2018
<u>AP</u>	HIKMA PHARMS	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N205917 001</u>	Nov 18, 2014
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N205917 002</u>	Nov 18, 2014
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N205917 003</u>	Nov 18, 2014
<u>AP</u>	HOSPIRA	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N201657 001</u>	Oct 21, 2014
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N201657 002</u>	Oct 21, 2014
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N201657 003</u>	Oct 21, 2014
<u>AP</u>	RISING	<u>0.005MG/ML (0.005MG/ML)</u>	<u>A203897 002</u>	Nov 02, 2017
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A203897 003</u>	Nov 02, 2017
<u>AP</u>	SANDOZ	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A091108 001</u>	Jul 27, 2011
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A091108 002</u>	Jul 27, 2011
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A091108 003</u>	Jul 27, 2011

ZEMPLAR

<u>AP</u>	+! ABBVIE	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N020819 002</u>	Feb 01, 2000
<u>AP</u>	+!	<u>0.005MG/ML (0.005MG/ML)</u>	<u>N020819 001</u>	Apr 17, 1998
<u>AP</u>	+!	<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N020819 003</u>	Feb 01, 2000

PAROMOMYCIN SULFATE

CAPSULE; ORAL

PAROMOMYCIN SULFATE

!	HERITAGE PHARMS	EQ 250MG BASE	A065173 001	Dec 14, 2007
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PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>	NOVITIUM PHARMA	<u>EQ 10MG BASE/5ML</u>	<u>A215003 001</u>	Sep 03, 2021
<u>AB</u>	+! APOTEX	<u>EQ 10MG BASE/5ML</u>	<u>N020710 001</u>	Jun 25, 1997

TABLET; ORAL

PAROXETINE

<u>AB</u>	PRINSTON INC	<u>EQ 10MG BASE</u>	<u>A203854 001</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203854 002</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A203854 003</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A203854 004</u>	Oct 31, 2014

PAROXETINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 10MG BASE</u>	<u>A075356 001</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A075356 002</u>	Jul 30, 2003

PRESCRIPTION DRUG PRODUCT LIST

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A075356 003</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A075356 004</u>	Jul 30, 2003
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A078406 001</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078406 002</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A078406 003</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078406 004</u>	Jul 25, 2007
<u>AB</u>	CHARTWELL RX	<u>EQ 10MG BASE</u>	<u>A076618 001</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076618 002</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A076618 003</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076618 004</u>	Aug 15, 2005
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A078902 001</u>	Mar 13, 2008
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078902 002</u>	Mar 13, 2008
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A078902 003</u>	Mar 13, 2008
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078902 004</u>	Mar 13, 2008
<u>AB</u>	OXFORD PHARMS	<u>EQ 10MG BASE</u>	<u>A076968 001</u>	Jun 21, 2010
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076968 002</u>	Jun 21, 2010
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A076968 003</u>	Jun 21, 2010
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076968 004</u>	Jun 21, 2010
<u>AB</u>	YILING	<u>EQ 10MG BASE</u>	<u>A211248 001</u>	Nov 02, 2021
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A211248 002</u>	Nov 02, 2021
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A211248 003</u>	Nov 02, 2021
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A211248 004</u>	Nov 02, 2021
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 10MG BASE</u>	<u>A077584 001</u>	Mar 07, 2007
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077584 002</u>	Mar 07, 2007
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A077584 003</u>	Mar 07, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077584 004</u>	Mar 07, 2007

PAXIL

<u>AB</u>	+	APOTEX	<u>EQ 10MG BASE</u>	<u>N020031 001</u>	Dec 29, 1992
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N020031 002</u>	Dec 29, 1992
<u>AB</u>	+		<u>EQ 30MG BASE</u>	<u>N020031 003</u>	Dec 29, 1992
<u>AB</u>	+	!	<u>EQ 40MG BASE</u>	<u>N020031 005</u>	Dec 29, 1992

TABLET, EXTENDED RELEASE; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA USA	<u>EQ 12.5MG BASE</u>	<u>A077873 001</u>	Jun 29, 2007
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A077873 002</u>	Jun 29, 2007
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A091427 001</u>	Apr 14, 2011
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 12.5MG BASE</u>	<u>A212645 001</u>	Aug 27, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A212645 002</u>	Aug 27, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A212645 003</u>	Aug 27, 2021
<u>AB</u>	CSPC OUYI	<u>EQ 12.5MG BASE</u>	<u>A213485 001</u>	Feb 16, 2021
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A213485 002</u>	Feb 16, 2021
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A213485 003</u>	Feb 16, 2021
<u>AB</u>	LANNETT CO INC	<u>EQ 12.5MG BASE</u>	<u>A204744 001</u>	Jun 10, 2016
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204744 002</u>	Jun 10, 2016
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A204744 003</u>	Jun 10, 2016
<u>AB</u>	LUPIN LTD	<u>EQ 12.5MG BASE</u>	<u>A204134 001</u>	Jan 20, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204134 002</u>	Jan 20, 2017
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A204134 003</u>	Jan 20, 2017
<u>AB</u>	PRINSTON INC	<u>EQ 12.5MG BASE</u>	<u>A209748 001</u>	Jan 04, 2024
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A209748 002</u>	Jan 04, 2024
<u>AB</u>	SCIECURE PHARMA INC	<u>EQ 12.5MG BASE</u>	<u>A209293 001</u>	Jun 12, 2018
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A209293 002</u>	Jun 12, 2018
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A209293 003</u>	Jun 12, 2018

PAXIL CR

<u>AB</u>	+	APOTEX	<u>EQ 12.5MG BASE</u>	<u>N020936 001</u>	Feb 16, 1999
<u>AB</u>	+		<u>EQ 25MG BASE</u>	<u>N020936 002</u>	Feb 16, 1999
<u>AB</u>	+	!	<u>EQ 37.5MG BASE</u>	<u>N020936 003</u>	Dec 06, 2000

PAROXETINE MESYLATE

CAPSULE; ORAL

BRISDELLE

<u>AB</u>	+	SEBELA IRELAND LTD	<u>EQ 7.5MG BASE</u>	<u>N204516 001</u>	Jun 28, 2013
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PAROXETINE MESYLATE

<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 7.5MG BASE</u>	<u>A207139 001</u>	Jun 20, 2017
<u>AB</u>	PRINSTON INC	<u>EQ 7.5MG BASE</u>	<u>A207188 001</u>	Aug 18, 2017

PRESCRIPTION DRUG PRODUCT LISTPASIREOTIDE DIASPARTATE

SOLUTION; SUBCUTANEOUS

SIGNIFOR

+	RECORDATI RARE	EQ 0.3MG BASE/ML (EQ 0.3MG BASE/ML)	N200677 001	Dec 14, 2012
+		EQ 0.6MG BASE/ML (EQ 0.6MG BASE/ML)	N200677 002	Dec 14, 2012
+	!	EQ 0.9MG BASE/ML (EQ 0.9MG BASE/ML)	N200677 003	Dec 14, 2012

PASIREOTIDE PAMOATE

FOR SUSPENSION; INTRAMUSCULAR

SIGNIFOR LAR KIT

+	RECORDATI RARE	EQ 10MG BASE/VIAL	N203255 004	Jun 29, 2018
+		EQ 20MG BASE/VIAL	N203255 001	Dec 15, 2014
+		EQ 30MG BASE/VIAL	N203255 005	Jun 29, 2018
+		EQ 40MG BASE/VIAL	N203255 002	Dec 15, 2014
+	!	EQ 60MG BASE/VIAL	N203255 003	Dec 15, 2014

PATIROMER SORBITE X CALCIUM

POWDER; ORAL

VELTASSA

+	VIFOR PHARMA	EQ 1GM BASE/PACKET	N205739 004	Oct 02, 2023
+		EQ 8.4GM BASE/PACKET	N205739 001	Oct 21, 2015
+		EQ 16.8GM BASE/PACKET	N205739 002	Oct 21, 2015
+	!	EQ 25.2GM BASE/PACKET	N205739 003	Oct 21, 2015

PATISIRAN SODIUM

SOLUTION; INTRAVENOUS

ONPATRO

+	!	ALNYLAM PHARMS INC	EQ 10MG BASE/5ML (EQ 2MG BASE/ML)	N210922 001	Aug 10, 2018
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PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

PAZOPANIB HYDROCHLORIDE

AB		APOTEX	EQ 200MG BASE	A217713 001	Oct 19, 2023	
AB		SUN PHARM	EQ 200MG BASE	A215837 001	Oct 19, 2023	
AB		TEVA PHARMS INC	EQ 200MG BASE	A217517 001	Oct 19, 2023	
AB	+	!	NOVARTIS	EQ 200MG BASE	N022465 001	Oct 19, 2009

PEGCETACOPLAN

SOLUTION; INTRAVITREAL

SYFOVRE

+	!	APELLIS PHARMS	15MG/0.1ML (15MG/0.1ML)	N217171 001	Feb 17, 2023
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SOLUTION; SUBCUTANEOUS

EMPAVELI

+	!	APELLIS PHARMS	1080MG/20ML (54MG/ML)	N215014 001	May 14, 2021
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PEMETREXED

SOLUTION; INTRAVENOUS

PEMETREXED

+	!	ACTAVIS	100MG/4ML (25MG/ML)	N208419 001	Aug 21, 2020
+	!		500MG/20ML (25MG/ML)	N208419 002	Aug 21, 2020
+	!		1GM/40ML (25MG/ML)	N208419 003	Aug 21, 2020

PEMFEXY

+	!	EAGLE PHARMS	500MG/20ML (25MG/ML)	N209472 001	Feb 08, 2020
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PEMETREXED DISODIUM

POWDER; INTRAVENOUS

ALIMTA

AP	+	!	LILLY	EQ 100MG BASE/VIAL	N021462 002	Sep 07, 2007
AP	+	!		EQ 500MG BASE/VIAL	N021462 001	Feb 04, 2004

PEMETREXED DISODIUM

AP		ACCORD HLTHCARE	EQ 100MG BASE/VIAL	A203485 001	May 25, 2022
AP			EQ 500MG BASE/VIAL	A203485 002	May 25, 2022
AP			EQ 1GM BASE/VIAL	A203485 003	May 25, 2022
AP		APOTEX	EQ 100MG BASE/VIAL	A203774 001	May 25, 2022
AP			EQ 500MG BASE/VIAL	A203774 002	May 25, 2022
AP			EQ 750MG BASE/VIAL	A209851 001	May 25, 2022
AP			EQ 1GM BASE/VIAL	A209085 001	May 25, 2022
AP		BAXTER HLTHCARE CORP	EQ 100MG BASE/VIAL	A214436 001	Aug 18, 2022
AP			EQ 500MG BASE/VIAL	A214436 002	Aug 18, 2022
AP		DR REDDYS	EQ 100MG BASE/VIAL	A202596 001	May 25, 2022
AP			EQ 500MG BASE/VIAL	A202596 002	May 25, 2022
AP			EQ 1GM BASE/VIAL	A202596 003	May 25, 2022
AP		EUGIA PHARMA	EQ 100MG BASE/VIAL	A214632 001	May 25, 2022
AP			EQ 500MG BASE/VIAL	A214632 002	May 25, 2022

PRESCRIPTION DRUG PRODUCT LIST

PEMETREXED DISODIUM

POWDER; INTRAVENOUS

PEMETREXED DISODIUM

<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A214632 003</u>	May 25, 2022
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 100MG BASE/VIAL</u>	<u>A090384 001</u>	May 25, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A090384 002</u>	May 25, 2022
<u>AP</u>	!	<u>EQ 750MG BASE/VIAL</u>	<u>A090384 003</u>	May 25, 2022
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A090384 004</u>	May 25, 2022
<u>AP</u>	JIANGSU HANSOH PHARM	<u>EQ 100MG BASE/VIAL</u>	<u>A208696 001</u>	May 25, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A208696 002</u>	May 25, 2022
<u>AP</u>	MEITHEAL	<u>EQ 100MG BASE/VIAL</u>	<u>A215479 001</u>	Dec 13, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A215479 002</u>	Dec 13, 2022
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A215479 003</u>	Dec 13, 2022
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A215479 004</u>	Dec 13, 2022
<u>AP</u>	NANG KUANG PHARM CO	<u>EQ 100MG BASE/VIAL</u>	<u>A207352 001</u>	May 25, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A207352 002</u>	May 25, 2022
<u>AP</u>	PRINSTON INC	<u>EQ 100MG BASE/VIAL</u>	<u>A216582 001</u>	Dec 11, 2023
<u>AP</u>	QILU PHARM HAINAN	<u>EQ 100MG BASE/VIAL</u>	<u>A204890 001</u>	May 25, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A204890 002</u>	May 25, 2022
<u>AP</u>	WAVERLEY PHARMA INC	<u>EQ 100MG BASE/VIAL</u>	<u>A211899 001</u>	May 25, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A211899 002</u>	May 25, 2022
<u>AP</u>	ZYDUS PHARMS	<u>EQ 100MG BASE/VIAL</u>	<u>A214073 001</u>	May 25, 2022
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A214073 002</u>	May 25, 2022
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A214073 003</u>	May 25, 2022

SOLUTION; INTRAVENOUS

PEMETREXED

+	SHILPA	EQ 100MG/10ML BASE (10MG/ML)	N215179 001	May 22, 2023
+		EQ 500MG/50ML BASE (10MG/ML)	N215179 002	May 22, 2023
+		EQ 1GM/100ML BASE (10MG/ML)	N215179 003	May 22, 2023

PEMETREXED DISODIUM

+	ACCORD HLTHCARE	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	N214408 001	Jul 19, 2022
+		EQ 500MG BASE/20ML (EQ 25MG BASE/ML)	N214408 002	Jul 19, 2022
+		EQ 850MG BASE/34ML (EQ 25MG BASE/ML)	N214408 003	Jul 19, 2022
+		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	N214408 004	Jul 19, 2022
+	HOSPIRA	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	N214218 001	Jun 22, 2022
+		EQ 500MG BASE/20ML (EQ 25MG BASE/ML)	N214218 002	Jun 22, 2022
+		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	N214218 003	Jun 22, 2022
+	SANDOZ	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	N214657 001	May 26, 2022
+		EQ 500MG BASE/20ML (EQ 25MG BASE/ML)	N214657 002	May 26, 2022
+		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	N214657 003	May 26, 2022

PEMETREXED DITROMETHAMINE

POWDER; INTRAVENOUS

PEMETREXED DITROMETHAMINE

+	HOSPIRA	EQ 100MG BASE/VIAL	N208746 001	Jun 10, 2022
+		EQ 500MG BASE/VIAL	N208746 002	Jun 10, 2022

PEMIGATINIB

TABLET; ORAL

PEMAZYRE

+	INCYTE CORP	4.5MG	N213736 001	Apr 17, 2020
+		9MG	N213736 002	Apr 17, 2020
+		13.5MG	N213736 003	Apr 17, 2020

PENCICLOVIR

CREAM; TOPICAL

DENAVIR

<u>AB</u>	+	MYLAN	<u>1%</u>	<u>N020629 001</u>	Sep 24, 1996
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PENCICLOVIR

<u>AB</u>		TEVA PHARMS USA	<u>1%</u>	<u>A212710 001</u>	Nov 09, 2022
<u>AB</u>		TORRENT	<u>1%</u>	<u>A216981 001</u>	Aug 07, 2023

PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE

<u>AB</u>	+	VALEANT PHARMS INTL	<u>250MG</u>	<u>N019853 001</u>	
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PENICILLAMINE

<u>AB</u>		ANI PHARMS	<u>250MG</u>	<u>A209921 001</u>	May 07, 2019
<u>AB</u>		APOTEX	<u>250MG</u>	<u>A213310 001</u>	Apr 28, 2020
<u>AB</u>		BRECKENRIDGE	<u>250MG</u>	<u>A215409 001</u>	Aug 23, 2021
<u>AB</u>		DR REDDYS	<u>250MG</u>	<u>A211867 001</u>	Aug 04, 2020
<u>AB</u>		GRANULES	<u>250MG</u>	<u>A211735 001</u>	Dec 02, 2020
<u>AB</u>		INVAGEN PHARMS	<u>250MG</u>	<u>A213293 001</u>	Aug 19, 2021
<u>AB</u>		NAVINTA LLC	<u>250MG</u>	<u>A214363 001</u>	Oct 08, 2021

PRESCRIPTION DRUG PRODUCT LIST

PENICILLAMINE

CAPSULE; ORAL

PENICILLAMINE

AB	PAR PHARM INC	250MG	A211231 001	Dec 23, 2019
AB	WATSON LABS INC	250MG	A210976 001	Jun 24, 2019

TABLET; ORAL

DEPEN

AB	+ !	MYLAN SPECIALITY LP	250MG	N019854 001
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PENICILLAMINE

AB	LUPIN LTD	250MG	A212933 001	Nov 30, 2020
AB	PAR PHARM INC	250MG	A211196 001	Dec 23, 2019

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

BC	+ !	KING PHARMS LLC	600,000 UNITS/ML	N050141 001
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PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

	+ !	KING PHARMS LLC	300,000 UNITS/ML; 300,000 UNITS/ML	N050138 001
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BICILLIN C-R 900/300

	+ !	KING PHARMS LLC	900,000 UNITS/2ML; 300,000 UNITS/2ML	N050138 003
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PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

AP	ACS DOBFAR SPA	20,000,000 UNITS/VIAL	A205043 002	Oct 26, 2018
AP		5,000,000 UNITS/VIAL	A205043 001	Oct 26, 2018
AP	HQ SPECTL PHARMA	5,000,000 UNITS/VIAL	A065149 002	Jul 23, 2009
AP		20,000,000 UNITS/VIAL	A065149 003	Jul 23, 2009
AP	ISTITUTO BIO ITA SPA	5,000,000 UNITS/VIAL	A065448 001	Aug 18, 2009
AP		20,000,000 UNITS/VIAL	A065448 002	Aug 18, 2009
AP	SANDOZ	5,000,000 UNITS/VIAL	A065079 002	Aug 30, 2002
AP		20,000,000 UNITS/VIAL	A065079 003	Aug 30, 2002

PFIZERPEN

AP	!	PFIZER	5,000,000 UNITS/VIAL	A060657 002
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AP	!		20,000,000 UNITS/VIAL	A060657 003
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PENICILLIN G POTASSIUM

		HQ SPECTL PHARMA	1,000,000 UNITS/VIAL	A065149 001 Jul 23, 2009
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PENICILLIN G POTASSIUM IN PLASTIC CONTAINER

	+ !	BAXTER HLTHCARE	20,000 UNITS/ML	N050638 001 Jun 25, 1990
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	+ !		40,000 UNITS/ML	N050638 002 Jun 25, 1990
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	+ !		60,000 UNITS/ML	N050638 003 Jun 25, 1990
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PENICILLIN G PROCAINE

INJECTABLE; INJECTION

PENICILLIN G PROCAINE

	!	KING PHARMS LLC	300,000 UNITS/ML	A060101 002
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	!		600,000 UNITS/ML	A060101 001
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PENICILLIN G SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

PENICILLIN G SODIUM

	!	SANDOZ	5,000,000 UNITS/VIAL	A065068 001 Feb 26, 2001
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PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN-VK

		TEVA	EQ 125MG BASE/5ML	A060456 001
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	!		EQ 250MG BASE/5ML	A060456 002
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TABLET; ORAL

PENICILLIN V POTASSIUM

AB	AUROBINDO PHARMA	EQ 250MG BASE	A065435 001	Apr 29, 2008
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AB		EQ 500MG BASE	A065435 002	Apr 29, 2008
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AB	CHARTWELL RX	EQ 250MG BASE	A062936 001	Nov 25, 1988
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AB		EQ 500MG BASE	A062935 001	Nov 23, 1988
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AB	HIKMA PHARMS	EQ 250MG BASE	A090549 001	Oct 11, 2013
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AB		EQ 500MG BASE	A090549 002	Oct 11, 2013
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AB	SANDOZ	EQ 250MG BASE	A064071 001	Nov 30, 1995
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AB	!		EQ 500MG BASE	A064071 002
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PENICILLIN-VK

AB	TEVA	EQ 250MG BASE	A060711 002
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AB		EQ 500MG BASE	A060711 003
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PRESCRIPTION DRUG PRODUCT LIST

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

AN	+!	FRESENIUS KABI USA	300MG/VIAL	N019887 001	Jun 15, 1989
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PENTAMIDINE ISETHIONATE

AN		SETON PHARMS	300MG/VIAL	A206667 001	Apr 24, 2019
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AN		X-GEN PHARMS INC	300MG/VIAL	A206983 001	Jan 20, 2023
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INJECTABLE; INJECTION

PENTAM

AP	+!	FRESENIUS KABI USA	300MG/VIAL	N019264 001	Oct 16, 1984
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PENTAMIDINE ISETHIONATE

AP		AVET LIFESCIENCES	300MG/VIAL	A213806 001	Jan 07, 2021
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AP		SETON PHARMS	300MG/VIAL	A206666 001	Sep 28, 2017
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AP		XGEN PHARMS	300MG/VIAL	A206982 001	Mar 17, 2022
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PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUM

AP	+!	RISING	50MG/ML	A083246 001	
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PENTOBARBITAL SODIUM

AP		BPI LABS	50MG/ML	A206677 001	Nov 27, 2017
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AP		HIKMA	50MG/ML	A203619 001	Nov 13, 2017
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AP		SAGENT PHARMS INC	50MG/ML	A206404 001	May 23, 2016
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PENTOSAN POLYSULFATE SODIUM

CAPSULE; ORAL

ELMIRON

	+!	JANSSEN PHARMS	100MG	N020193 001	Sep 26, 1996
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PENTOSTATIN

INJECTABLE; INJECTION

NIPENT

AP	+!	HOSPIRA INC	10MG/VIAL	N020122 001	Oct 11, 1991
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PENTOSTATIN

AP		WEST-WARD PHARMS	10MG/VIAL	A077841 001	Aug 07, 2007
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INT

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

AB	!	APOTEX	400MG	A075191 001	Jun 09, 1999
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AB		VALEANT PHARMS	400MG	A075028 001	Jul 20, 1998
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PERAMIVIR

SOLUTION; INTRAVENOUS

RAPIVAB

	+!	BIOCRIST	200MG/20ML (10MG/ML)	N206426 001	Dec 19, 2014
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PERAMPANEL

SUSPENSION; ORAL

FYCOMPA

	+!	CATALYST PHARMS	0.5MG/ML	N208277 001	Apr 29, 2016
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TABLET; ORAL

FYCOMPA

	+	CATALYST PHARMS	2MG	N202834 001	Oct 22, 2012
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	+		4MG	N202834 002	Oct 22, 2012
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	+		6MG	N202834 003	Oct 22, 2012
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	+		8MG	N202834 004	Oct 22, 2012
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	+		10MG	N202834 005	Oct 22, 2012
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	+!		12MG	N202834 006	Oct 22, 2012
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PERFLUOROHEXYLOCTANE

SOLUTION/DROPS; OPHTHALMIC

MIEBO

	+!	BAUSCH AND LOMB INC	1.338GM/ML	N216675 001	May 18, 2023
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PERFLUTREN

INJECTABLE; INTRAVENOUS

DEFINITY

	+!	LANTHEUS MEDCL	13.04MG/2ML (6.52MG/ML)	N021064 001	Jul 31, 2001
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DEFINITY RT

	+!	LANTHEUS MEDCL	13.04MG/2ML (6.52MG/ML)	N021064 002	Nov 17, 2020
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PRESCRIPTION DRUG PRODUCT LIST

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE

AUROBINDO PHARMA	2MG
	4MG
!	8MG

A079070	001	Nov 10, 2009
A079070	002	Nov 10, 2009
A079070	003	Nov 10, 2009

PERMETHRIN

CREAM; TOPICAL

ELIMITE

AB	+	AUROBINDO PHARMA	5%
		USA	

N019855	001	Aug 25, 1989
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PERMETHRIN

AB		ACTAVIS LABS	5%
AB		DR REDDYS LABS EU	5%
AB		ENCUBE ETHICALS	5%
AB	!	PADAGIS ISRAEL	5%

A074806	001	Jan 23, 1998
A209732	001	Aug 01, 2023
A211303	001	Apr 03, 2019
A076369	001	Apr 21, 2003

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

AB		APPCO	2MG
AB			4MG
AB			8MG
AB			16MG
AB		RISING	2MG
AB			4MG
AB			8MG
AB			16MG
AB		SANDOZ	2MG
AB			4MG
AB			8MG
AB	!		16MG
AB		VINTAGE PHARMS	2MG
AB			4MG
AB			8MG
AB			16MG
AB		WATSON LABS INC	2MG
AB			4MG
AB			8MG
AB			16MG
AB		WILSHIRE PHARMS INC	2MG
AB			4MG
AB			8MG
AB			16MG
AB		ZYDUS PHARMS	2MG
AB			4MG
AB			8MG
AB			16MG

A210163	001	May 18, 2022
A210163	002	May 18, 2022
A210163	003	May 18, 2022
A210163	004	May 18, 2022
A205056	001	Mar 01, 2019
A205056	002	Mar 01, 2019
A205056	003	Mar 01, 2019
A205056	004	Mar 01, 2019
A089685	002	Dec 08, 1988
A089685	003	Dec 08, 1988
A089685	001	Dec 08, 1988
A089685	004	Dec 08, 1988
A040226	001	Dec 31, 1998
A040226	002	Dec 31, 1998
A040226	003	Dec 31, 1998
A040226	004	Dec 31, 1998
A207582	001	Oct 17, 2016
A207582	002	Oct 17, 2016
A207582	003	Oct 17, 2016
A207582	004	Oct 17, 2016
A205973	001	Dec 17, 2015
A205973	002	Dec 17, 2015
A205973	003	Dec 17, 2015
A205973	004	Dec 17, 2015
A205232	001	Apr 06, 2020
A205232	002	Apr 06, 2020
A205232	003	Apr 06, 2020
A205232	004	Apr 06, 2020

PEXIDARTINIB HYDROCHLORIDE

CAPSULE; ORAL

TURALIO

+	DAIICHI SANKYO INC	EQ 125MG BASE
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N211810	002	Oct 14, 2022
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PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

PHENDIMETRAZINE TARTRATE

+	VIRTUS	105MG
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N018074	001	
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TABLET; ORAL

BONTRIL PDM

AA	+	VALEANT	35MG
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A085272	001	
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PHENDIMETRAZINE TARTRATE

AA		CHARTWELL	35MG
AA		ELITE LABS INC	35MG
AA		KVK TECH	35MG
AA		VIRTUS	35MG

A089452	001	Oct 30, 1991
A040762	001	Jan 28, 2008
A091042	001	Aug 31, 2010
A085588	001	

PHENELZINE SULFATE

TABLET; ORAL

NARDIL

AB	+	PARKE DAVIS	EQ 15MG BASE
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N011909	002	
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PHENELZINE SULFATE

AB		NOVEL LABS INC	EQ 15MG BASE
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A200181	001	Dec 08, 2010
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PRESCRIPTION DRUG PRODUCT LIST

PHENOBARBITAL SODIUM

POWDER; INTRAVENOUS

SEZABY

+! SUN PHARM INDS INC 100MG/VIAL

N215910 001 Nov 17, 2022

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIBENZYLINEAB + CONCORDIA 10MGN008708 001PHENOXYBENZAMINE HYDROCHLORIDEAB AMNEAL 10MGA212568 001 Oct 27, 2020AB AUROBINDO PHARMA 10MGA215600 001 May 08, 2023AB NOVITIUM PHARMA 10MGA215042 001 Jul 19, 2022AB ! PAR PHARM INC 10MGA204522 001 Jan 24, 2017AB RISING 10MGA204551 001 Jun 20, 2023PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

ADIPEX-PAA +! TEVA 37.5MGA088023 001 Aug 02, 1983PHENTERMINE HYDROCHLORIDEAA AUROLIFE PHARMA LLC 15MGA204318 001 Nov 09, 2016AA 30MGA204318 002 Nov 09, 2016AA ELITE LABS 15MGA202248 001 Sep 28, 2012AA 30MGA202248 002 Sep 28, 2012AA ELITE LABS INC 37.5MGA040228 001 Jun 19, 1997AA INVAGEN PHARMS 15MGA202858 001 Feb 14, 2014AA 30MGA202858 002 Feb 14, 2014AA 30MGA204414 001 May 05, 2014AA 37.5MGA202846 001 Feb 05, 2014AA KVK TECH 15MGA040886 002 Mar 31, 2008AA 30MGA040875 001 Mar 21, 2008AA 30MGA040886 001 Mar 31, 2008AA 37.5MGA040887 001 Apr 24, 2008AA NUVO PHARM 15MGA205019 001 Dec 05, 2014AA 30MGA205019 002 Dec 05, 2014AA 37.5MGA205017 001 Sep 25, 2014AA +! SANDOZ 15MGA087190 002AA +! 30MGA086945 001 Jul 20, 1983AA +! 30MGA087190 001

TABLET; ORAL

ADIPEX-PAA +! TEVA 37.5MGA085128 001LOMAIRAAA ! AVANTHI INC 8MGA203495 001 Sep 12, 2016PHENTERMINE HYDROCHLORIDEAA AUROLIFE PHARMA LLC 37.5MGA203068 001 Aug 06, 2014AA ELITE LABS 37.5MGA200272 001 Jan 31, 2011AA ELITE LABS INC 37.5MGA040190 001 May 30, 1997AA INVAGEN PHARMS 37.5MGA202942 001 Feb 05, 2014AA KVK TECH 37.5MGA040876 001 Mar 31, 2008AA KVK TECH INC 8MGA203436 001 Mar 17, 2017AA MERRO PHARM USA 37.5MGA206342 001 Nov 18, 2016AA NUVO PHARM 37.5MGA205008 001 Sep 25, 2014AA PRINSTON INC 37.5MGA040377 001 Jan 04, 2002AA SUN PHARM 37.5MGA040526 001 Oct 23, 2003

INDUSTRIES

TABLET, ORALLY DISINTEGRATING; ORAL

PHENTERMINE HYDROCHLORIDE

ZYDUS PHARMS 15MG

A204663 001 Jun 28, 2017

30MG

A204663 002 Jun 28, 2017

37.5MG

A204663 003 Jun 28, 2017

PHENTERMINE HYDROCHLORIDE; TOPIRAMATE

CAPSULE, EXTENDED RELEASE; ORAL

QSYMIA

+ VIVUS EQ 3.75MG BASE; 23MG

N022580 001 Jul 17, 2012

+ EQ 7.5MG BASE; 46MG

N022580 002 Jul 17, 2012

+ EQ 11.25MG BASE; 69MG

N022580 003 Jul 17, 2012

+! EQ 15MG BASE; 92MG

N022580 004 Jul 17, 2012

PRESCRIPTION DRUG PRODUCT LIST

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

<u>AP</u>	!	HIKMA	<u>5MG/VIAL</u>	<u>A040235</u>	<u>001</u>	Mar 11, 1998
<u>AP</u>		PRECISION DOSE INC	<u>5MG/VIAL</u>	<u>A207686</u>	<u>001</u>	Jul 14, 2017
		ORAVERSE				
	+	SEPTODONT HOLDING	0.4MG/1.7ML	N022159	001	May 09, 2008
		SOLUTION;OPHTHALMIC				
		RYZUMVI				
	+	OCUPHIRE	EQ 0.75% BASE	N217064	001	Sep 25, 2023

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

<u>AP1</u>		ACCORD HLTHCARE	<u>10MG/ML (10MG/ML)</u>	<u>A213237</u>	<u>001</u>	Jul 01, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A213237</u>	<u>002</u>	Jul 01, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A213237</u>	<u>003</u>	Jul 01, 2020
<u>AP1</u>		AMNEAL	<u>10MG/ML (10MG/ML)</u>	<u>A211079</u>	<u>001</u>	Jul 05, 2018
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A211078</u>	<u>001</u>	Jul 19, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A211078</u>	<u>002</u>	Jul 19, 2018
<u>AP1</u>		CAPLIN	<u>10MG/ML (10MG/ML)</u>	<u>A213318</u>	<u>001</u>	Jun 11, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A213318</u>	<u>002</u>	Jun 11, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A213318</u>	<u>003</u>	Jun 11, 2020
<u>AP1</u>		EUGIA PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A210697</u>	<u>001</u>	Nov 13, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210697</u>	<u>002</u>	Nov 13, 2020
<u>AP1</u>		FRESENIUS KABI USA	<u>50MG/5ML (10MG/ML)</u>	<u>A210666</u>	<u>001</u>	Jan 30, 2019
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210666</u>	<u>002</u>	Jan 30, 2019
<u>AP1</u>		GLAND PHARMA LTD	<u>10MG/ML (10MG/ML)</u>	<u>A211920</u>	<u>003</u>	Apr 08, 2021
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A211920</u>	<u>001</u>	Jun 05, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A211920</u>	<u>002</u>	Jun 05, 2020
<u>AP1</u>		MANKIND PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>A217069</u>	<u>001</u>	Aug 30, 2022
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A217069</u>	<u>002</u>	Aug 30, 2022
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A217069</u>	<u>003</u>	Aug 30, 2022
<u>AP1</u>		MEITHEAL	<u>50MG/5ML (10MG/ML)</u>	<u>A210333</u>	<u>001</u>	Apr 27, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210333</u>	<u>002</u>	Apr 27, 2018
<u>AP1</u>		PAR STERILE PRODUCTS	<u>50MG/5ML (10MG/ML)</u>	<u>A210025</u>	<u>002</u>	Dec 21, 2018
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A210025</u>	<u>003</u>	Dec 21, 2018
<u>AP1</u>		PROVEPHARM SAS	<u>10MG/ML (10MG/ML)</u>	<u>A211081</u>	<u>001</u>	Jul 17, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A211081</u>	<u>002</u>	Jul 17, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A211081</u>	<u>003</u>	Jul 17, 2020
<u>AP1</u>		SAGENT PHARMS INC	<u>10MG/ML (10MG/ML)</u>	<u>A209967</u>	<u>001</u>	Jan 16, 2020
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A209967</u>	<u>002</u>	Jan 16, 2020
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A209967</u>	<u>003</u>	Jan 16, 2020
<u>AP1</u>		SANDOZ	<u>10MG/ML (10MG/ML)</u>	<u>A208905</u>	<u>001</u>	Jan 31, 2019
<u>AP1</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A208905</u>	<u>002</u>	Jan 31, 2019
<u>AP1</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A208905</u>	<u>003</u>	Jan 31, 2019
		<u>VAZCULEP</u>				
<u>AP1</u>	+	EXELA PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>N204300</u>	<u>001</u>	Jun 27, 2014
<u>AP1</u>	+		<u>50MG/5ML (10MG/ML)</u>	<u>N204300</u>	<u>002</u>	Jun 27, 2014
<u>AP1</u>	+		<u>100MG/10ML (10MG/ML)</u>	<u>N204300</u>	<u>003</u>	Jun 27, 2014

PHENYLEPHRINE HYDROCHLORIDE

<u>AP2</u>		AVET LIFSCIENCES	<u>10MG/ML (10MG/ML)</u>	<u>A209968</u>	<u>001</u>	Feb 28, 2023
<u>AP2</u>		BE PHARMS	<u>10MG/ML (10MG/ML)</u>	<u>A217521</u>	<u>001</u>	Jun 26, 2023
<u>AP2</u>			<u>50MG/5ML (10MG/ML)</u>	<u>A217521</u>	<u>002</u>	Jun 26, 2023
<u>AP2</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A217521</u>	<u>003</u>	Jun 26, 2023
<u>AP2</u>		EUGIA PHARMA	<u>10MG/ML (10MG/ML)</u>	<u>A210696</u>	<u>001</u>	Jan 07, 2021
<u>AP2</u>		FRESENIUS KABI USA	<u>10MG/ML (10MG/ML)</u>	<u>A210665</u>	<u>001</u>	Jan 29, 2019
<u>AP2</u>	+	HIKMA	<u>10MG/ML (10MG/ML)</u>	<u>N203826</u>	<u>001</u>	Dec 20, 2012
<u>AP2</u>		MEITHEAL	<u>10MG/ML (10MG/ML)</u>	<u>A210334</u>	<u>001</u>	Apr 27, 2018
		BIORPHEN				
	+	DR REDDYS LABS SA	0.5MG/5ML (0.1MG/ML)	N212909	001	Oct 21, 2019
		IMPPHENTIV				
	+	HIKMA	0.5MG/5ML (0.1MG/ML)	N203826	004	Mar 09, 2023
	+		1MG/10ML (0.1MG/ML)	N203826	005	Mar 09, 2023
		PHENYLEPHRINE HYDROCHLORIDE				
	+	HIKMA	50MG/5ML (10MG/ML)	N203826	002	Jun 19, 2019
	+		100MG/10ML (10MG/ML)	N203826	003	Jun 19, 2019

SOLUTION/DROPS;OPHTHALMIC

PHENYLEPHRINE HYDROCHLORIDE

<u>AT</u>	+	ALCON	<u>2.5%</u>	<u>N207926</u>	<u>001</u>	Jan 15, 2015
<u>AT</u>	+		<u>10%</u>	<u>N207926</u>	<u>002</u>	Jan 15, 2015
<u>AT</u>	!	MANKIND PHARMA	<u>2.5%</u>	<u>A216859</u>	<u>001</u>	Sep 29, 2022

PRESCRIPTION DRUG PRODUCT LIST

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

PHENYLEPHRINE HYDROCHLORIDE

AT	!		<u>10%</u>	<u>A216496</u>	<u>001</u>	Jan 11, 2023
AT	+	PARAGON BIOTECK	<u>2.5%</u>	<u>N203510</u>	<u>001</u>	Mar 21, 2013
AT	+		<u>10%</u>	<u>N203510</u>	<u>002</u>	Mar 21, 2013

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

AA		HIKMA	<u>5MG/5ML;6.25MG/5ML</u>	<u>A040675</u>	<u>001</u>	Dec 23, 2014
AA	!	PHARM ASSOC	<u>5MG/5ML;6.25MG/5ML</u>	<u>A040654</u>	<u>001</u>	Dec 07, 2006
AA		AMNEAL PHARMS	<u>5MG/5ML;6.25MG/5ML</u>	<u>A040902</u>	<u>001</u>	Aug 25, 2009

PHENYLEPHRINE HYDROCHLORIDE; TROPICAMIDE

SPRAY, METERED;OPHTHALMIC

MYDCOMBI

	+	EYENOVIA	<u>2.5%;1%</u>	N215352	001	May 05, 2023
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PHENYTOIN

SUSPENSION;ORAL

DILANTIN-125

AB	+	VIATRIS	<u>125MG/5ML</u>	<u>N008762</u>	<u>001</u>	
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PHENYTOIN

AB		PAI HOLDINGS PHARM	<u>125MG/5ML</u>	<u>A040420</u>	<u>001</u>	Apr 19, 2002
AB		TARO	<u>125MG/5ML</u>	<u>A040521</u>	<u>001</u>	Mar 08, 2004
AB		VISTAPHARM	<u>125MG/5ML</u>	<u>A040342</u>	<u>001</u>	Jan 31, 2001
AB			<u>125MG/5ML</u>	<u>A040342</u>	<u>002</u>	Aug 18, 2005

TABLET, CHEWABLE;ORAL

DILANTIN

AB	+	PHARMACIA	<u>50MG</u>	<u>A084427</u>	<u>001</u>	
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PHENYTOIN

AB		EPIC PHARMA LLC	<u>50MG</u>	<u>A040884</u>	<u>001</u>	Nov 28, 2014
AB		RISING	<u>50MG</u>	<u>A200691</u>	<u>001</u>	Dec 26, 2012
AB		TARO	<u>50MG</u>	<u>A200565</u>	<u>001</u>	Apr 17, 2014

PHENYTOIN SODIUM

CAPSULE;ORAL

DILANTIN

AB	+	VIATRIS	<u>100MG EXTENDED</u>	<u>A084349</u>	<u>002</u>	
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EXTENDED PHENYTOIN SODIUM

AB		AMNEAL PHARMS NY	<u>100MG EXTENDED</u>	<u>A040765</u>	<u>001</u>	Nov 12, 2008
AB		SUN PHARM INDS	<u>200MG EXTENDED</u>	<u>A040731</u>	<u>001</u>	Jun 30, 2008
AB			<u>300MG EXTENDED</u>	<u>A040731</u>	<u>002</u>	Jun 30, 2008
AB		TARO	<u>100MG EXTENDED</u>	<u>A040684</u>	<u>001</u>	Sep 05, 2006

PHENYTEK

AB		MYLAN	<u>200MG EXTENDED</u>	<u>A040298</u>	<u>002</u>	Dec 06, 2001
AB	!		<u>300MG EXTENDED</u>	<u>A040298</u>	<u>003</u>	Dec 06, 2001

PHENYTOIN SODIUM

AB		AUROBINDO PHARMA	<u>100MG EXTENDED</u>	<u>A204309</u>	<u>001</u>	Jun 10, 2015
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DILANTIN

	+	VIATRIS	<u>30MG EXTENDED</u>	A084349	001	
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INJECTABLE;INJECTION

PHENYTOIN SODIUM

AP		ACELLA	<u>50MG/ML</u>	<u>A040573</u>	<u>001</u>	Sep 13, 2006
AP	+	WEST-WARD PHARMS INT	<u>50MG/ML</u>	<u>A084307</u>	<u>001</u>	

PHYTONADIONE

INJECTABLE;INJECTION

PHYTONADIONE

AB1		DR REDDYS	<u>10MG/ML</u>	<u>A207719</u>	<u>001</u>	May 22, 2019
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VITAMIN K1

AB1	+	HOSPIRA	<u>10MG/ML</u>	<u>A087955</u>	<u>001</u>	Jul 25, 1983
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PHYTONADIONE

BP	+	INTL MEDICATION	<u>1MG/0.5ML</u>	A083722	001	
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VITAMIN K1

BP	+	HOSPIRA	<u>1MG/0.5ML</u>	A087954	001	Jul 25, 1983
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PHYTONADIONE

	!	CIPLA	<u>1MG/0.5ML</u>	A214596	001	Apr 22, 2022
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	!		<u>10MG/ML</u>	A214596	002	Apr 22, 2022
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TABLET;ORAL

PHYTONADIONE

AB		AGNITIO	<u>5MG</u>	<u>A213336</u>	<u>001</u>	Oct 12, 2023
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PRESCRIPTION DRUG PRODUCT LIST

PHYTONADIONE

TABLET; ORAL

PHYTONADIONE

AB	AMNEAL PHARMS CO	5MG	A209373 001	May 11, 2018
AB	SCIEGEN PHARMS INC	5MG	A213329 001	Jul 28, 2023
AB	! ZYDUS	5MG	A210189 001	Feb 20, 2019

PIFLUFOLASTAT F-18

SOLUTION; INTRAVENOUS

PYLARIFY

+	PROGENICS PHARMS INC	50ML (1-80mCi/ML)	N214793 001	May 26, 2021
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PILOCARPINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

ISOPTO CARPINE

AT	+	SANDOZ	1%	N200890 001	Jun 22, 2010
AT	+		2%	N200890 002	Jun 22, 2010
AT	+		4%	N200890 003	Jun 22, 2010

PILOCARPINE HYDROCHLORIDE

AT		AMNEAL	1%	A214193 001	Sep 21, 2020
AT			2%	A214193 002	Sep 21, 2020
AT			4%	A214193 003	Sep 21, 2020
AT		RISING	1%	A204398 001	Sep 27, 2017
AT			2%	A204398 002	Sep 27, 2017
AT			4%	A204398 003	Sep 27, 2017
AT		SOMERSET THERAPS LLC	1%	A210384 001	Nov 25, 2019
AT			2%	A210384 002	Nov 25, 2019
AT			4%	A210384 003	Nov 25, 2019

QLOSI

+	ORASIS PHARMS	0.4%	N217836 001	Oct 17, 2023
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VUITY

+	ABEVIE	1.25%	N214028 001	Oct 28, 2021
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TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

AB		AUROBINDO PHARMA LTD	5MG	A212377 001	Aug 13, 2019
AB			7.5MG	A212377 002	Aug 13, 2019
AB		IMPAX LABS	5MG	A077248 001	Mar 31, 2006
AB			7.5MG	A077248 002	Mar 31, 2006
AB		INNOGENIX	5MG	A076963 001	Dec 22, 2004
AB			7.5MG	A076963 002	Feb 27, 2007
AB		LANNETT CO INC	5MG	A077220 001	Oct 14, 2005
AB			7.5MG	A077220 002	May 06, 2009
AB		PADAGIS US	5MG	A076746 001	Nov 16, 2004

SALAGEN

AB	+	CONCORDIA	5MG	N020237 001	Mar 22, 1994
AB	+		7.5MG	N020237 002	Apr 18, 2003

PIMAVANSERIN TARTRATE

CAPSULE; ORAL

NUPLAZID

+	ACADIA PHARMS INC	EQ 34MG BASE	N210793 001	Jun 28, 2018
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TABLET; ORAL

NUPLAZID

+	ACADIA PHARMS INC	EQ 10MG BASE	N207318 002	Jun 28, 2018
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PIMECROLIMUS

CREAM; TOPICAL

ELIDEL

AB	+	BAUSCH	1%	N021302 001	Dec 13, 2001
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PIMECROLIMUS

AB		ACTAVIS LABS UT INC	1%	A209345 001	Dec 27, 2018
AB		GLENMARK PHARMS LTD	1%	A211769 001	Aug 29, 2019

PIMOZIDE

TABLET; ORAL

PIMOZIDE

		PAR PHARM	1MG	A204521 001	Sep 28, 2015
	!		2MG	A204521 002	Sep 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

PINDOLOL

TABLET; ORAL

PINDOLOL

<u>AB</u>	ANI PHARMS	<u>5MG</u>	<u>A073609 002</u>	Mar 29, 1993
<u>AB</u>		<u>10MG</u>	<u>A073609 001</u>	Mar 29, 1993
<u>AB</u>	AUROBINDO PHARMA USA	<u>5MG</u>	<u>A074019 001</u>	Sep 03, 1992
<u>AB</u>	!	<u>10MG</u>	<u>A074019 002</u>	Sep 03, 1992
<u>AB</u>	BAYSHORE PHARMS LLC	<u>5MG</u>	<u>A211712 001</u>	Aug 01, 2019
<u>AB</u>		<u>10MG</u>	<u>A211712 002</u>	Aug 01, 2019
<u>AB</u>	NOSTRUM LABS INC	<u>5MG</u>	<u>A205415 001</u>	Jan 13, 2016
<u>AB</u>		<u>10MG</u>	<u>A205415 002</u>	Jan 13, 2016
<u>AB</u>	SUN PHARM INDUSTRIES	<u>5MG</u>	<u>A074063 001</u>	Jan 27, 1994
<u>AB</u>		<u>10MG</u>	<u>A074063 002</u>	Jan 27, 1994

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOS

<u>AB</u>	+	TAKEDA PHARMS USA	<u>EQ 15MG BASE</u>	<u>N021073 001</u>	Jul 15, 1999
<u>AB</u>	+		<u>EQ 30MG BASE</u>	<u>N021073 002</u>	Jul 15, 1999
<u>AB</u>	+	!	<u>EQ 45MG BASE</u>	<u>N021073 003</u>	Jul 15, 1999

PIOGLITAZONE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 15MG BASE</u>	<u>A200044 001</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A200044 002</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A200044 003</u>	Feb 13, 2013
<u>AB</u>	ANNORA PHARMA	<u>EQ 15MG BASE</u>	<u>A204133 001</u>	Apr 07, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A204133 002</u>	Apr 07, 2014
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A204133 003</u>	Apr 07, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 15MG BASE</u>	<u>A200268 001</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A200268 002</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A200268 003</u>	Feb 13, 2013
<u>AB</u>	CHARTWELL RX	<u>EQ 15MG BASE</u>	<u>A076798 001</u>	Oct 26, 2012
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A076798 002</u>	Oct 26, 2012
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A076798 003</u>	Oct 26, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 15MG BASE</u>	<u>A202467 001</u>	Feb 06, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202467 002</u>	Feb 06, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A202467 003</u>	Feb 06, 2013
<u>AB</u>	PRINSTON INC	<u>EQ 15MG BASE</u>	<u>A207806 001</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A207806 002</u>	Apr 17, 2018
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A207806 003</u>	Apr 17, 2018
<u>AB</u>	PURACAP PHARM LLC	<u>EQ 15MG BASE</u>	<u>A206738 001</u>	Oct 06, 2017
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A206738 002</u>	Oct 06, 2017
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A206738 003</u>	Oct 06, 2017
<u>AB</u>	SANDOZ	<u>EQ 15MG BASE</u>	<u>A078670 001</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A078670 002</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A078670 003</u>	Feb 13, 2013
<u>AB</u>	SHUANGCHENG	<u>EQ 15MG BASE</u>	<u>A210165 001</u>	Jan 22, 2021
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A210165 002</u>	Jan 22, 2021
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A210165 003</u>	Jan 22, 2021
<u>AB</u>	TEVA PHARMS USA	<u>EQ 15MG BASE</u>	<u>A077210 001</u>	Jan 10, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A077210 002</u>	Jan 10, 2014
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A077210 003</u>	Jan 10, 2014
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 15MG BASE</u>	<u>A202456 001</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A202456 002</u>	Feb 13, 2013
<u>AB</u>		<u>EQ 45MG BASE</u>	<u>A202456 003</u>	Feb 13, 2013

PIPERACILLIN SODIUM

INJECTABLE; INJECTION

PIPERACILLIN

!	ISTITUTO BIO ITA SPA	EQ 2GM BASE/VIAL	A065114 001	Nov 14, 2003
!		EQ 3GM BASE/VIAL	A065114 002	Nov 14, 2003
!		EQ 4GM BASE/VIAL	A065114 003	Nov 14, 2003
!		EQ 40GM BASE/VIAL	A065157 001	Jul 12, 2004

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

<u>AP</u>	ASTRAL	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A212287 001</u>	Jul 29, 2019
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A212287 002</u>	Jul 29, 2019
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A212287 003</u>	Jul 29, 2019
<u>AP</u>	EUGIA PHARMA	<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A217409 002</u>	Oct 12, 2023

PRESCRIPTION DRUG PRODUCT LIST

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A217409 001</u>	Oct 12, 2023
<u>AP</u>	EUGIA PHARMA SPECLTS	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065498 001</u>	May 23, 2011
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065498 002</u>	May 23, 2011
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065498 003</u>	May 23, 2011
<u>AP</u>	FRESENIUS KABI	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A203719 001</u>	May 18, 2018
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A203719 002</u>	May 18, 2018
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A203719 003</u>	May 18, 2018
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A203720 001</u>	May 11, 2018
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A206204 001</u>	May 11, 2018
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065523 001</u>	May 31, 2011
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065523 002</u>	May 31, 2011
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065523 003</u>	May 31, 2011
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A090498 001</u>	May 31, 2011
<u>AP</u>	PROVEPHARM SAS	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A207847 001</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A207847 002</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A207848 002</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A207847 003</u>	Jan 13, 2017
<u>AP</u>		<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A207848 001</u>	May 11, 2018
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A208674 001</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A208674 002</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A208674 003</u>	Feb 16, 2021
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A208675 001</u>	Feb 16, 2021
<u>AP</u>	SANDOZ	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065362 001</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065363 001</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065362 002</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065363 002</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065362 003</u>	Oct 21, 2010
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065363 003</u>	Oct 21, 2010
<u>AP</u>	!	<u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u>	<u>A203557 001</u>	Oct 29, 2014
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A203557 002</u>	Jul 08, 2021
<u>AP</u>	SHANDONG	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A204959 001</u>	Aug 10, 2018
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A204959 002</u>	Aug 10, 2018
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A204959 003</u>	Aug 10, 2018
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A217243 001</u>	Apr 14, 2023
<u>AP</u>	STERISCIENCE	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065458 001</u>	Aug 15, 2014
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065458 002</u>	Aug 15, 2014
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065458 003</u>	Aug 15, 2014
<u>AP</u>	! WOCKHARDT BIO AG	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A206996 001</u>	Mar 22, 2017
<u>AP</u>	!	<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A206996 002</u>	Mar 22, 2017
<u>AP</u>	!	<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A206996 003</u>	Mar 22, 2017
<u>AP</u>	!	<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A207146 001</u>	Mar 17, 2017
	ZOSYN IN PLASTIC CONTAINER			
	+! BAXTER HLTHCARE CORP	EQ 40MG BASE/ML;EQ 5MG BASE/ML	N050750 001	Feb 24, 1998
	+!	EQ 60MG BASE/ML;EQ 7.5MG BASE/ML	N050750 002	Feb 24, 1998
	+!	EQ 4GM BASE/100ML;EQ 500MG BASE/100ML	N050750 003	Feb 24, 1998

PIRFENIDONE

CAPSULE; ORAL

ESBRIET

<u>AB</u>	+! GENENTECH INC	<u>267MG</u>	<u>N022535 001</u>	Oct 15, 2014
	<u>PIRFENIDONE</u>			
<u>AB</u>	ACCORD HLTHCARE	<u>267MG</u>	<u>A212731 001</u>	Jan 20, 2022
<u>AB</u>	AMNEAL	<u>267MG</u>	<u>A212569 001</u>	Jan 03, 2022
<u>AB</u>	LAURUS	<u>267MG</u>	<u>A212724 001</u>	Jul 19, 2022
<u>AB</u>	SANDOZ	<u>267MG</u>	<u>A212600 001</u>	Jun 13, 2022
<u>AB</u>	SCIEGEN PHARMS INC	<u>267MG</u>	<u>A212077 001</u>	Aug 01, 2022

TABLET; ORAL

ESBRIET

<u>AB</u>	+ GENENTECH INC	<u>267MG</u>	<u>N208780 001</u>	Jan 11, 2017
<u>AB</u>	+!	<u>801MG</u>	<u>N208780 003</u>	Jan 11, 2017
	<u>PIRFENIDONE</u>			
<u>AB</u>	ACCORD HLTHCARE	<u>267MG</u>	<u>A212730 001</u>	Jan 25, 2022
<u>AB</u>		<u>801MG</u>	<u>A212730 002</u>	Jan 25, 2022
<u>AB</u>	AIZANT	<u>267MG</u>	<u>A212747 001</u>	Jul 21, 2022
<u>AB</u>		<u>534MG</u>	<u>A212747 002</u>	Jul 21, 2022
<u>AB</u>		<u>801MG</u>	<u>A212747 003</u>	Jul 21, 2022
<u>AB</u>	ALEMBIC	<u>267MG</u>	<u>A212708 001</u>	May 20, 2022

PRESCRIPTION DRUG PRODUCT LIST

PIRFENIDONE

TABLET; ORAL

PIRFENIDONE

<u>AB</u>		<u>801MG</u>	<u>A212708</u>	<u>002</u>	May 20, 2022
<u>AB</u>	AMNEAL	<u>267MG</u>	<u>A212570</u>	<u>001</u>	Mar 25, 2022
<u>AB</u>		<u>801MG</u>	<u>A212570</u>	<u>002</u>	Mar 25, 2022
<u>AB</u>	APOTEX	<u>267MG</u>	<u>A212709</u>	<u>001</u>	May 31, 2023
<u>AB</u>		<u>801MG</u>	<u>A212709</u>	<u>002</u>	May 31, 2023
<u>AB</u>	HETERO LABS LTD V	<u>267MG</u>	<u>A212674</u>	<u>001</u>	Sep 21, 2022
<u>AB</u>		<u>801MG</u>	<u>A212674</u>	<u>002</u>	Sep 21, 2022
<u>AB</u>	LAURUS	<u>267MG</u>	<u>A212722</u>	<u>001</u>	Jul 19, 2022
<u>AB</u>		<u>534MG</u>	<u>A212722</u>	<u>002</u>	Jul 19, 2022
<u>AB</u>		<u>801MG</u>	<u>A212722</u>	<u>003</u>	Jul 19, 2022
<u>AB</u>	MICRO LABS	<u>267MG</u>	<u>A212680</u>	<u>001</u>	May 18, 2022
<u>AB</u>		<u>801MG</u>	<u>A212680</u>	<u>002</u>	May 18, 2022
<u>AB</u>	MSN	<u>267MG</u>	<u>A212772</u>	<u>001</u>	May 24, 2022
<u>AB</u>		<u>801MG</u>	<u>A212772</u>	<u>002</u>	May 24, 2022
<u>AB</u>	SANDOZ	<u>267MG</u>	<u>A212560</u>	<u>001</u>	Apr 28, 2022
<u>AB</u>		<u>801MG</u>	<u>A212560</u>	<u>002</u>	Apr 28, 2022
<u>AB</u>	SCIEGEN PHARMS INC	<u>267MG</u>	<u>A212078</u>	<u>001</u>	Aug 01, 2022
<u>AB</u>		<u>801MG</u>	<u>A212078</u>	<u>002</u>	Aug 01, 2022
<u>AB</u>	TEVA PHARMS USA	<u>267MG</u>	<u>A212759</u>	<u>001</u>	Jan 25, 2022
<u>AB</u>		<u>801MG</u>	<u>A212759</u>	<u>002</u>	Jan 25, 2022

PIROXICAM

CAPSULE; ORAL

FELDENE

<u>AB</u>	+	PFIZER	<u>10MG</u>	<u>N018147</u>	<u>002</u>	Apr 06, 1982
<u>AB</u>	+	!	<u>20MG</u>	<u>N018147</u>	<u>003</u>	Apr 06, 1982

PIROXICAM

<u>AB</u>	CADILA	<u>10MG</u>	<u>A205585</u>	<u>001</u>	Jul 17, 2018
<u>AB</u>		<u>20MG</u>	<u>A205585</u>	<u>002</u>	Jul 17, 2018
<u>AB</u>	FLAMINGO PHARMS	<u>10MG</u>	<u>A207938</u>	<u>001</u>	Sep 09, 2016
<u>AB</u>		<u>20MG</u>	<u>A207938</u>	<u>002</u>	Sep 09, 2016
<u>AB</u>	MICRO LABS	<u>10MG</u>	<u>A206152</u>	<u>001</u>	Dec 29, 2017
<u>AB</u>		<u>20MG</u>	<u>A206152</u>	<u>002</u>	Dec 29, 2017
<u>AB</u>	NOSTRUM LABS INC	<u>10MG</u>	<u>A074118</u>	<u>002</u>	Jun 15, 1993
<u>AB</u>		<u>20MG</u>	<u>A074118</u>	<u>001</u>	Jun 15, 1993
<u>AB</u>	STRIDES PHARMA	<u>10MG</u>	<u>A206136</u>	<u>001</u>	Jun 20, 2017
<u>AB</u>		<u>10MG</u>	<u>A210347</u>	<u>001</u>	Jan 26, 2018
<u>AB</u>		<u>20MG</u>	<u>A206136</u>	<u>002</u>	Jun 20, 2017
<u>AB</u>		<u>20MG</u>	<u>A210347</u>	<u>002</u>	Jan 26, 2018
<u>AB</u>	TEVA	<u>10MG</u>	<u>A074131</u>	<u>001</u>	Dec 11, 1992
<u>AB</u>		<u>20MG</u>	<u>A074131</u>	<u>002</u>	Dec 11, 1992
<u>AB</u>	UNICHEM	<u>10MG</u>	<u>A208340</u>	<u>001</u>	Apr 13, 2017
<u>AB</u>		<u>20MG</u>	<u>A208340</u>	<u>002</u>	Apr 13, 2017

PIRTOBRUTINIB

TABLET; ORAL

JAYPIRCA

	+	LOXO ONCOL	50MG	N216059	001	Jan 27, 2023
	+	!	100MG	N216059	002	Jan 27, 2023

PITAVASTATIN CALCIUM

TABLET; ORAL

LIVALO

<u>AB</u>	+	KOWA CO	<u>EQ 1MG BASE</u>	<u>N022363</u>	<u>001</u>	Aug 03, 2009
<u>AB</u>	+		<u>EQ 2MG BASE</u>	<u>N022363</u>	<u>002</u>	Aug 03, 2009
<u>AB</u>	+	!	<u>EQ 4MG BASE</u>	<u>N022363</u>	<u>003</u>	Aug 03, 2009

PITAVASTATIN CALCIUM

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 1MG BASE</u>	<u>A206015</u>	<u>001</u>	Dec 20, 2016
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A206015</u>	<u>002</u>	Dec 20, 2016
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A206015</u>	<u>003</u>	Dec 20, 2016
<u>AB</u>	LUPIN LTD	<u>EQ 1MG BASE</u>	<u>A206029</u>	<u>001</u>	Nov 20, 2023
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A206029</u>	<u>002</u>	Nov 20, 2023
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A206029</u>	<u>003</u>	Nov 20, 2023
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>A206070</u>	<u>001</u>	Apr 04, 2019
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A206070</u>	<u>002</u>	Apr 04, 2019
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A206070</u>	<u>003</u>	Apr 04, 2019
<u>AB</u>	ORIENT PHARMA CO LTD	<u>EQ 1MG BASE</u>	<u>A205932</u>	<u>001</u>	Feb 03, 2017
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205932</u>	<u>002</u>	Feb 03, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205932</u>	<u>003</u>	Feb 03, 2017
<u>AB</u>	SAWAI USA	<u>EQ 1MG BASE</u>	<u>A205955</u>	<u>001</u>	Feb 03, 2017

PRESCRIPTION DRUG PRODUCT LIST

PITAVASTATIN CALCIUM

TABLET; ORAL

PITAVASTATIN CALCIUM

<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205955 002</u>	Feb 03, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205955 003</u>	Feb 03, 2017
<u>AB</u>	ZYDUS PHARMS	<u>EQ 1MG BASE</u>	<u>A206047 001</u>	Feb 22, 2023
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A206047 002</u>	Feb 22, 2023
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A206047 003</u>	Feb 22, 2023

PITAVASTATIN MAGNESIUM

TABLET; ORAL

ZYPITAMAG

+	MEDICURE	EQ 2MG BASE	N208379 002	Jul 14, 2017
+	!	EQ 4MG BASE	N208379 003	Jul 14, 2017

PITOLISANT HYDROCHLORIDE

TABLET; ORAL

WAKIX

+	HARMONY	EQ 4.45MG BASE	N211150 001	Aug 14, 2019
+	!	EQ 17.8MG BASE	N211150 002	Aug 14, 2019

PLAZOMICIN SULFATE

SOLUTION; INTRAVENOUS

ZEMDRI

+	CIPLA USA	EQ 500MG BASE/10ML (EQ 50MG BASE/ML)	N210303 001	Jun 25, 2018
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PLECANATIDE

TABLET; ORAL

TRULANCE

+	SALIX	3MG	N208745 001	Jan 19, 2017
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PLERIXAFOR

SOLUTION; SUBCUTANEOUS

MOZOBI

<u>AP</u>	+	GENZYME	<u>24MG/1.2ML (20MG/ML)</u>	<u>N022311 001</u>	Dec 15, 2008
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PLERIXAFOR

<u>AP</u>		AMNEAL	<u>24MG/1.2ML (20MG/ML)</u>	<u>A215334 001</u>	Jul 24, 2023
<u>AP</u>		DR REDDYS	<u>24MG/1.2ML (20MG/ML)</u>	<u>A205182 001</u>	Jul 24, 2023
<u>AP</u>		EUGIA PHARMA	<u>24MG/1.2ML (20MG/ML)</u>	<u>A213672 001</u>	Jul 24, 2023
<u>AP</u>		KINDOS	<u>24MG/1.2ML (20MG/ML)</u>	<u>A215698 001</u>	Jul 24, 2023
<u>AP</u>		MSN	<u>24MG/1.2ML (20MG/ML)</u>	<u>A211901 001</u>	Jul 24, 2023
<u>AP</u>		TEVA PHARMS USA INC	<u>24MG/1.2ML (20MG/ML)</u>	<u>A205197 001</u>	Jul 24, 2023
<u>AP</u>		ZYDUS PHARMS	<u>24MG/1.2ML (20MG/ML)</u>	<u>A208980 001</u>	Jul 26, 2023

PODOFILOX

GEL; TOPICAL

CONDYLOX

<u>AB</u>	+	ALLERGAN	<u>0.5%</u>	<u>N020529 001</u>	Mar 13, 1997
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PODOFILOX

<u>AB</u>		PADAGIS US	<u>0.5%</u>	<u>A211871 001</u>	Nov 22, 2023
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SOLUTION; TOPICAL

CONDYLOX

<u>AT</u>	+	TEVA BRANDED PHARM	<u>0.5%</u>	<u>N019795 001</u>	Dec 13, 1990
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PODOFILOX

<u>AT</u>		PADAGIS US	<u>0.5%</u>	<u>A075600 001</u>	Jan 29, 2002
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POLIDOCANOL

SOLUTION; INTRAVENOUS

ASCLERA

+	CHEMISCH FBRK	10MG/2ML (5MG/ML)	N021201 001	Mar 30, 2010
	KRSSLR			

+	!		20MG/2ML (10MG/ML)	N021201 002	Mar 30, 2010
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VARITHENA

+	!	PROVENSIS	180MG/18ML (10MG/ML)	N205098 001	Nov 25, 2013
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POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

NULYTELY

<u>AA</u>	+	BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N019797 001</u>	Apr 22, 1991
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NULYTELY-FLAVORED

<u>AA</u>	+	BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N019797 002</u>	Nov 18, 1994
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PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

<u>AA</u>		STRIDES PHARMA	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A204559 001</u>	Apr 13, 2015
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PRESCRIPTION DRUG PRODUCT LIST

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION; ORAL

COLYTE WITH FLAVOR PACKS

AA	MYLAN SPECIALITY LP	<u>240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT</u>	N018983 012	Oct 08, 1998
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GOLYTELY

AA	+! BRAINTREE	<u>236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT</u>	N019011 001	Jul 13, 1984
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PEG 3350 AND ELECTROLYTES

AA	NOVEL LABS INC	<u>236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT</u>	A090231 001	Jun 01, 2009
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AA		<u>240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT</u>	A090186 001	Jun 01, 2009
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POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

AA	STRIDES PHARMA	<u>236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT</u>	A204558 001	Dec 21, 2018
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POLYMYXIN B SULFATE

INJECTABLE; INJECTION

POLYMYXIN B SULFATE

AP	BEOWULF ASSET	<u>EQ 500,000 UNITS BASE/VIAL</u>	A063000 001	Sep 30, 1994
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AP	EUGIA PHARMA	<u>EQ 500,000 UNITS BASE/VIAL</u>	A206589 001	Apr 04, 2016
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AP	FRESENIUS KABI USA	<u>EQ 500,000 UNITS BASE/VIAL</u>	A065372 001	Jan 10, 2008
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AP	GLAND	<u>EQ 500,000 UNITS BASE/VIAL</u>	A207322 001	Apr 14, 2016
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AP	! XELLIA PHARMS APS	<u>EQ 500,000 UNITS BASE/VIAL</u>	A202766 001	Jan 15, 2014
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POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS; OPHTHALMIC

TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE

AT	AKORN	<u>10,000 UNITS/ML; EQ 1MG BASE/ML</u>	A065006 001	Dec 17, 1998
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AT	BAUSCH AND LOMB	<u>10,000 UNITS/ML; EQ 1MG BASE/ML</u>	A064120 001	Feb 14, 1997
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AT	! SANDOZ	<u>10,000 UNITS/ML; EQ 1MG BASE/ML</u>	A064211 001	Apr 13, 1998
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POMALIDOMIDE

CAPSULE; ORAL

POMALYST

+	BRISTOL	1MG	N204026 001	Feb 08, 2013
+		2MG	N204026 002	Feb 08, 2013
+		3MG	N204026 003	Feb 08, 2013
+	!	4MG	N204026 004	Feb 08, 2013

PONATINIB HYDROCHLORIDE

TABLET; ORAL

ICLUSIG

+	TAKEDA PHARMS USA	EQ 10MG BASE	N203469 004	Dec 18, 2020
+		EQ 15MG BASE	N203469 001	Dec 14, 2012
+		EQ 30MG BASE	N203469 003	Apr 23, 2015
+	!	EQ 45MG BASE	N203469 002	Dec 14, 2012

PONESIMOD

TABLET; ORAL

PONVORY

+	JANSSEN PHARMS	2MG	N213498 001	Mar 18, 2021
+		3MG	N213498 002	Mar 18, 2021
+		4MG	N213498 003	Mar 18, 2021
+		5MG	N213498 004	Mar 18, 2021
+		6MG	N213498 005	Mar 18, 2021
+		7MG	N213498 006	Mar 18, 2021
+		8MG	N213498 007	Mar 18, 2021
+		9MG	N213498 008	Mar 18, 2021
+		10MG	N213498 009	Mar 18, 2021
+		20MG	N213498 010	Mar 18, 2021

PORFIMER SODIUM

INJECTABLE; INJECTION

PHOTOFRIN

	PINNACLE BIOLGS	75MG/VIAL	N020451 001	Dec 27, 1995
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POSACONAZOLE

FOR SUSPENSION, DELAYED RELEASE; ORAL

NOXAFIL POWDERMIX KIT

+	MSD MERCK CO	300MG	N214770 001	May 31, 2021
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SOLUTION; INTRAVENOUS

NOXAFIL

AP	+! MERCK SHARP DOHME	<u>300MG/16.7ML (18MG/ML)</u>	N205596 001	Mar 13, 2014
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PRESCRIPTION DRUG PRODUCT LIST

POSACONAZOLE

SOLUTION; INTRAVENOUS

POSACONAZOLE

<u>AP</u>	EUGIA PHARMA	<u>300MG/16.7ML (18MG/ML)</u>	<u>A214842 001</u>	Dec 26, 2023
<u>AP</u>	FRESENIUS KABI USA	<u>300MG/16.7ML (18MG/ML)</u>	<u>A209983 001</u>	Dec 26, 2023
<u>AP</u>	GLAND PHARMA LTD	<u>300MG/16.7ML (18MG/ML)</u>	<u>A217553 001</u>	Dec 26, 2023
<u>AP</u>	MYLAN LABS LTD	<u>300MG/16.7ML (18MG/ML)</u>	<u>A211500 001</u>	Dec 26, 2023
<u>AP</u>	PAR STERILE PRODUCTS	<u>300MG/16.7ML (18MG/ML)</u>	<u>A208768 001</u>	May 25, 2022

SUSPENSION; ORAL

NOXAFIL

<u>AB</u>	+! SCHERING	<u>40MG/ML</u>	<u>N022003 001</u>	Sep 15, 2006
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POSACONAZOLE

<u>AB</u>	HIKMA	<u>40MG/ML</u>	<u>A208773 001</u>	May 15, 2020
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TABLET, DELAYED RELEASE; ORAL

NOXAFIL

<u>AB</u>	+! MERCK SHARP DOHME	<u>100MG</u>	<u>N205053 001</u>	Nov 25, 2013
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POSACONAZOLE

<u>AB</u>	AET PHARMA	<u>100MG</u>	<u>A213454 001</u>	Feb 01, 2021
<u>AB</u>	AMNEAL	<u>100MG</u>	<u>A216626 001</u>	Dec 22, 2022
<u>AB</u>	BIOCON PHARMA	<u>100MG</u>	<u>A214476 001</u>	Feb 04, 2022
<u>AB</u>	DR REDDYS	<u>100MG</u>	<u>A212500 001</u>	Apr 07, 2022
<u>AB</u>	HETERO LABS LTD III	<u>100MG</u>	<u>A214321 001</u>	Dec 08, 2022
<u>AB</u>	I 3 PHARMS	<u>100MG</u>	<u>A216488 001</u>	Aug 28, 2023
<u>AB</u>	SINOTHERAPEUTICS INC	<u>100MG</u>	<u>A212411 001</u>	Aug 21, 2019
<u>AB</u>	SPECGX LLC	<u>100MG</u>	<u>A212226 001</u>	May 10, 2022
<u>AB</u>	WESTMINSTER PHARMS	<u>100MG</u>	<u>A216326 001</u>	Jun 20, 2023

POTASSIUM ACETATE

INJECTABLE; INJECTION

POTASSIUM ACETATE

<u>AP</u>	EXELA PHARMA	<u>2MEQ/ML</u>	<u>A206203 001</u>	Dec 29, 2015
<u>AP</u>		<u>2MEQ/ML</u>	<u>A212692 001</u>	Oct 20, 2021
<u>AP</u>	+! HOSPIRA	<u>2MEQ/ML</u>	<u>N018896 001</u>	Jul 20, 1984

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

KLOR-CON

<u>AB</u>	UPSHER SMITH LABS	<u>8MEQ</u>	<u>A203106 001</u>	Jul 10, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A203106 002</u>	Jul 10, 2015

POTASSIUM CHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>8MEQ</u>	<u>A077419 001</u>	Jun 02, 2008
<u>AB</u>	!	<u>10MEQ</u>	<u>A077419 002</u>	Jun 02, 2008
<u>AB</u>	ADARE PHARMS INC	<u>8MEQ</u>	<u>A208864 001</u>	Mar 17, 2017
<u>AB</u>		<u>10MEQ</u>	<u>A208864 002</u>	Mar 17, 2017
<u>AB</u>	AMNEAL PHARMS	<u>10MEQ</u>	<u>A202128 001</u>	Feb 22, 2013
<u>AB</u>	ANCHEN PHARMS	<u>8MEQ</u>	<u>A202886 001</u>	Dec 26, 2013
<u>AB</u>		<u>10MEQ</u>	<u>A202886 002</u>	Dec 26, 2013
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MEQ</u>	<u>A202868 001</u>	Jan 19, 2016
<u>AB</u>	GRANULES	<u>8MEQ</u>	<u>A214686 001</u>	Feb 16, 2021
<u>AB</u>		<u>10MEQ</u>	<u>A214686 002</u>	Feb 16, 2021
<u>AB</u>	LUPIN LTD	<u>8MEQ</u>	<u>A203002 001</u>	Dec 18, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A203002 002</u>	Dec 18, 2015
<u>AB</u>	NOVEL LABS INC	<u>8MEQ</u>	<u>A204828 001</u>	Aug 16, 2016
<u>AB</u>		<u>10MEQ</u>	<u>A204828 002</u>	Aug 16, 2016
<u>AB</u>	PADAGIS US	<u>8MEQ</u>	<u>A200185 001</u>	May 18, 2011
<u>AB</u>		<u>10MEQ</u>	<u>A200185 002</u>	May 18, 2011
<u>AB</u>	PRINSTON INC	<u>8MEQ</u>	<u>A209026 001</u>	Jun 11, 2019
<u>AB</u>		<u>10MEQ</u>	<u>A209026 002</u>	Jun 11, 2019
<u>AB</u>	STRIDES PHARMA	<u>8MEQ</u>	<u>A205549 001</u>	Dec 08, 2015
<u>AB</u>		<u>10MEQ</u>	<u>A205549 002</u>	Dec 08, 2015
<u>AB</u>	ZYDUS PHARMS	<u>8MEQ</u>	<u>A208445 001</u>	Mar 11, 2019
<u>AB</u>		<u>10MEQ</u>	<u>A208445 002</u>	Mar 11, 2019

FOR SOLUTION; ORAL

KLOR-CON

<u>AA</u>	UPSHER SMITH LABS	<u>20MEQ</u>	<u>A209662 001</u>	Oct 23, 2017
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POTASSIUM CHLORIDE

<u>AA</u>	AMNEAL	<u>20MEQ</u>	<u>A210902 001</u>	May 23, 2019
<u>AA</u>	BELCHER	<u>20MEQ</u>	<u>A212183 001</u>	May 06, 2019
<u>AA</u>	EPIC PHARMA LLC	<u>20MEQ</u>	<u>A210200 001</u>	Nov 23, 2018
<u>AA</u>	GRANULES	<u>20MEQ</u>	<u>A213467 001</u>	Jan 27, 2022
<u>AA</u>	NOVEL LABS INC	<u>20MEQ</u>	<u>A210241 001</u>	Nov 21, 2018
<u>AA</u>	NOVITIUM PHARMA	<u>20MEQ</u>	<u>A212816 001</u>	Jul 12, 2023

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

FOR SOLUTION;ORAL

POTASSIUM CHLORIDE

<u>AA</u>	+	PHARMA RES SOFTWARE	<u>20MEQ</u>	<u>N208019</u>	<u>001</u>	Aug 19, 2015
<u>AA</u>		RUBICON	<u>20MEQ</u>	<u>A214108</u>	<u>001</u>	Mar 24, 2022
<u>AA</u>		STRIDES PHARMA	<u>20MEQ</u>	<u>A211667</u>	<u>001</u>	Mar 11, 2021
		POKONZA				
	+	PHARMA RES SOFTWARE	10MEQ	N208019	002	Aug 25, 2023
		POTASSIUM CHLORIDE				
	+	PHARMA RES SOFTWARE	40MEQ	N208019	003	Aug 25, 2023

INJECTABLE;INJECTION

POTASSIUM CHLORIDE

<u>AP</u>		B BRAUN	<u>2MEQ/ML</u>	<u>A085870</u>	<u>001</u>	
<u>AP</u>	+	HOSPIRA	<u>2MEQ/ML</u>	<u>A080205</u>	<u>001</u>	
<u>AP</u>		NEXUS PHARMS	<u>2MEQ/ML</u>	<u>A217704</u>	<u>001</u>	Aug 14, 2023
		<u>POTASSIUM CHLORIDE 10MEQ</u>				
<u>AP</u>		FRESENIUS KABI USA	<u>14.9MG/ML</u>	<u>A211087</u>	<u>001</u>	Sep 09, 2020
<u>AP</u>			<u>746MG/100ML</u>	<u>A211087</u>	<u>002</u>	Sep 09, 2020
<u>AP</u>		NEXUS	<u>14.9MG/ML</u>	<u>A214727</u>	<u>001</u>	Mar 18, 2021
<u>AP</u>			<u>745MG/100ML</u>	<u>A214727</u>	<u>002</u>	Mar 18, 2021

POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>14.9MG/ML</u>	<u>N019904</u>	<u>001</u>	Dec 26, 1989
<u>AP</u>	+		<u>746MG/100ML</u>	<u>N019904</u>	<u>005</u>	Dec 17, 1990
<u>AP</u>	+	ICU MEDICAL INC	<u>14.9MG/ML</u>	<u>N020161</u>	<u>005</u>	Nov 30, 1992
<u>AP</u>	+		<u>745MG/100ML</u>	<u>N020161</u>	<u>001</u>	Nov 30, 1992

POTASSIUM CHLORIDE 20MEQ

<u>AP</u>		FRESENIUS KABI USA	<u>29.8MG/ML</u>	<u>A211087</u>	<u>003</u>	Sep 09, 2020
<u>AP</u>			<u>1.49GM/100ML</u>	<u>A211087</u>	<u>005</u>	May 07, 2021
<u>AP</u>		NEXUS	<u>29.8MG/ML</u>	<u>A214727</u>	<u>003</u>	Mar 18, 2021
<u>AP</u>			<u>1.49GM/100ML</u>	<u>A214727</u>	<u>004</u>	Mar 18, 2021

POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>29.8MG/ML</u>	<u>N019904</u>	<u>002</u>	Dec 26, 1989
<u>AP</u>	+		<u>1.49GM/100ML</u>	<u>N019904</u>	<u>006</u>	Dec 17, 1990
<u>AP</u>	+	ICU MEDICAL INC	<u>29.8MG/ML</u>	<u>N020161</u>	<u>006</u>	Aug 11, 1998
<u>AP</u>	+		<u>1.49GM/100ML</u>	<u>N020161</u>	<u>002</u>	Nov 30, 1992

POTASSIUM CHLORIDE 40MEQ

<u>AP</u>		FRESENIUS KABI USA	<u>2.98GM/100ML</u>	<u>A211087</u>	<u>004</u>	Sep 09, 2020
<u>AP</u>		NEXUS	<u>2.98GM/100ML</u>	<u>A214727</u>	<u>005</u>	Mar 18, 2021

POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>2.98GM/100ML</u>	<u>N019904</u>	<u>004</u>	Dec 26, 1989
<u>AP</u>	+	ICU MEDICAL INC	<u>2.98GM/100ML</u>	<u>N020161</u>	<u>004</u>	Aug 11, 1998

POTASSIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>		FRESENIUS KABI USA	<u>2MEQ/ML</u>	<u>A088901</u>	<u>001</u>	Jan 25, 1985
<u>AP</u>			<u>2MEQ/ML</u>	<u>A088908</u>	<u>001</u>	Jan 25, 1985
		POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER				
	+	BAXTER HLTHCARE	2.24GM/100ML	N019904	003	Dec 26, 1989

SOLUTION;ORAL

POTASSIUM CHLORIDE

<u>AA</u>		AMNEAL	<u>20MEQ/15ML</u>	<u>A210041</u>	<u>001</u>	Jul 19, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A210041</u>	<u>002</u>	Jul 19, 2018
<u>AA</u>		ANDA REPOSITORY	<u>20MEQ/15ML</u>	<u>A214892</u>	<u>001</u>	Dec 30, 2022
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A214892</u>	<u>002</u>	Dec 30, 2022
<u>AA</u>		APOTEX	<u>20MEQ/15ML</u>	<u>A211067</u>	<u>001</u>	Aug 08, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A211067</u>	<u>002</u>	Aug 08, 2018
<u>AA</u>		BELCHER	<u>20MEQ/15ML</u>	<u>A216156</u>	<u>001</u>	Mar 07, 2023
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A216156</u>	<u>002</u>	Mar 07, 2023
<u>AA</u>	+	GENUS LIFESCIENCES	<u>20MEQ/15ML</u>	<u>N206814</u>	<u>001</u>	Dec 22, 2014
<u>AA</u>	+		<u>40MEQ/15ML</u>	<u>N206814</u>	<u>002</u>	Dec 22, 2014
<u>AA</u>		GRANULES	<u>20MEQ/15ML</u>	<u>A213392</u>	<u>001</u>	Jan 29, 2021
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A213392</u>	<u>002</u>	Jan 29, 2021
<u>AA</u>		NOVEL LABS INC	<u>20MEQ/15ML</u>	<u>A209786</u>	<u>001</u>	Aug 29, 2018
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A209786</u>	<u>002</u>	Aug 29, 2018
<u>AA</u>		PHARM ASSOC	<u>20MEQ/15ML</u>	<u>A210766</u>	<u>001</u>	Mar 29, 2019
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A210766</u>	<u>002</u>	Mar 29, 2019
<u>AA</u>		RUBICON	<u>20MEQ/15ML</u>	<u>A214656</u>	<u>001</u>	Jan 13, 2022
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A214656</u>	<u>002</u>	Jan 13, 2022
<u>AA</u>		SAPTALIS PHARMS	<u>20MEQ/15ML</u>	<u>A211648</u>	<u>001</u>	May 21, 2021
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A211648</u>	<u>002</u>	May 21, 2021
<u>AA</u>		WES PHARMA INC	<u>20MEQ/15ML</u>	<u>A213062</u>	<u>001</u>	Jun 06, 2022
<u>AA</u>			<u>40MEQ/15ML</u>	<u>A213062</u>	<u>002</u>	Jun 06, 2022

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE;ORAL

KLOR-CON M10

<u>AB1</u>	UPSHER SMITH LABS	<u>10MEO</u>	<u>A074726 002</u>	Aug 09, 2000
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KLOR-CON M15

<u>AB1</u>	UPSHER SMITH LABS	<u>15MEO</u>	<u>A074726 003</u>	Jun 06, 2003
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KLOR-CON M20

<u>AB1</u>	! UPSHER SMITH LABS	<u>20MEO</u>	<u>A074726 001</u>	Nov 20, 1998
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POTASSIUM CHLORIDE

<u>AB1</u>	ACTAVIS LABS FL INC	<u>10MEO</u>	<u>A075604 001</u>	Apr 10, 2002
<u>AB1</u>		<u>20MEO</u>	<u>A075604 002</u>	Apr 10, 2002
<u>AB1</u>	ADARE PHARMS INC	<u>10MEO</u>	<u>A076368 002</u>	Jun 05, 2019
<u>AB1</u>		<u>20MEO</u>	<u>A076368 001</u>	Aug 18, 2004
<u>AB1</u>	AMNEAL	<u>10MEO</u>	<u>A212861 001</u>	May 08, 2020
<u>AB1</u>		<u>20MEO</u>	<u>A212861 003</u>	May 08, 2020
<u>AB1</u>	ASCENT PHARMS INC	<u>10MEO</u>	<u>A214422 001</u>	Dec 29, 2020
<u>AB1</u>		<u>15MEO</u>	<u>A214422 002</u>	Dec 29, 2020
<u>AB1</u>		<u>20MEO</u>	<u>A214422 003</u>	Dec 29, 2020
<u>AB1</u>	GLENMARK PHARMS LTD	<u>10MEO</u>	<u>A203562 001</u>	Jul 26, 2016
<u>AB1</u>		<u>20MEO</u>	<u>A203562 002</u>	Jul 26, 2016
<u>AB1</u>	GRANULES	<u>10MEO</u>	<u>A214452 001</u>	Oct 21, 2020
<u>AB1</u>		<u>20MEO</u>	<u>A214452 002</u>	Oct 21, 2020
<u>AB1</u>	GUANGZHOU NOVAKEN	<u>10MEO</u>	<u>A214395 001</u>	Jan 28, 2021
<u>AB1</u>		<u>20MEO</u>	<u>A214395 002</u>	Jan 28, 2021
<u>AB1</u>	NOVEL LABS INC	<u>10MEO</u>	<u>A206347 001</u>	Jan 21, 2016
<u>AB1</u>		<u>20MEO</u>	<u>A206347 002</u>	Jan 21, 2016
<u>AB1</u>	PRINSTON INC	<u>10MEO</u>	<u>A209922 001</u>	Apr 30, 2019
<u>AB1</u>		<u>15MEO</u>	<u>A209922 002</u>	Apr 30, 2019
<u>AB1</u>		<u>20MEO</u>	<u>A209922 003</u>	Apr 30, 2019
<u>AB1</u>	RUBICON	<u>10MEO</u>	<u>A216321 001</u>	Jun 16, 2023
<u>AB1</u>		<u>15MEO</u>	<u>A216321 002</u>	Jun 16, 2023
<u>AB1</u>		<u>20MEO</u>	<u>A216321 003</u>	Jun 16, 2023
<u>AB1</u>	ZYDUS PHARMS	<u>10MEO</u>	<u>A210395 001</u>	Sep 17, 2020
<u>AB1</u>		<u>20MEO</u>	<u>A210395 002</u>	Sep 17, 2020

KLOR-CON

<u>AB2</u>	+ UPSHER SMITH LABS	<u>8MEO</u>	<u>N019123 001</u>	Apr 17, 1986
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<u>AB2</u>	+!	<u>10MEO</u>	<u>N019123 002</u>	Apr 17, 1986
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POTASSIUM CHLORIDE

<u>AB2</u>	AUROBINDO PHARMA	<u>8MEO</u>	<u>A210921 001</u>	Dec 19, 2018
<u>AB2</u>		<u>10MEO</u>	<u>A210921 002</u>	Dec 19, 2018
<u>AB2</u>	GRANULES	<u>8MEO</u>	<u>A211797 001</u>	Mar 04, 2020
<u>AB2</u>		<u>10MEO</u>	<u>A211797 002</u>	Mar 04, 2020
<u>AB2</u>	MYLAN	<u>8MEO</u>	<u>A204662 001</u>	Aug 21, 2014
<u>AB2</u>		<u>10MEO</u>	<u>A204662 002</u>	Aug 21, 2014
<u>AB2</u>	NOVEL LABS INC	<u>8MEO</u>	<u>A206759 001</u>	Aug 09, 2016
<u>AB2</u>		<u>10MEO</u>	<u>A206759 002</u>	Aug 09, 2016
<u>AB2</u>	PADAGIS US	<u>8MEO</u>	<u>A205993 001</u>	Nov 05, 2015
<u>AB2</u>		<u>10MEO</u>	<u>A205993 002</u>	Nov 05, 2015
<u>AB2</u>	RISING	<u>8MEO</u>	<u>A217412 001</u>	Dec 19, 2023
<u>AB2</u>		<u>10MEO</u>	<u>A217412 002</u>	Dec 19, 2023
<u>AB2</u>	STRIDES PHARMA	<u>8MEO</u>	<u>A210733 001</u>	Aug 31, 2018
<u>AB2</u>		<u>10MEO</u>	<u>A210733 002</u>	Aug 31, 2018
<u>AB2</u>	YICHANG HUMANWELL	<u>8MEO</u>	<u>A209314 001</u>	Jun 22, 2018
<u>AB2</u>		<u>10MEO</u>	<u>A209314 002</u>	Jun 22, 2018
<u>AB3</u>	RUBICON	<u>10MEO</u>	<u>A215725 001</u>	Jul 25, 2022
<u>AB3</u>		<u>20MEO</u>	<u>A215725 002</u>	Jul 25, 2022
<u>AB3</u>	TWI PHARMS	<u>10MEO</u>	<u>A209688 001</u>	Jan 12, 2018
<u>AB3</u>	!	<u>20MEO</u>	<u>A209688 002</u>	Jan 12, 2018
<u>AB3</u>	YICHANG HUMANWELL	<u>10MEO</u>	<u>A212561 001</u>	Sep 30, 2019
<u>AB3</u>		<u>20MEO</u>	<u>A212561 002</u>	Sep 30, 2019

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AP</u>	ICU MEDICAL INC	<u>149MG/100ML; 450MG/100ML</u>	<u>A078446 001</u>	Sep 10, 2008
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POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%

<u>AP</u>	+!	<u>150MG/100ML; 450MG/100ML</u>	<u>N017648 005</u>	Nov 26, 2002
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<u>AP</u>	FRESENIUS KABI USA	<u>150MG/100ML; 450MG/100ML</u>	<u>A212347 001</u>	Sep 17, 2020
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POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%

<u>AP</u>	+!	<u>150MG/100ML; 900MG/100ML</u>	<u>N017648 001</u>	
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<u>AP</u>	FRESENIUS KABI USA	<u>150MG/100ML; 900MG/100ML</u>	<u>A212347 003</u>	Jun 02, 2021
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PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>150MG/100ML;900MG/100ML</u>	<u>N019708 004</u>	Sep 29, 1989
<u>POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%</u>				
<u>AP</u>	+! BAXTER HLTHCARE	<u>300MG/100ML;900MG/100ML</u>	<u>N017648 002</u>	
<u>AP</u>	FRESENIUS KABI USA	<u>300MG/100ML;900MG/100ML</u>	<u>A212347 002</u>	Sep 17, 2020
<u>POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
<u>AP</u>	ICU MEDICAL INC	<u>149MG/100ML;900MG/100ML</u>	<u>N019686 001</u>	Oct 17, 1988
<u>POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
<u>AP</u>	ICU MEDICAL INC	<u>298MG/100ML;900MG/100ML</u>	<u>N019686 002</u>	Oct 17, 1988

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CITRATE

<u>AB</u>	ANI PHARMS	<u>10MEQ</u>	<u>A212779 001</u>	Jan 14, 2020
<u>AB</u>		<u>15MEQ</u>	<u>A212779 002</u>	Jan 14, 2020
<u>AB</u>	ASCENT PHARMS INC	<u>5MEQ</u>	<u>A214420 001</u>	Feb 05, 2021
<u>AB</u>		<u>10MEQ</u>	<u>A214420 002</u>	Feb 05, 2021
<u>AB</u>		<u>15MEQ</u>	<u>A214420 003</u>	Feb 05, 2021
<u>AB</u>	BIONPHARMA	<u>10MEQ</u>	<u>A212799 001</u>	Jun 29, 2020
<u>AB</u>		<u>15MEQ</u>	<u>A212799 002</u>	Jun 29, 2020
<u>AB</u>	EYWA	<u>5MEQ</u>	<u>A214426 001</u>	Feb 19, 2021
<u>AB</u>		<u>10MEQ</u>	<u>A214426 002</u>	Feb 19, 2021
<u>AB</u>		<u>15MEQ</u>	<u>A214426 003</u>	Feb 19, 2021
<u>AB</u>	RISING	<u>5MEQ</u>	<u>A077440 001</u>	Jun 09, 2006
<u>AB</u>		<u>10MEQ</u>	<u>A077440 002</u>	Jun 09, 2006
<u>AB</u>	STRIDES PHARMA	<u>5MEQ</u>	<u>A206813 001</u>	Sep 11, 2017
<u>AB</u>		<u>10MEQ</u>	<u>A206813 002</u>	Sep 11, 2017
<u>AB</u>		<u>15MEQ</u>	<u>A206813 003</u>	Sep 11, 2017
<u>AB</u>	TEVA PHARMS USA INC	<u>5MEQ</u>	<u>A209758 001</u>	Mar 05, 2018
<u>AB</u>		<u>10MEQ</u>	<u>A209758 002</u>	Mar 05, 2018
<u>AB</u>		<u>15MEQ</u>	<u>A209758 003</u>	Mar 05, 2018
<u>AB</u>	ZYDUS PHARMS	<u>5MEQ</u>	<u>A203546 001</u>	Aug 06, 2014
<u>AB</u>		<u>10MEQ</u>	<u>A203546 002</u>	Aug 06, 2014
<u>AB</u>		<u>15MEQ</u>	<u>A203546 003</u>	Aug 06, 2014

UROCIIT-K

<u>AB</u>	+ MISSION PHARMA	<u>5MEQ</u>	<u>N019071 001</u>	Aug 30, 1985
<u>AB</u>	+	<u>10MEQ</u>	<u>N019071 002</u>	Aug 31, 1992
<u>AB</u>	+!	<u>15MEQ</u>	<u>N019071 003</u>	Dec 30, 2009

POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC

SOLUTION; INTRAVENOUS

POTASSIUM PHOSPHATES

<u>AP</u>	AM REGENT	<u>1.18GM/5ML (236MG/ML); 1.12GM/5ML (224MG/ML)</u>	<u>A216274 001</u>	Oct 02, 2023
<u>AP</u>		<u>3.54GM/15ML (236MG/ML); 3.36GM/15ML (224MG/ML)</u>	<u>A216274 002</u>	Oct 02, 2023
<u>AP</u>		<u>11.8GM/50ML (236MG/ML); 11.2GM/50ML (224MG/ML)</u>	<u>A216274 003</u>	Oct 02, 2023
<u>AP</u>	AMNEAL	<u>1.18GM/5ML (236MG/ML); 1.12GM/5ML (224MG/ML)</u>	<u>A216344 001</u>	Oct 10, 2023
<u>AP</u>		<u>3.54GM/15ML (236MG/ML); 3.36GM/15ML (224MG/ML)</u>	<u>A216344 002</u>	Oct 10, 2023
<u>AP</u>		<u>11.8GM/50ML (236MG/ML); 11.2GM/50ML (224MG/ML)</u>	<u>A216344 003</u>	Oct 10, 2023
<u>AP</u>	+! FRESENIUS KABI USA	<u>1.18GM/5ML (236MG/ML); 1.12GM/5ML (224MG/ML)</u>	<u>N212832 001</u>	Nov 26, 2019
<u>AP</u>	+	<u>3.54GM/15ML (236MG/ML); 3.36GM/15ML (224MG/ML)</u>	<u>N212832 002</u>	Nov 26, 2019
<u>AP</u>	+	<u>11.8GM/50ML (236MG/ML); 11.2GM/50ML (224MG/ML)</u>	<u>N212832 003</u>	Nov 26, 2019
	+! CMP DEV LLC	<u>4.5GM/15ML (300MG/ML); 2.65GM/15ML (175MG/ML)</u>	<u>N212121 001</u>	Sep 19, 2019

POVIDONE-IODINE

SOLUTION/DROPS; OPHTHALMIC

BETADINE

+	ALCON PHARMS LTD	5%	<u>N018634 001</u>	Dec 17, 1986
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PRESCRIPTION DRUG PRODUCT LIST

PRALATREXATE

SOLUTION; INTRAVENOUS

FOLOTYN

+	ACROTECH BIOPHARMA	20MG/ML (20MG/ML)	N022468	001	Sep 24, 2009
+	!	40MG/2ML (20MG/ML)	N022468	002	Sep 24, 2009

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PROTOPAM CHLORIDE

+	BAXTER HLTHCARE CORP	1GM/VIAL	N014134	001	
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PRALSETINIB

CAPSULE; ORAL

GAVRETO

+	GENENTECH INC	100MG	N213721	001	Sep 04, 2020
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PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.125MG</u>	<u>A202633</u>	<u>001</u>	Oct 26, 2012
<u>AB</u>		<u>0.25MG</u>	<u>A202633</u>	<u>002</u>	Oct 26, 2012
<u>AB</u>		<u>0.5MG</u>	<u>A202633</u>	<u>003</u>	Oct 26, 2012
<u>AB</u>		<u>0.75MG</u>	<u>A202633</u>	<u>004</u>	Oct 26, 2012
<u>AB</u>		<u>1MG</u>	<u>A202633</u>	<u>005</u>	Oct 26, 2012
<u>AB</u>		<u>1.5MG</u>	<u>A202633</u>	<u>006</u>	Oct 26, 2012
<u>AB</u>	GLENMARK GENERICS	<u>0.125MG</u>	<u>A090781</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>	!	<u>0.25MG</u>	<u>A090781</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090781</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090781</u>	<u>006</u>	Sep 11, 2015
<u>AB</u>		<u>1MG</u>	<u>A090781</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A090781</u>	<u>005</u>	Oct 08, 2010
<u>AB</u>	RISING	<u>0.25MG</u>	<u>A211088</u>	<u>001</u>	Oct 03, 2018
<u>AB</u>		<u>0.5MG</u>	<u>A211088</u>	<u>002</u>	Oct 03, 2018
<u>AB</u>		<u>0.75MG</u>	<u>A211088</u>	<u>003</u>	Oct 03, 2018
<u>AB</u>		<u>1MG</u>	<u>A211088</u>	<u>004</u>	Oct 03, 2018
<u>AB</u>		<u>1.5MG</u>	<u>A211088</u>	<u>005</u>	Oct 03, 2018
<u>AB</u>	SCIEGEN PHARMS INC	<u>0.125MG</u>	<u>A203855</u>	<u>001</u>	Oct 28, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A203855</u>	<u>002</u>	Oct 28, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A203855</u>	<u>003</u>	Oct 28, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A203855</u>	<u>004</u>	Oct 28, 2014
<u>AB</u>		<u>1MG</u>	<u>A203855</u>	<u>005</u>	Oct 28, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A203855</u>	<u>006</u>	Oct 28, 2014
<u>AB</u>	STRIDES PHARMA	<u>0.125MG</u>	<u>A202702</u>	<u>001</u>	Jun 03, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A202702</u>	<u>002</u>	Jun 03, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A202702</u>	<u>003</u>	Jun 03, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A202702</u>	<u>004</u>	Jun 03, 2014
<u>AB</u>		<u>1MG</u>	<u>A202702</u>	<u>005</u>	Jun 03, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A202702</u>	<u>006</u>	Jun 03, 2014
<u>AB</u>	TORRENT PHARMS	<u>0.125MG</u>	<u>A090865</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A090865</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090865</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090865</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		<u>1MG</u>	<u>A090865</u>	<u>005</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A090865</u>	<u>006</u>	Oct 08, 2010
<u>AB</u>	ZENNOVA	<u>0.125MG</u>	<u>A090151</u>	<u>001</u>	Apr 30, 2012
<u>AB</u>		<u>0.25MG</u>	<u>A090151</u>	<u>002</u>	Apr 30, 2012
<u>AB</u>		<u>0.5MG</u>	<u>A090151</u>	<u>003</u>	Apr 30, 2012
<u>AB</u>		<u>0.75MG</u>	<u>A090151</u>	<u>006</u>	Apr 30, 2012
<u>AB</u>		<u>1MG</u>	<u>A090151</u>	<u>004</u>	Apr 30, 2012
<u>AB</u>		<u>1.5MG</u>	<u>A090151</u>	<u>005</u>	Apr 30, 2012
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.125MG</u>	<u>A078920</u>	<u>001</u>	Jul 06, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A078920</u>	<u>002</u>	Jul 06, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078920</u>	<u>003</u>	Jul 06, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A078920</u>	<u>006</u>	Nov 23, 2022
<u>AB</u>		<u>1MG</u>	<u>A078920</u>	<u>004</u>	Jul 06, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A078920</u>	<u>005</u>	Jul 06, 2010

TABLET, EXTENDED RELEASE; ORAL

MIRAPEX ER

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>0.375MG</u>	<u>N022421</u>	<u>001</u>	Feb 19, 2010
<u>AB</u>	+		<u>0.75MG</u>	<u>N022421</u>	<u>002</u>	Feb 19, 2010
<u>AB</u>	+		<u>1.5MG</u>	<u>N022421</u>	<u>003</u>	Feb 19, 2010

PRESCRIPTION DRUG PRODUCT LIST

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

MIRAPEX ER

<u>AB</u>	+	<u>2.25MG</u>	<u>N022421</u>	<u>006</u>	Jun 17, 2011
<u>AB</u>	+	<u>3MG</u>	<u>N022421</u>	<u>004</u>	Feb 19, 2010
<u>AB</u>	+	<u>3.75MG</u>	<u>N022421</u>	<u>007</u>	Jun 17, 2011
<u>AB</u>	+	<u>4.5MG</u>	<u>N022421</u>	<u>005</u>	Feb 19, 2010

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.375MG</u>	<u>A201963</u>	<u>001</u>	Apr 21, 2016
<u>AB</u>		<u>0.75MG</u>	<u>A201963</u>	<u>002</u>	Apr 21, 2016
<u>AB</u>		<u>1.5MG</u>	<u>A201963</u>	<u>003</u>	Apr 21, 2016
<u>AB</u>		<u>2.25MG</u>	<u>A203615</u>	<u>001</u>	Oct 14, 2016
<u>AB</u>		<u>3MG</u>	<u>A201963</u>	<u>004</u>	Apr 21, 2016
<u>AB</u>		<u>3.75MG</u>	<u>A203615</u>	<u>002</u>	Jan 03, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A201963</u>	<u>005</u>	Apr 21, 2016
<u>AB</u>	ALEMBIC	<u>0.375MG</u>	<u>A204518</u>	<u>001</u>	Jan 02, 2019
<u>AB</u>		<u>0.75MG</u>	<u>A204518</u>	<u>002</u>	Jan 02, 2019
<u>AB</u>		<u>1.5MG</u>	<u>A204518</u>	<u>003</u>	Jan 02, 2019
<u>AB</u>		<u>2.25MG</u>	<u>A204518</u>	<u>004</u>	Jan 02, 2019
<u>AB</u>		<u>3MG</u>	<u>A204518</u>	<u>005</u>	Jan 02, 2019
<u>AB</u>		<u>3.75MG</u>	<u>A204518</u>	<u>006</u>	Jan 02, 2019
<u>AB</u>		<u>4.5MG</u>	<u>A204518</u>	<u>007</u>	Jan 02, 2019
<u>AB</u>	DR REDDYS	<u>0.375MG</u>	<u>A203354</u>	<u>001</u>	Aug 07, 2015
<u>AB</u>		<u>0.75MG</u>	<u>A203354</u>	<u>002</u>	Aug 07, 2015
<u>AB</u>		<u>1.5MG</u>	<u>A203354</u>	<u>003</u>	Aug 07, 2015
<u>AB</u>		<u>3MG</u>	<u>A203354</u>	<u>004</u>	Aug 07, 2015
<u>AB</u>		<u>4.5MG</u>	<u>A203354</u>	<u>005</u>	Aug 07, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>0.375MG</u>	<u>A206156</u>	<u>001</u>	Jun 24, 2016
<u>AB</u>		<u>0.75MG</u>	<u>A206156</u>	<u>002</u>	Jun 24, 2016
<u>AB</u>		<u>1.5MG</u>	<u>A206156</u>	<u>003</u>	Jun 24, 2016
<u>AB</u>		<u>2.25MG</u>	<u>A206156</u>	<u>004</u>	Jun 24, 2016
<u>AB</u>		<u>3MG</u>	<u>A206156</u>	<u>005</u>	Jun 24, 2016
<u>AB</u>		<u>3.75MG</u>	<u>A206156</u>	<u>007</u>	Jan 23, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A206156</u>	<u>006</u>	Jun 24, 2016
<u>AB</u>	NOVAST LABS	<u>0.375MG</u>	<u>A213444</u>	<u>001</u>	Feb 03, 2022
<u>AB</u>		<u>0.75MG</u>	<u>A213444</u>	<u>002</u>	Feb 03, 2022
<u>AB</u>		<u>1.5MG</u>	<u>A213444</u>	<u>003</u>	Feb 03, 2022
<u>AB</u>		<u>2.25MG</u>	<u>A213444</u>	<u>004</u>	Feb 03, 2022
<u>AB</u>		<u>3MG</u>	<u>A213444</u>	<u>005</u>	Feb 03, 2022
<u>AB</u>		<u>3.75MG</u>	<u>A213444</u>	<u>006</u>	Feb 03, 2022
<u>AB</u>		<u>4.5MG</u>	<u>A213444</u>	<u>007</u>	Feb 03, 2022
<u>AB</u>	SANDOZ	<u>0.375MG</u>	<u>A202353</u>	<u>001</u>	Dec 04, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A202353</u>	<u>002</u>	Dec 04, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A202353</u>	<u>003</u>	Dec 04, 2014
<u>AB</u>		<u>3MG</u>	<u>A202353</u>	<u>004</u>	Dec 04, 2014
<u>AB</u>		<u>4.5MG</u>	<u>A202353</u>	<u>005</u>	Dec 04, 2014
<u>AB</u>	XIAMEN LP PHARM CO	<u>0.375MG</u>	<u>A212797</u>	<u>001</u>	Jun 11, 2021
<u>AB</u>		<u>0.75MG</u>	<u>A212797</u>	<u>002</u>	Jun 11, 2021
<u>AB</u>	ZYDUS PHARMS	<u>0.375MG</u>	<u>A202891</u>	<u>001</u>	Dec 12, 2017
<u>AB</u>		<u>0.75MG</u>	<u>A202891</u>	<u>002</u>	Dec 12, 2017
<u>AB</u>		<u>1.5MG</u>	<u>A202891</u>	<u>003</u>	Dec 12, 2017
<u>AB</u>		<u>2.25MG</u>	<u>A202891</u>	<u>004</u>	Dec 12, 2017
<u>AB</u>		<u>3MG</u>	<u>A202891</u>	<u>005</u>	Dec 12, 2017
<u>AB</u>		<u>3.75MG</u>	<u>A202891</u>	<u>006</u>	Dec 12, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A202891</u>	<u>007</u>	Dec 12, 2017

PRAMLINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS

SYMLIN

+	ASTRAZENECA AB	EQ 1.5MG BASE/1.5ML (EQ 1MG BASE/ML)	N021332	002	Sep 25, 2007
+	!	EQ 2.7MG BASE/2.7ML (EQ 1MG BASE/ML)	N021332	003	Sep 25, 2007

PRASTERONE

INSERT; VAGINAL

INTRAROSA

+	!	MILLICENT	6.5MG	N208470	001	Nov 16, 2016
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PRASUGREL HYDROCHLORIDE

TABLET; ORAL

EFFIENT

<u>AB</u>	+	COSETTE	<u>EQ 5MG BASE</u>	<u>N022307</u>	<u>001</u>	Jul 10, 2009
<u>AB</u>	+	!	<u>EQ 10MG BASE</u>	<u>N022307</u>	<u>002</u>	Jul 10, 2009

PRESCRIPTION DRUG PRODUCT LIST

PRASUGREL HYDROCHLORIDE

TABLET; ORAL

PRASUGREL

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A205987 001</u>	Feb 02, 2018
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205987 002</u>	Feb 02, 2018
<u>AB</u>	AMNEAL PHARMS	<u>EQ 5MG BASE</u>	<u>A205913 001</u>	Jun 19, 2018
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205913 002</u>	Jun 19, 2018
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 5MG BASE</u>	<u>A205888 001</u>	Oct 16, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205888 002</u>	Oct 16, 2017
<u>AB</u>	HEC PHARM	<u>EQ 5MG BASE</u>	<u>A206021 001</u>	Jan 16, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206021 002</u>	Jan 16, 2019
<u>AB</u>	MYLAN	<u>EQ 5MG BASE</u>	<u>A205927 001</u>	Jul 12, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205927 002</u>	Jul 12, 2017
<u>AB</u>	PANACEA	<u>EQ 5MG BASE</u>	<u>A205897 001</u>	Oct 16, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205897 002</u>	Oct 16, 2017

PRASUGREL HYDROCHLORIDE

<u>AB</u>	UNICHEM	<u>EQ 5MG BASE</u>	<u>A213315 001</u>	Aug 28, 2023
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A213315 002</u>	Aug 28, 2023

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVASTATIN SODIUM

<u>AB</u>	ACCORD HLTHCARE	<u>10MG</u>	<u>A207068 001</u>	Nov 17, 2016
<u>AB</u>		<u>20MG</u>	<u>A207068 002</u>	Nov 17, 2016
<u>AB</u>		<u>40MG</u>	<u>A207068 003</u>	Nov 17, 2016
<u>AB</u>		<u>80MG</u>	<u>A207068 004</u>	Nov 17, 2016
<u>AB</u>	APNAR PHARMA LP	<u>10MG</u>	<u>A077491 002</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A077491 003</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A077491 004</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A077491 001</u>	Feb 11, 2008
<u>AB</u>	APOTEX	<u>10MG</u>	<u>A076341 001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076341 002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076341 003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076341 004</u>	Dec 28, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A203367 001</u>	Feb 02, 2017
<u>AB</u>		<u>20MG</u>	<u>A203367 002</u>	Feb 02, 2017
<u>AB</u>		<u>40MG</u>	<u>A203367 003</u>	Feb 02, 2017
<u>AB</u>		<u>80MG</u>	<u>A203367 004</u>	Feb 02, 2017
<u>AB</u>	BIOCON PHARMA	<u>10MG</u>	<u>A209869 001</u>	Apr 13, 2018
<u>AB</u>		<u>20MG</u>	<u>A209869 002</u>	Apr 13, 2018
<u>AB</u>		<u>40MG</u>	<u>A209869 003</u>	Apr 13, 2018
<u>AB</u>		<u>80MG</u>	<u>A209869 004</u>	Apr 13, 2018
<u>AB</u>	CHARTWELL RX	<u>10MG</u>	<u>A077917 001</u>	Jan 08, 2008
<u>AB</u>		<u>20MG</u>	<u>A077917 002</u>	Jan 08, 2008
<u>AB</u>		<u>40MG</u>	<u>A077917 003</u>	Jan 08, 2008
<u>AB</u>		<u>80MG</u>	<u>A077917 004</u>	Jan 08, 2008
<u>AB</u>	CIPLA	<u>10MG</u>	<u>A077904 001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A077904 002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A077904 003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A077904 004</u>	Mar 22, 2016
<u>AB</u>	DR REDDYS LABS INC	<u>10MG</u>	<u>A076714 001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076714 002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076714 003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076714 004</u>	Dec 28, 2007
<u>AB</u>	GLENMARK GENERICS	<u>10MG</u>	<u>A077987 001</u>	May 11, 2007
<u>AB</u>		<u>20MG</u>	<u>A077987 002</u>	May 11, 2007
<u>AB</u>		<u>40MG</u>	<u>A077987 003</u>	May 11, 2007
<u>AB</u>		<u>80MG</u>	<u>A077987 004</u>	Dec 28, 2007
<u>AB</u>	TEVA	<u>10MG</u>	<u>A076056 001</u>	Apr 24, 2006
<u>AB</u>		<u>20MG</u>	<u>A076056 002</u>	Apr 24, 2006
<u>AB</u>		<u>40MG</u>	<u>A076056 003</u>	Apr 24, 2006
<u>AB</u>	! TEVA PHARMS	<u>80MG</u>	<u>A077793 001</u>	Jan 15, 2008
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A076939 004</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076939 003</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076939 002</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076939 001</u>	Dec 28, 2007
<u>AB</u>	ZYDUS PHARMS USA	<u>20MG</u>	<u>A077751 002</u>	Apr 30, 2008
<u>AB</u>		<u>40MG</u>	<u>A077751 003</u>	Apr 30, 2008
<u>AB</u>		<u>80MG</u>	<u>A077751 004</u>	Apr 30, 2008

PRESCRIPTION DRUG PRODUCT LIST

PRAZIOUANTEL

TABLET; ORAL

BILTRICIDE

AB	+	BAYER HLTHCARE	600MG	N018714	001	Dec 29, 1982
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PRAZIOUANTEL

AB		PAR PHARM INC	600MG	A208820	001	Nov 27, 2017
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PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIPRESS

AB	+	PFIZER	EQ 1MG BASE	N017442	002	
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AB	+		EQ 2MG BASE	N017442	003	
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AB	+		EQ 5MG BASE	N017442	001	
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PRAZOSIN HYDROCHLORIDE

AB		ALEMBIC	EQ 1MG BASE	A217268	001	Mar 06, 2023
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AB			EQ 2MG BASE	A217268	002	Mar 06, 2023
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AB			EQ 5MG BASE	A217268	003	Mar 06, 2023
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AB		APPCO	EQ 1MG BASE	A213406	001	Oct 21, 2022
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AB			EQ 2MG BASE	A213406	002	Oct 21, 2022
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AB			EQ 5MG BASE	A213406	003	Oct 21, 2022
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AB		AUROBINDO PHARMA LTD	EQ 1MG BASE	A213052	001	Mar 31, 2023
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AB			EQ 2MG BASE	A213052	002	Mar 31, 2023
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AB			EQ 5MG BASE	A213052	003	Mar 31, 2023
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AB		GRANULES	EQ 1MG BASE	A214608	001	Dec 23, 2021
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AB			EQ 2MG BASE	A214608	002	Dec 23, 2021
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AB			EQ 5MG BASE	A214608	003	Dec 23, 2021
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AB		LANNETT CO INC	EQ 1MG BASE	A214083	001	Jan 03, 2024
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AB			EQ 2MG BASE	A214083	002	Jan 03, 2024
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AB			EQ 5MG BASE	A214083	003	Jan 03, 2024
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AB		MANKIND PHARMA	EQ 1MG BASE	A215697	001	Dec 30, 2022
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AB			EQ 2MG BASE	A215697	002	Dec 30, 2022
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AB			EQ 5MG BASE	A215697	003	Dec 30, 2022
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AB		MYLAN	EQ 1MG BASE	A072575	003	May 16, 1989
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AB			EQ 2MG BASE	A072575	002	May 16, 1989
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AB			EQ 5MG BASE	A072575	001	May 16, 1989
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AB		NOVITIUM PHARMA	EQ 1MG BASE	A210971	001	Oct 03, 2018
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AB			EQ 2MG BASE	A210971	002	Oct 03, 2018
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AB			EQ 5MG BASE	A210971	003	Oct 03, 2018
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AB		TEVA PHARMS	EQ 1MG BASE	A071745	002	Sep 12, 1988
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AB			EQ 2MG BASE	A071745	003	Sep 12, 1988
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AB			EQ 5MG BASE	A071745	001	Sep 12, 1988
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PREDNICARBATE

OINTMENT; TOPICAL

PREDNICARBATE

!	FOUGERA PHARMS	0.1%	A077236	001	Mar 09, 2007
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PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

AA	!	CHARTWELL RX	15MG/5ML	A040323	001	May 13, 1999
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AA		HIKMA	15MG/5ML	A040401	001	Feb 27, 2003
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AA		LANNETT CO INC	15MG/5ML	A040775	001	Sep 21, 2007
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AA		PHARM ASSOC	15MG/5ML	A040399	001	Mar 05, 2003
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AA			15MG/5ML	A040571	001	Aug 25, 2005
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PRELONE

AA		TEVA	15MG/5ML	A089081	001	Feb 04, 1986
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TABLET; ORAL

PREDNISOLONE

AB		AUROBINDO PHARMA LTD	5MG	A215673	001	Mar 17, 2023
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AB	+	WATSON LABS	5MG	A080354	001	
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BX		CHARTWELL MOLECULAR	5MG	A080531	002	
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PREDNISOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

OMNIPRED

AB	+	SANDOZ	1%	N017469	001	
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PRED FORTE

AB	+	ABEVIE	1%	N017011	001	
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PRED MILD

+	ABEVIE	0.12%	N017100	001	
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PRESCRIPTION DRUG PRODUCT LIST

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION;ORAL

PEDIAAPRED

AA	+ !	SETON PHARM	EQ 5MG BASE/5ML	N019157 001	May 28, 1986
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PREDNISOLONE SODIUM PHOSPHATE

AA		AMNEAL	EQ 15MG BASE/5ML	A216715 001	Oct 25, 2022
AA		CHARTWELL RX	EQ 5MG BASE/5ML	A075988 001	May 25, 2004
AA		EDENBRIDGE PHARMS	EQ 10MG BASE/5ML	A203559 001	Dec 20, 2016
AA			EQ 15MG BASE/5ML	A203559 003	Feb 06, 2023
AA			EQ 20MG BASE/5ML	A203559 002	Dec 20, 2016
AA			EQ 25MG BASE/5ML	A203559 004	Feb 06, 2023
AA		HIKMA	EQ 5MG BASE/5ML	A075183 001	Mar 26, 2003
AA	!	MISSION PHARMA	EQ 25MG BASE/5ML	A091396 001	Sep 13, 2010
AA	!	PHARM ASSOC	EQ 10MG BASE/5ML	A078465 001	Mar 07, 2008
AA	!		EQ 15MG BASE/5ML	A076913 001	Apr 25, 2005
AA	!		EQ 20MG BASE/5ML	A078988 001	Jun 09, 2008
			EQ 30MG BASE/5ML	A204962 001	Mar 11, 2020

SOLUTION/DROPS;OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

!	BAUSCH AND LOMB	EQ 0.9% PHOSPHATE	A040070 001	Jul 29, 1994
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TABLET, ORALLY DISINTEGRATING;ORAL

ORAPRED ODT

+	CONCORDIA PHARMS INC	EQ 10MG BASE	N021959 001	Jun 01, 2006
+		EQ 15MG BASE	N021959 002	Jun 01, 2006
+ !		EQ 30MG BASE	N021959 003	Jun 01, 2006

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

!	BAUSCH AND LOMB	EQ 0.23% PHOSPHATE;10%	A074449 001	Dec 29, 1995
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PREDNISONE

SOLUTION;ORAL

PREDNISONE

+ !	HIKMA	5MG/5ML	A088703 001	Nov 08, 1984
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PREDNISONE INTENSOL

!	HIKMA	5MG/ML	A088810 001	Feb 20, 1985
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TABLET;ORAL

PREDNISONE

AB		AMNEAL	1MG	A213385 001	Jun 16, 2020
AB			5MG	A213385 002	Jun 16, 2020
AB			10MG	A213386 001	Jun 24, 2020
AB			20MG	A213386 002	Jun 24, 2020
AB		AUROBINDO PHARMA LTD	1MG	A215671 001	Nov 16, 2021
AB			2.5MG	A215672 001	Mar 28, 2022
AB			5MG	A215672 002	Mar 28, 2022
AB			10MG	A215672 003	Mar 28, 2022
AB			20MG	A215672 004	Mar 28, 2022
AB			50MG	A215672 005	Mar 28, 2022
AB		GENEYORK PHARMS	1MG	A211496 001	Dec 28, 2018
AB			2.5MG	A210525 004	Dec 07, 2018
AB			5MG	A210525 005	Dec 07, 2018
AB			10MG	A210525 001	Dec 04, 2018
AB			20MG	A210525 002	Dec 04, 2018
AB			50MG	A210525 003	Dec 04, 2018
AB		GRANULATION TECH	5MG	A212629 001	Dec 05, 2023
AB			10MG	A212629 002	Dec 05, 2023
AB			20MG	A212629 003	Dec 05, 2023
AB	+ !	HIKMA	1MG	A087800 001	Apr 22, 1982
AB	+ !		2.5MG	A087801 001	Apr 22, 1982
AB	+ !		5MG	A080352 001	
AB	+ !		10MG	A084122 001	
AB	+ !		20MG	A087342 001	
AB	+ !		50MG	A084283 001	
AB		JUBILANT CADISTA	1MG	A040611 001	Jun 06, 2005
AB			2.5MG	A040362 004	Apr 17, 2023
AB			5MG	A040362 002	Aug 29, 2001
AB			10MG	A040362 001	Aug 29, 2001
AB			20MG	A040362 003	Jun 29, 2005
AB			50MG	A040362 005	Apr 17, 2023
AB		MYLAN	5MG	A080292 001	
AB			10MG	A088832 001	Dec 04, 1985

PRESCRIPTION DRUG PRODUCT LIST

PREDNISONE

TABLET; ORAL

PREDNISONE

<u>AB</u>		<u>20MG</u>	<u>A083677 001</u>	
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A215246 001</u>	Jul 06, 2021
<u>AB</u>		<u>2.5MG</u>	<u>A211575 001</u>	Nov 15, 2019
<u>AB</u>		<u>5MG</u>	<u>A211575 002</u>	Nov 15, 2019
<u>AB</u>		<u>10MG</u>	<u>A211575 003</u>	Nov 15, 2019
<u>AB</u>		<u>20MG</u>	<u>A211575 004</u>	Nov 15, 2019
<u>AB</u>		<u>50MG</u>	<u>A211575 005</u>	Nov 15, 2019
<u>AB</u>	STRIDES PHARMA	<u>1MG</u>	<u>A210785 001</u>	Sep 02, 2020
<u>AB</u>		<u>2.5MG</u>	<u>A209727 001</u>	Nov 20, 2020
<u>AB</u>		<u>5MG</u>	<u>A209727 002</u>	Nov 20, 2020
<u>AB</u>		<u>10MG</u>	<u>A208412 001</u>	Feb 11, 2021
<u>AB</u>		<u>20MG</u>	<u>A208412 002</u>	Feb 11, 2021
<u>AB</u>		<u>50MG</u>	<u>A208412 003</u>	Jan 11, 2022
<u>AB</u>	SUN PHARM INDUSTRIES	<u>5MG</u>	<u>A089247 002</u>	Dec 04, 1985
<u>AB</u>		<u>10MG</u>	<u>A089247 003</u>	Dec 04, 1985
<u>AB</u>		<u>20MG</u>	<u>A089247 001</u>	Dec 04, 1985
<u>AB</u>	VINTAGE PHARMS	<u>1MG</u>	<u>A040584 001</u>	Dec 21, 2004
<u>AB</u>		<u>2.5MG</u>	<u>A040581 001</u>	Dec 21, 2004
<u>AB</u>		<u>5MG</u>	<u>A040256 001</u>	Jul 12, 2002
<u>AB</u>		<u>10MG</u>	<u>A040256 002</u>	Jul 12, 2002
<u>AB</u>		<u>20MG</u>	<u>A040392 001</u>	Feb 12, 2003
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>A080356 001</u>	
<u>AB</u>		<u>10MG</u>	<u>A085162 001</u>	
<u>AB</u>		<u>20MG</u>	<u>A085161 001</u>	

TABLET, DELAYED RELEASE; ORAL

PREDNISONE

<u>AB</u>	ACTAVIS LABS FL INC	<u>1MG</u>	<u>A204867 001</u>	Apr 25, 2017
<u>AB</u>		<u>2MG</u>	<u>A204867 002</u>	Apr 25, 2017
<u>AB</u>		<u>5MG</u>	<u>A204867 003</u>	Apr 25, 2017

RAYOS

<u>AB</u>	+	HORIZON	<u>1MG</u>	<u>N202020 001</u>	Jul 26, 2012
<u>AB</u>	+		<u>2MG</u>	<u>N202020 002</u>	Jul 26, 2012
<u>AB</u>	+	!	<u>5MG</u>	<u>N202020 003</u>	Jul 26, 2012

PREGABALIN

CAPSULE; ORAL

LYRICA

<u>AB</u>	+	UPJOHN	<u>25MG</u>	<u>N021446 001</u>	Dec 30, 2004
<u>AB</u>	+		<u>50MG</u>	<u>N021446 002</u>	Dec 30, 2004
<u>AB</u>	+		<u>75MG</u>	<u>N021446 003</u>	Dec 30, 2004
<u>AB</u>	+		<u>100MG</u>	<u>N021446 004</u>	Dec 30, 2004
<u>AB</u>	+		<u>150MG</u>	<u>N021446 005</u>	Dec 30, 2004
<u>AB</u>	+		<u>200MG</u>	<u>N021446 006</u>	Dec 30, 2004
<u>AB</u>	+		<u>225MG</u>	<u>N021446 007</u>	Dec 30, 2004
<u>AB</u>	+	!	<u>300MG</u>	<u>N021446 008</u>	Dec 30, 2004

PREGABALIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>25MG</u>	<u>A091025 001</u>	Jul 09, 2020
<u>AB</u>		<u>50MG</u>	<u>A091025 002</u>	Jul 09, 2020
<u>AB</u>		<u>75MG</u>	<u>A091025 003</u>	Jul 09, 2020
<u>AB</u>		<u>100MG</u>	<u>A091025 004</u>	Jul 09, 2020
<u>AB</u>		<u>150MG</u>	<u>A091025 005</u>	Jul 09, 2020
<u>AB</u>		<u>200MG</u>	<u>A091025 006</u>	Jul 09, 2020
<u>AB</u>		<u>225MG</u>	<u>A091025 007</u>	Jul 09, 2020
<u>AB</u>		<u>300MG</u>	<u>A091025 008</u>	Jul 09, 2020
<u>AB</u>	ADAPTIS	<u>25MG</u>	<u>A216197 001</u>	Jul 18, 2022
<u>AB</u>		<u>50MG</u>	<u>A216197 002</u>	Jul 18, 2022
<u>AB</u>		<u>75MG</u>	<u>A216197 003</u>	Jul 18, 2022
<u>AB</u>		<u>100MG</u>	<u>A216197 004</u>	Jul 18, 2022
<u>AB</u>		<u>150MG</u>	<u>A216197 005</u>	Jul 18, 2022
<u>AB</u>		<u>200MG</u>	<u>A216197 006</u>	Jul 18, 2022
<u>AB</u>		<u>225MG</u>	<u>A216197 007</u>	Jul 18, 2022
<u>AB</u>		<u>300MG</u>	<u>A216197 008</u>	Jul 18, 2022
<u>AB</u>	ALEMBIC	<u>25MG</u>	<u>A203459 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A203459 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A203459 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A203459 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A203459 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A203459 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A203459 007</u>	Jul 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<u>AB</u>		<u>300MG</u>	<u>A203459</u>	<u>008</u>	Jul 19, 2019
<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A207799</u>	<u>007</u>	Sep 30, 2019
<u>AB</u>		<u>50MG</u>	<u>A207799</u>	<u>008</u>	Sep 30, 2019
<u>AB</u>		<u>75MG</u>	<u>A207799</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A207799</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A207799</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A207799</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A207799</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A207799</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>	AMNEAL PHARMS CO	<u>25MG</u>	<u>A209743</u>	<u>001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209743</u>	<u>002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209743</u>	<u>003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209743</u>	<u>004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209743</u>	<u>005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209743</u>	<u>006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209743</u>	<u>007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209743</u>	<u>008</u>	Jul 19, 2019
<u>AB</u>	APOTEX	<u>25MG</u>	<u>A211685</u>	<u>001</u>	Jul 07, 2021
<u>AB</u>		<u>50MG</u>	<u>A211685</u>	<u>002</u>	Jul 07, 2021
<u>AB</u>		<u>75MG</u>	<u>A211685</u>	<u>003</u>	Jul 07, 2021
<u>AB</u>		<u>100MG</u>	<u>A211685</u>	<u>004</u>	Jul 07, 2021
<u>AB</u>		<u>150MG</u>	<u>A211685</u>	<u>005</u>	Jul 07, 2021
<u>AB</u>		<u>200MG</u>	<u>A211685</u>	<u>006</u>	Jul 07, 2021
<u>AB</u>		<u>225MG</u>	<u>A211685</u>	<u>007</u>	Jul 07, 2021
<u>AB</u>		<u>300MG</u>	<u>A211685</u>	<u>008</u>	Jul 07, 2021
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A205321</u>	<u>001</u>	Mar 29, 2023
<u>AB</u>		<u>50MG</u>	<u>A205321</u>	<u>002</u>	Mar 29, 2023
<u>AB</u>		<u>75MG</u>	<u>A205321</u>	<u>003</u>	Mar 29, 2023
<u>AB</u>		<u>100MG</u>	<u>A205321</u>	<u>004</u>	Mar 29, 2023
<u>AB</u>		<u>150MG</u>	<u>A205321</u>	<u>005</u>	Mar 29, 2023
<u>AB</u>		<u>200MG</u>	<u>A205321</u>	<u>006</u>	Mar 29, 2023
<u>AB</u>		<u>225MG</u>	<u>A205321</u>	<u>007</u>	Mar 29, 2023
<u>AB</u>		<u>300MG</u>	<u>A205321</u>	<u>008</u>	Mar 29, 2023
<u>AB</u>	CADILA PHARMS LTD	<u>25MG</u>	<u>A206452</u>	<u>001</u>	Jul 12, 2023
<u>AB</u>		<u>50MG</u>	<u>A206452</u>	<u>002</u>	Jul 12, 2023
<u>AB</u>		<u>75MG</u>	<u>A206452</u>	<u>003</u>	Jul 12, 2023
<u>AB</u>		<u>100MG</u>	<u>A206452</u>	<u>004</u>	Jul 12, 2023
<u>AB</u>		<u>150MG</u>	<u>A206452</u>	<u>005</u>	Jul 12, 2023
<u>AB</u>		<u>200MG</u>	<u>A206452</u>	<u>006</u>	Jul 12, 2023
<u>AB</u>		<u>225MG</u>	<u>A206452</u>	<u>007</u>	Jul 12, 2023
<u>AB</u>		<u>300MG</u>	<u>A206452</u>	<u>008</u>	Jul 12, 2023
<u>AB</u>	CHANGZHOU PHARM	<u>50MG</u>	<u>A214322</u>	<u>001</u>	Jul 15, 2021
<u>AB</u>		<u>75MG</u>	<u>A214322</u>	<u>002</u>	Jul 15, 2021
<u>AB</u>		<u>100MG</u>	<u>A214322</u>	<u>003</u>	Jul 15, 2021
<u>AB</u>		<u>150MG</u>	<u>A214322</u>	<u>004</u>	Jul 15, 2021
<u>AB</u>		<u>200MG</u>	<u>A214322</u>	<u>005</u>	Jul 15, 2021
<u>AB</u>		<u>225MG</u>	<u>A214322</u>	<u>006</u>	Jul 15, 2021
<u>AB</u>		<u>300MG</u>	<u>A214322</u>	<u>007</u>	Jul 15, 2021
<u>AB</u>	CHARTWELL RX	<u>50MG</u>	<u>A212865</u>	<u>001</u>	Mar 20, 2020
<u>AB</u>		<u>75MG</u>	<u>A212865</u>	<u>002</u>	Mar 20, 2020
<u>AB</u>		<u>100MG</u>	<u>A212865</u>	<u>003</u>	Mar 20, 2020
<u>AB</u>		<u>150MG</u>	<u>A212865</u>	<u>004</u>	Mar 20, 2020
<u>AB</u>	CIPLA	<u>25MG</u>	<u>A212280</u>	<u>001</u>	Jan 10, 2020
<u>AB</u>		<u>50MG</u>	<u>A212280</u>	<u>002</u>	Jan 10, 2020
<u>AB</u>		<u>75MG</u>	<u>A212280</u>	<u>003</u>	Jan 10, 2020
<u>AB</u>		<u>100MG</u>	<u>A212280</u>	<u>004</u>	Jan 10, 2020
<u>AB</u>		<u>150MG</u>	<u>A212280</u>	<u>005</u>	Jan 10, 2020
<u>AB</u>		<u>200MG</u>	<u>A212280</u>	<u>006</u>	Jan 10, 2020
<u>AB</u>		<u>225MG</u>	<u>A212280</u>	<u>007</u>	Jan 10, 2020
<u>AB</u>		<u>300MG</u>	<u>A212280</u>	<u>008</u>	Jan 10, 2020
<u>AB</u>	CREEKWOOD PHARMS	<u>25MG</u>	<u>A213423</u>	<u>001</u>	Mar 23, 2020
<u>AB</u>		<u>50MG</u>	<u>A213423</u>	<u>002</u>	Mar 23, 2020
<u>AB</u>		<u>75MG</u>	<u>A213423</u>	<u>003</u>	Mar 23, 2020
<u>AB</u>		<u>100MG</u>	<u>A213423</u>	<u>004</u>	Mar 23, 2020
<u>AB</u>		<u>150MG</u>	<u>A213423</u>	<u>005</u>	Mar 23, 2020
<u>AB</u>		<u>200MG</u>	<u>A213423</u>	<u>006</u>	Mar 23, 2020
<u>AB</u>		<u>225MG</u>	<u>A213423</u>	<u>007</u>	Mar 23, 2020
<u>AB</u>		<u>300MG</u>	<u>A213423</u>	<u>008</u>	Mar 23, 2020
<u>AB</u>	DR REDDYS	<u>25MG</u>	<u>A209664</u>	<u>001</u>	Jul 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<u>AB</u>		<u>50MG</u>	<u>A209664 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209664 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209664 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209664 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209664 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209664 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209664 008</u>	Jul 19, 2019
<u>AB</u>	ESKAYEF	<u>25MG</u>	<u>A212988 001</u>	Mar 08, 2022
<u>AB</u>		<u>50MG</u>	<u>A212988 002</u>	Mar 08, 2022
<u>AB</u>		<u>75MG</u>	<u>A212988 003</u>	Mar 08, 2022
<u>AB</u>		<u>100MG</u>	<u>A212988 004</u>	Mar 08, 2022
<u>AB</u>		<u>150MG</u>	<u>A212988 005</u>	Mar 08, 2022
<u>AB</u>		<u>200MG</u>	<u>A212988 006</u>	Mar 08, 2022
<u>AB</u>		<u>225MG</u>	<u>A212988 007</u>	Mar 08, 2022
<u>AB</u>		<u>300MG</u>	<u>A212988 008</u>	Mar 08, 2022
<u>AB</u>	HETERO LABS LTD III	<u>25MG</u>	<u>A206912 001</u>	Oct 08, 2019
<u>AB</u>		<u>50MG</u>	<u>A206912 002</u>	Oct 08, 2019
<u>AB</u>		<u>75MG</u>	<u>A206912 003</u>	Oct 08, 2019
<u>AB</u>		<u>100MG</u>	<u>A206912 004</u>	Oct 08, 2019
<u>AB</u>		<u>150MG</u>	<u>A206912 005</u>	Oct 08, 2019
<u>AB</u>		<u>200MG</u>	<u>A206912 006</u>	Oct 08, 2019
<u>AB</u>		<u>225MG</u>	<u>A206912 007</u>	Oct 08, 2019
<u>AB</u>		<u>300MG</u>	<u>A206912 008</u>	Oct 08, 2019
<u>AB</u>	INVAGEN PHARMS	<u>25MG</u>	<u>A211384 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A211384 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A211384 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A211384 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A211384 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A211384 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A211384 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A211384 008</u>	Jul 19, 2019
<u>AB</u>	MSN	<u>25MG</u>	<u>A209357 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A209357 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A209357 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A209357 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A209357 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A209357 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A209357 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A209357 008</u>	Jul 19, 2019
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A207883 001</u>	Sep 01, 2022
<u>AB</u>		<u>50MG</u>	<u>A207883 002</u>	Sep 01, 2022
<u>AB</u>		<u>75MG</u>	<u>A207883 003</u>	Sep 01, 2022
<u>AB</u>		<u>100MG</u>	<u>A207883 004</u>	Sep 01, 2022
<u>AB</u>		<u>150MG</u>	<u>A207883 005</u>	Sep 01, 2022
<u>AB</u>		<u>200MG</u>	<u>A207883 006</u>	Sep 01, 2022
<u>AB</u>		<u>225MG</u>	<u>A207883 007</u>	Sep 01, 2022
<u>AB</u>		<u>300MG</u>	<u>A207883 008</u>	Sep 01, 2022
<u>AB</u>	RENATA	<u>50MG</u>	<u>A210585 001</u>	Dec 26, 2019
<u>AB</u>		<u>75MG</u>	<u>A210585 002</u>	Dec 26, 2019
<u>AB</u>		<u>100MG</u>	<u>A210585 003</u>	Dec 26, 2019
<u>AB</u>		<u>150MG</u>	<u>A210585 004</u>	Dec 26, 2019
<u>AB</u>		<u>200MG</u>	<u>A210585 005</u>	Dec 26, 2019
<u>AB</u>		<u>300MG</u>	<u>A210585 006</u>	Dec 26, 2019
<u>AB</u>	RISING	<u>25MG</u>	<u>A210432 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A210432 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A210432 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A210432 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A210432 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A210432 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A210432 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A210432 008</u>	Jul 19, 2019
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A208677 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A208677 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A208677 003</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A208677 004</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A208677 005</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A208677 006</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A208677 007</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A208677 008</u>	Jul 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

PREGABALIN

CAPSULE; ORAL

PREGABALIN

<u>AB</u>	SUN PHARM	<u>25MG</u>	<u>A091157 001</u>	Nov 29, 2019
<u>AB</u>		<u>50MG</u>	<u>A091157 002</u>	Nov 29, 2019
<u>AB</u>		<u>75MG</u>	<u>A091157 003</u>	Nov 29, 2019
<u>AB</u>		<u>100MG</u>	<u>A091157 004</u>	Nov 29, 2019
<u>AB</u>		<u>150MG</u>	<u>A091157 005</u>	Nov 29, 2019
<u>AB</u>		<u>200MG</u>	<u>A091157 006</u>	Nov 29, 2019
<u>AB</u>		<u>225MG</u>	<u>A091157 007</u>	Nov 29, 2019
<u>AB</u>		<u>300MG</u>	<u>A091157 008</u>	Nov 29, 2019
<u>AB</u>	TEVA PHARMS	<u>25MG</u>	<u>A091219 001</u>	Jul 19, 2019
<u>AB</u>		<u>50MG</u>	<u>A091219 002</u>	Jul 19, 2019
<u>AB</u>		<u>75MG</u>	<u>A091224 001</u>	Jul 19, 2019
<u>AB</u>		<u>100MG</u>	<u>A091224 002</u>	Jul 19, 2019
<u>AB</u>		<u>150MG</u>	<u>A091224 003</u>	Jul 19, 2019
<u>AB</u>		<u>200MG</u>	<u>A091224 004</u>	Jul 19, 2019
<u>AB</u>		<u>225MG</u>	<u>A091224 005</u>	Jul 19, 2019
<u>AB</u>		<u>300MG</u>	<u>A091224 006</u>	Jul 19, 2019
<u>AB</u>	YILING	<u>75MG</u>	<u>A210891 001</u>	Sep 30, 2019
<u>AB</u>		<u>300MG</u>	<u>A210891 002</u>	Sep 30, 2019

SOLUTION; ORAL

LYRICA

<u>AA</u>	+! UPJOHN	<u>20MG/ML</u>	<u>N022488 001</u>	Jan 04, 2010
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PREGABALIN

<u>AA</u>	ALKEM LABS LTD	<u>20MG/ML</u>	<u>A207623 001</u>	Jul 19, 2019
<u>AA</u>	PATRIN	<u>20MG/ML</u>	<u>A212604 001</u>	Feb 18, 2022

TABLET, EXTENDED RELEASE; ORAL

LYRICA CR

<u>AB</u>	+ UPJOHN	<u>82.5MG</u>	<u>N209501 001</u>	Oct 11, 2017
<u>AB</u>	+	<u>165MG</u>	<u>N209501 002</u>	Oct 11, 2017
<u>AB</u>	+!	<u>330MG</u>	<u>N209501 003</u>	Oct 11, 2017

PREGABALIN

<u>AB</u>	ALVOGEN	<u>82.5MG</u>	<u>A211593 001</u>	Apr 13, 2021
<u>AB</u>		<u>165MG</u>	<u>A211593 002</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A211593 003</u>	Apr 13, 2021
<u>AB</u>	APOTEX	<u>165MG</u>	<u>A213313 001</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A213313 002</u>	Apr 13, 2021
<u>AB</u>	EPIC PHARMA LLC	<u>82.5MG</u>	<u>A214496 001</u>	Jun 01, 2023
<u>AB</u>		<u>165MG</u>	<u>A214496 002</u>	Jun 01, 2023
<u>AB</u>		<u>330MG</u>	<u>A214496 003</u>	Jun 01, 2023
<u>AB</u>	MSN	<u>82.5MG</u>	<u>A213226 001</u>	Apr 13, 2021
<u>AB</u>		<u>165MG</u>	<u>A213226 002</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A213226 003</u>	Apr 13, 2021
<u>AB</u>	RUBICON	<u>82.5MG</u>	<u>A215249 001</u>	Mar 22, 2022
<u>AB</u>		<u>165MG</u>	<u>A215249 002</u>	Mar 22, 2022
<u>AB</u>		<u>330MG</u>	<u>A215249 003</u>	Mar 22, 2022
<u>AB</u>	SUN PHARM	<u>82.5MG</u>	<u>A211889 001</u>	Apr 13, 2021
<u>AB</u>		<u>165MG</u>	<u>A211889 002</u>	Apr 13, 2021
<u>AB</u>		<u>330MG</u>	<u>A211889 003</u>	Apr 13, 2021

PRETOMANID

TABLET; ORAL

PRETOMANID

	+! MYLAN IRELAND LTD	200MG	N212862 001	Aug 14, 2019
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PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRILOCAINE HYDROCHLORIDE

	! SEPTODONT INC	4%	A079235 001	Sep 29, 2010
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PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE

<u>AB</u>	+! SANOFI AVENTIS US	<u>EQ 15MG BASE</u>	<u>N008316 001</u>	
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PRIMAQUINE PHOSPHATE

<u>AB</u>	BAYSHORE PHARMS LLC	<u>EQ 15MG BASE</u>	<u>A204476 001</u>	Feb 25, 2014
<u>AB</u>	NOVAST LABS	<u>EQ 15MG BASE</u>	<u>A206043 001</u>	Jun 23, 2016

PRESCRIPTION DRUG PRODUCT LIST

PRIMIDONE

TABLET; ORAL

MYSOLINE

<u>AB</u>	+	VALEANT	<u>50MG</u>	<u>N009170</u>	<u>003</u>	
<u>AB</u>	+		<u>250MG</u>	<u>N009170</u>	<u>002</u>	

PRIMIDONE

<u>AB</u>		AMNEAL PHARM	<u>50MG</u>	<u>A040866</u>	<u>001</u>	Apr 23, 2008
<u>AB</u>			<u>250MG</u>	<u>A040866</u>	<u>002</u>	Apr 23, 2008
<u>AB</u>		ANDA REPOSITORY	<u>50MG</u>	<u>A040626</u>	<u>001</u>	Sep 29, 2005
<u>AB</u>			<u>250MG</u>	<u>A040626</u>	<u>002</u>	Sep 29, 2005
<u>AB</u>		LANNETT	<u>50MG</u>	<u>A084903</u>	<u>002</u>	May 24, 2001
<u>AB</u>			<u>250MG</u>	<u>A084903</u>	<u>001</u>	
<u>AB</u>		OXFORD PHARMS	<u>50MG</u>	<u>A040586</u>	<u>001</u>	Feb 24, 2005
<u>AB</u>			<u>250MG</u>	<u>A040586</u>	<u>002</u>	Feb 24, 2005
<u>AB</u>		RUBICON	<u>50MG</u>	<u>A214896</u>	<u>001</u>	Jun 28, 2022
<u>AB</u>			<u>250MG</u>	<u>A214896</u>	<u>002</u>	Jun 28, 2022
<u>AB</u>		WATSON LABS	<u>250MG</u>	<u>A083551</u>	<u>001</u>	
<u>AB</u>		RUBICON	125MG	A214896	003	Dec 28, 2022

PROBENECID

TABLET; ORAL

PROBALAN

<u>AB</u>		LANNETT	<u>500MG</u>	<u>A080966</u>	<u>001</u>	
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PROBENECID

<u>AB</u>		RISING	<u>500MG</u>	<u>A217020</u>	<u>001</u>	Nov 20, 2023
<u>AB</u>	!	WATSON LABS TEVA	<u>500MG</u>	<u>A084442</u>	<u>004</u>	Mar 29, 1983

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

<u>AP</u>	!	HOSPIRA	<u>100MG/ML</u>	<u>A089069</u>	<u>001</u>	Feb 12, 1986
<u>AP</u>	!		<u>500MG/ML</u>	<u>A089070</u>	<u>001</u>	Feb 12, 1986
<u>AP</u>		INTL MEDICATION	<u>100MG/ML</u>	<u>A088636</u>	<u>001</u>	Jul 31, 1984
<u>AP</u>		NEXUS	<u>100MG/ML</u>	<u>A206332</u>	<u>001</u>	Oct 13, 2017
<u>AP</u>			<u>500MG/ML</u>	<u>A206332</u>	<u>002</u>	Oct 13, 2017

PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL

MATULANE

	+	LEADIANT BIOSCI INC	EQ 50MG BASE	N016785	001	
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PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPRO

<u>AB</u>		PADAGIS US	<u>25MG</u>	<u>A040246</u>	<u>001</u>	Jun 28, 2000
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PROCHLORPERAZINE

<u>AB</u>	!	COSETTE	<u>25MG</u>	<u>A040058</u>	<u>001</u>	Nov 24, 1993
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PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

<u>AP</u>		AMNEAL	<u>EQ 5MG BASE/ML</u>	<u>A214192</u>	<u>001</u>	Nov 28, 2022
<u>AP</u>	!	AVET LIFESCIENCES	<u>EQ 5MG BASE/ML</u>	<u>A204147</u>	<u>001</u>	Oct 15, 2013
<u>AP</u>		CAPLIN	<u>EQ 5MG BASE/ML</u>	<u>A214379</u>	<u>001</u>	Apr 22, 2021
<u>AP</u>		EUGIA PHARMA	<u>EQ 5MG BASE/ML</u>	<u>A213873</u>	<u>001</u>	Jul 14, 2022
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A214107</u>	<u>001</u>	Sep 22, 2021
<u>AP</u>		HIKMA	<u>EQ 5MG BASE/ML</u>	<u>A089903</u>	<u>001</u>	Aug 29, 1989
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A213630</u>	<u>001</u>	Nov 22, 2022
<u>AP</u>		MYLAN LABS LTD	<u>EQ 5MG BASE/ML</u>	<u>A210710</u>	<u>001</u>	Oct 25, 2018
<u>AP</u>		NEXUS	<u>EQ 5MG BASE/ML</u>	<u>A204860</u>	<u>001</u>	Feb 15, 2019
<u>AP</u>		SAGENT	<u>EQ 5MG BASE/ML</u>	<u>A040540</u>	<u>001</u>	May 28, 2004
<u>AP</u>		VIWIT PHARM	<u>EQ 5MG BASE/ML</u>	<u>A213626</u>	<u>001</u>	Sep 28, 2021

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

<u>AB</u>		AMNEAL	<u>EQ 5MG BASE</u>	<u>A216598</u>	<u>001</u>	Apr 17, 2023
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A216598</u>	<u>002</u>	Apr 17, 2023
<u>AB</u>		BIONPHARMA	<u>EQ 5MG BASE</u>	<u>A217478</u>	<u>001</u>	Apr 04, 2023
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A217478</u>	<u>002</u>	Apr 04, 2023
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A216595</u>	<u>001</u>	Mar 17, 2023
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A216595</u>	<u>002</u>	Mar 17, 2023
<u>AB</u>		NOVITIUM PHARMA	<u>EQ 5MG BASE</u>	<u>A216202</u>	<u>001</u>	Jun 13, 2022
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A216202</u>	<u>002</u>	Jun 13, 2022
<u>AB</u>		ZYDUS	<u>EQ 5MG BASE</u>	<u>A216495</u>	<u>001</u>	Aug 08, 2022

PRESCRIPTION DRUG PRODUCT LIST

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A216495 002</u>	Aug 08, 2022
	<u>PROCOMP</u>			
<u>AB</u>	JUBILANT CADISTA	<u>EQ 5MG BASE</u>	<u>A040268 001</u>	Feb 27, 1998
<u>AB</u>	!	<u>EQ 10MG BASE</u>	<u>A040268 002</u>	Feb 27, 1998

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

<u>AB</u>	AMNEAL PHARMS NY	<u>100MG</u>	<u>A207724 001</u>	Sep 07, 2017
<u>AB</u>		<u>200MG</u>	<u>A207724 002</u>	Sep 07, 2017
<u>AB</u>	BIONPHARMA	<u>100MG</u>	<u>A200900 001</u>	Aug 16, 2013
<u>AB</u>		<u>200MG</u>	<u>A200900 002</u>	Aug 16, 2013
<u>AB</u>	DR REDDYS	<u>100MG</u>	<u>A208801 001</u>	Feb 28, 2017
<u>AB</u>		<u>200MG</u>	<u>A208801 002</u>	Feb 28, 2017
<u>AB</u>	EUGIA PHARMA	<u>100MG</u>	<u>A211285 001</u>	Oct 26, 2018
<u>AB</u>		<u>200MG</u>	<u>A211285 002</u>	Oct 26, 2018
<u>AB</u>	SOFGEN PHARMS	<u>100MG</u>	<u>A200456 001</u>	Sep 28, 2012
<u>AB</u>		<u>200MG</u>	<u>A200456 002</u>	Sep 28, 2012
<u>AB</u>	XIROMED	<u>100MG</u>	<u>A205229 001</u>	Oct 20, 2017
<u>AB</u>		<u>200MG</u>	<u>A205229 002</u>	Oct 20, 2017

PROMETRIUM

<u>AB</u>	+ VIRTUS	<u>100MG</u>	<u>N019781 001</u>	May 14, 1998
<u>AB</u>	+	<u>200MG</u>	<u>N019781 002</u>	Oct 15, 1999

GEL; VAGINAL

CRINONE

	+! ALLERGAN	4%	N020701 001	Jul 31, 1997
	+!	8%	N020701 002	Jul 31, 1997

INJECTABLE; INJECTION

PROGESTERONE

<u>AO</u>	ACCORD HLTHCARE	<u>50MG/ML</u>	<u>A217707 001</u>	Nov 21, 2023
<u>AO</u>	! EUGIA PHARMA	<u>50MG/ML</u>	<u>A210965 001</u>	Dec 06, 2018
<u>AO</u>	FRESENIUS KABI USA	<u>50MG/ML</u>	<u>A075906 001</u>	Apr 25, 2001
<u>AO</u>	HIKMA FARMACEUTICA	<u>50MG/ML</u>	<u>A091033 001</u>	Oct 28, 2010
<u>AO</u>	XIROMED	<u>50MG/ML</u>	<u>A215634 001</u>	Jan 05, 2022

INSERT; VAGINAL

ENDOMETRIN

	+! FERRING	100MG	N022057 001	Jun 21, 2007
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PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMETHAZINE HYDROCHLORIDE

<u>AP</u>	+! WEST-WARD PHARMS	<u>25MG/ML</u>	<u>A083312 001</u>	
	INT			
<u>AP</u>	+!	<u>50MG/ML</u>	<u>A083312 002</u>	
<u>AP</u>	XGEN PHARMS	<u>25MG/ML</u>	<u>A040737 001</u>	Apr 24, 2008
<u>AP</u>		<u>50MG/ML</u>	<u>A040737 002</u>	Apr 24, 2008

SUPPOSITORY; RECTAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	ANNORA PHARMA	<u>12.5MG</u>	<u>A216446 001</u>	Nov 02, 2022
<u>AB</u>		<u>25MG</u>	<u>A216446 002</u>	Nov 02, 2022
<u>AB</u>	COSETTE	<u>12.5MG</u>	<u>A040428 002</u>	Mar 31, 2003
<u>AB</u>	!	<u>25MG</u>	<u>A040428 001</u>	Feb 05, 2002
<u>AB</u>	PADAGIS ISRAEL	<u>12.5MG</u>	<u>A040500 001</u>	Jun 30, 2003
<u>AB</u>		<u>25MG</u>	<u>A040500 002</u>	Jun 30, 2003
<u>AB</u>	TARO	<u>12.5MG</u>	<u>A040603 001</u>	Oct 26, 2006
<u>AB</u>		<u>25MG</u>	<u>A040603 002</u>	Oct 26, 2006

PROMETHEGAN

	! COSETTE	50MG	A087165 001	Aug 14, 1987
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SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AA</u>	CHARTWELL RX	<u>6.25MG/5ML</u>	<u>A040026 001</u>	Sep 25, 1998
<u>AA</u>	NOSTRUM LABS INC	<u>6.25MG/5ML</u>	<u>A040891 001</u>	Mar 13, 2009
<u>AA</u>	QUAGEN	<u>6.25MG/5ML</u>	<u>A213890 001</u>	Jul 12, 2021
<u>AA</u>	TARO	<u>6.25MG/5ML</u>	<u>A040718 001</u>	Apr 04, 2007
<u>AA</u>	TRIS PHARMA INC	<u>6.25MG/5ML</u>	<u>A091675 001</u>	Jun 28, 2012

PROMETHAZINE PLAIN

<u>AA</u>	+! WOCHKHARDT BIO AG	<u>6.25MG/5ML</u>	<u>A087953 001</u>	Nov 15, 1982
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TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS NY	<u>12.5MG</u>	<u>A091179 001</u>	Dec 13, 2010
<u>AB</u>		<u>25MG</u>	<u>A091179 002</u>	Dec 13, 2010

PRESCRIPTION DRUG PRODUCT LIST

PROMETHAZINE HYDROCHLORIDE

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>		<u>50MG</u>	<u>A091179</u>	<u>003</u>	Dec 13, 2010
<u>AB</u>	KVK TECH	<u>12.5MG</u>	<u>A040712</u>	<u>002</u>	May 04, 2007
<u>AB</u>		<u>25MG</u>	<u>A040712</u>	<u>001</u>	Jul 31, 2006
<u>AB</u>		<u>50MG</u>	<u>A040712</u>	<u>003</u>	Jul 31, 2006
<u>AB</u>	PRINSTON INC	<u>12.5MG</u>	<u>A040622</u>	<u>001</u>	Jul 18, 2006
<u>AB</u>		<u>25MG</u>	<u>A040622</u>	<u>002</u>	Jul 18, 2006
<u>AB</u>		<u>50MG</u>	<u>A040622</u>	<u>003</u>	Jul 18, 2006
<u>AB</u>	QUAGEN	<u>12.5MG</u>	<u>A040673</u>	<u>001</u>	Mar 05, 2008
<u>AB</u>		<u>25MG</u>	<u>A040673</u>	<u>002</u>	Mar 05, 2008
<u>AB</u>		<u>50MG</u>	<u>A040673</u>	<u>003</u>	Mar 05, 2008
<u>AB</u>	+ SANDOZ	<u>25MG</u>	<u>A084176</u>	<u>003</u>	
<u>AB</u>	+!	<u>50MG</u>	<u>A084176</u>	<u>001</u>	
<u>AB</u>	STRIDES PHARMA	<u>12.5MG</u>	<u>A209177</u>	<u>001</u>	Jun 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A209177</u>	<u>002</u>	Jun 30, 2017
<u>AB</u>		<u>50MG</u>	<u>A209177</u>	<u>003</u>	Jun 30, 2017
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A083426</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A083711</u>	<u>001</u>	
<u>AB</u>	ZYDUS PHARMS USA	<u>12.5MG</u>	<u>A040596</u>	<u>001</u>	Nov 18, 2005
<u>AB</u>		<u>25MG</u>	<u>A040596</u>	<u>002</u>	Nov 18, 2005
<u>AB</u>		<u>50MG</u>	<u>A040596</u>	<u>003</u>	Nov 18, 2005

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>225MG</u>	<u>A213096</u>	<u>001</u>	Feb 21, 2023
<u>AB</u>		<u>325MG</u>	<u>A213096</u>	<u>002</u>	Feb 21, 2023
<u>AB</u>		<u>425MG</u>	<u>A213096</u>	<u>003</u>	Feb 21, 2023
<u>AB</u>	GLENMARK PHARMS LTD	<u>225MG</u>	<u>A205268</u>	<u>001</u>	Sep 08, 2017
<u>AB</u>		<u>325MG</u>	<u>A205268</u>	<u>002</u>	Sep 08, 2017
<u>AB</u>		<u>425MG</u>	<u>A205268</u>	<u>003</u>	Sep 08, 2017
<u>AB</u>	RISING	<u>225MG</u>	<u>A205956</u>	<u>001</u>	Jul 02, 2018
<u>AB</u>		<u>325MG</u>	<u>A205956</u>	<u>002</u>	Jul 02, 2018
<u>AB</u>		<u>425MG</u>	<u>A205956</u>	<u>003</u>	Jul 02, 2018
<u>AB</u>	SCINOPHARM TAIWAN	<u>225MG</u>	<u>A212928</u>	<u>001</u>	Jun 18, 2020
<u>AB</u>		<u>325MG</u>	<u>A212928</u>	<u>002</u>	Jun 18, 2020
<u>AB</u>		<u>425MG</u>	<u>A212928</u>	<u>003</u>	Jun 18, 2020
<u>AB</u>	STRIDES PHARMA	<u>225MG</u>	<u>A078540</u>	<u>001</u>	Oct 18, 2010
<u>AB</u>		<u>325MG</u>	<u>A078540</u>	<u>002</u>	Oct 18, 2010
<u>AB</u>	!	<u>425MG</u>	<u>A078540</u>	<u>003</u>	Oct 18, 2010
<u>AB</u>	UPSHER SMITH LABS	<u>225MG</u>	<u>A212744</u>	<u>001</u>	Jun 25, 2020
<u>AB</u>		<u>325MG</u>	<u>A212744</u>	<u>002</u>	Jun 25, 2020
<u>AB</u>		<u>425MG</u>	<u>A212744</u>	<u>003</u>	Jun 25, 2020
<u>AB</u>	WATSON LABS INC	<u>225MG</u>	<u>A202688</u>	<u>001</u>	Aug 24, 2015
<u>AB</u>		<u>325MG</u>	<u>A202688</u>	<u>002</u>	Aug 24, 2015
<u>AB</u>		<u>425MG</u>	<u>A202688</u>	<u>003</u>	Aug 24, 2015
<u>AB</u>	ZYDUS	<u>225MG</u>	<u>A214184</u>	<u>001</u>	Apr 21, 2021
<u>AB</u>		<u>325MG</u>	<u>A214184</u>	<u>002</u>	Apr 21, 2021
<u>AB</u>		<u>425MG</u>	<u>A214184</u>	<u>003</u>	Apr 21, 2021
BX	SINOTHERAPEUTICS INC	225MG	A210339	001	Jan 04, 2019
BX		325MG	A210339	002	Jan 04, 2019
BX		425MG	A210339	003	Jan 04, 2019

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS	<u>150MG</u>	<u>A076550</u>	<u>001</u>	Apr 23, 2004
<u>AB</u>		<u>225MG</u>	<u>A076550</u>	<u>002</u>	Apr 23, 2004
<u>AB</u>		<u>300MG</u>	<u>A076550</u>	<u>003</u>	Apr 23, 2004
<u>AB</u>	AUROBINDO PHARMA	<u>150MG</u>	<u>A202445</u>	<u>001</u>	May 11, 2016
<u>AB</u>		<u>225MG</u>	<u>A202445</u>	<u>002</u>	May 11, 2016
<u>AB</u>		<u>300MG</u>	<u>A202445</u>	<u>003</u>	May 11, 2016
<u>AB</u>	STRIDES PHARMA	<u>150MG</u>	<u>A075938</u>	<u>001</u>	Oct 17, 2002
<u>AB</u>		<u>225MG</u>	<u>A075938</u>	<u>002</u>	Oct 17, 2002
<u>AB</u>	!	<u>300MG</u>	<u>A075938</u>	<u>003</u>	Oct 17, 2002
<u>AB</u>	SUN PHARM INDUSTRIES	<u>150MG</u>	<u>A075998</u>	<u>001</u>	Nov 29, 2001
<u>AB</u>		<u>225MG</u>	<u>A075998</u>	<u>002</u>	Nov 29, 2001
<u>AB</u>		<u>300MG</u>	<u>A075998</u>	<u>003</u>	Nov 29, 2001
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A075203</u>	<u>001</u>	Oct 24, 2000
<u>AB</u>		<u>225MG</u>	<u>A075203</u>	<u>002</u>	Oct 24, 2000

PRESCRIPTION DRUG PRODUCT LIST

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ALCAINE

AT +! ALCON LABS INC **0.5%** **A080027 001**

PROPARACAINE HYDROCHLORIDE

AT ! BAUSCH AND LOMB **0.5%** **A040074 001** Sep 29, 1995
AT RISING **0.5%** **A040277 001** Mar 16, 2000

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

AB +! FRESENIUS KABI USA **10MG/ML** **N019627 002** Jun 11, 1996

PROPOFOL

AB AVET LIFESCIENCES **10MG/ML** **A206408 001** Oct 12, 2021
AB DR REDDYS **10MG/ML** **A205067 001** Nov 15, 2018
AB HIKMA **10MG/ML** **A074848 001** Apr 19, 2005
AB HOSPIRA **10MG/ML** **A077908 001** Mar 17, 2006
AB INNOPHARMA **10MG/ML** **A205576 001** Sep 16, 2020
AB SAGENT PHARMS INC **10MG/ML** **A075102 001** Jan 04, 1999

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

INDERAL LA

AB + ANI PHARMS **60MG** **N018553 004** Mar 18, 1987
AB + **80MG** **N018553 002** Apr 19, 1983
AB + **120MG** **N018553 003** Apr 19, 1983
AB +! **160MG** **N018553 001** Apr 19, 1983

PROPRANOLOL HYDROCHLORIDE

AB ACTAVIS ELIZABETH **60MG** **A078494 001** Aug 10, 2007
AB **80MG** **A078494 002** Aug 10, 2007
AB **120MG** **A078494 003** Aug 10, 2007
AB **160MG** **A078494 004** Aug 10, 2007
AB ADARE PHARMS INC **60MG** **A078703 001** Jul 15, 2011
AB **80MG** **A078703 002** Jul 15, 2011
AB **120MG** **A078703 003** Jul 15, 2011
AB **160MG** **A078703 004** Jul 15, 2011
AB AMTA **60MG** **A212026 001** Jan 06, 2020
AB **80MG** **A212026 002** Jan 06, 2020
AB **120MG** **A212026 003** Jan 06, 2020
AB **160MG** **A212026 004** Jan 06, 2020
AB NORTEC DEV ASSOC **60MG** **A078065 001** Jan 26, 2007
AB **80MG** **A078065 002** Jan 26, 2007
AB **120MG** **A078065 003** Jan 26, 2007
AB **160MG** **A078065 004** Jan 26, 2007
AB ZYDUS PHARMS USA **60MG** **A090321 001** Mar 25, 2011
AB INC **80MG** **A090321 002** Mar 25, 2011
AB **120MG** **A090321 003** Mar 25, 2011
AB **160MG** **A090321 004** Mar 25, 2011

INNOPRAN XL

BX + ANI PHARMS **80MG** **N021438 001** Mar 12, 2003
BX +! **120MG** **N021438 002** Mar 12, 2003

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

AP FRESENIUS KABI USA **1MG/ML** **A075826 001** Aug 31, 2001
AP ! HIKMA FARMACEUTICA **1MG/ML** **A077760 001** Jan 31, 2008

SOLUTION; ORAL

HEMANGEOL

+! PIERRE FABRE DERMA **4.28MG/ML** **N205410 001** Mar 14, 2014
! HIKMA **20MG/5ML** **A070979 001** May 15, 1987
! **40MG/5ML** **A070690 001** May 15, 1987

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

AB IMPAX LABS INC **10MG** **A071972 001** Apr 06, 1988
AB **20MG** **A071972 002** Apr 06, 1988
AB **40MG** **A071972 003** Apr 06, 1988
AB **60MG** **A071976 002** May 13, 1986
AB ! **80MG** **A071976 001** Apr 06, 1988
AB INNOGENIX **10MG** **A070322 002** Oct 22, 1985
AB **20MG** **A070322 003** Oct 22, 1985
AB **40MG** **A070322 004** Oct 22, 1985
AB **60MG** **A070322 005** Sep 24, 1986
AB **80MG** **A070322 001** Aug 04, 1986

PRESCRIPTION DRUG PRODUCT LIST

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

<u>AB</u>	IPCA LABS LTD	<u>10MG</u>	<u>A078955 001</u>	Jun 02, 2008
<u>AB</u>		<u>20MG</u>	<u>A078955 002</u>	Jun 02, 2008
<u>AB</u>		<u>40MG</u>	<u>A078955 003</u>	Jun 02, 2008
<u>AB</u>		<u>60MG</u>	<u>A078955 004</u>	Jun 02, 2008
<u>AB</u>		<u>80MG</u>	<u>A078955 005</u>	Jun 02, 2008
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A070213 002</u>	Nov 19, 1985
<u>AB</u>		<u>20MG</u>	<u>A070213 003</u>	Nov 19, 1985
<u>AB</u>		<u>40MG</u>	<u>A070213 001</u>	Nov 19, 1985
<u>AB</u>		<u>60MG</u>	<u>A070213 005</u>	Apr 08, 2011
<u>AB</u>		<u>80MG</u>	<u>A070213 004</u>	Nov 19, 1985
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078213 001</u>	Jan 10, 2008
<u>AB</u>		<u>20MG</u>	<u>A078213 002</u>	Jan 10, 2008
<u>AB</u>		<u>40MG</u>	<u>A078213 003</u>	Jan 10, 2008
<u>AB</u>		<u>60MG</u>	<u>A078213 004</u>	Jan 10, 2008
<u>AB</u>		<u>80MG</u>	<u>A078213 005</u>	Jan 10, 2008
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A070221 002</u>	Aug 01, 1986
<u>AB</u>		<u>20MG</u>	<u>A070221 003</u>	Aug 01, 1986
<u>AB</u>		<u>40MG</u>	<u>A070221 004</u>	Aug 01, 1986
<u>AB</u>		<u>60MG</u>	<u>A070221 005</u>	Sep 24, 1986
<u>AB</u>		<u>80MG</u>	<u>A070221 001</u>	Apr 14, 1986
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A070175 001</u>	May 13, 1986
<u>AB</u>		<u>20MG</u>	<u>A070176 001</u>	May 13, 1986
<u>AB</u>		<u>40MG</u>	<u>A070177 001</u>	May 13, 1986
<u>AB</u>		<u>80MG</u>	<u>A070178 001</u>	May 13, 1986

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

<u>AB</u>	DAVA PHARMS INC	<u>50MG</u>	<u>N006188 001</u>	
<u>AB</u>	MACLEODS PHARMS LTD	<u>50MG</u>	<u>A208867 001</u>	May 10, 2023
BD	ACTAVIS ELIZABETH	50MG	A080172 001	
BD	QUAGEN	50MG	A080154 001	

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

!	FRESENIUS KABI USA	10MG/ML	A089454 001	Apr 07, 1987
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PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

PROTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A202220 001</u>	Nov 19, 2012
<u>AB</u>		<u>10MG</u>	<u>A202220 002</u>	Nov 19, 2012
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A078913 001</u>	Sep 16, 2008
<u>AB</u>	!	<u>10MG</u>	<u>A078913 002</u>	Sep 16, 2008
<u>AB</u>	SIGMAPHARM LABS LLC	<u>5MG</u>	<u>A090462 001</u>	May 03, 2010
<u>AB</u>		<u>10MG</u>	<u>A090462 002</u>	May 03, 2010

PRUCALOPRIDE SUCCINATE

TABLET; ORAL

MOTEGRITY

+	TAKEDA PHARMS USA	EQ 1MG BASE	N210166 001	Dec 14, 2018
+	!	EQ 2MG BASE	N210166 002	Dec 14, 2018

PYRAZINAMIDE

TABLET; ORAL

PYRAZINAMIDE

<u>AB</u>	!	HIKMA	<u>500MG</u>	<u>A081319 001</u>	Jun 30, 1992
<u>AB</u>		MACLEODS PHARMS LTD	<u>500MG</u>	<u>A212541 001</u>	Jul 27, 2020
<u>AB</u>	+	NOVITIUM PHARMA	<u>500MG</u>	<u>A080157 001</u>	

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION

MESTINON

<u>AP</u>	!	BAUSCH	<u>5MG/ML</u>	<u>N009830 001</u>	
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REGONOL

<u>AP</u>		SANDOZ	<u>5MG/ML</u>	<u>N017398 001</u>	
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SYRUP; ORAL

MESTINON

<u>AA</u>	!	BAUSCH	<u>60MG/5ML</u>	<u>N015193 001</u>	
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PYRIDOSTIGMINE BROMIDE

<u>AA</u>		AMNEAL	<u>60MG/5ML</u>	<u>A212702 001</u>	Jan 10, 2020
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PRESCRIPTION DRUG PRODUCT LIST

PYRIDOSTIGMINE BROMIDE

SYRUP; ORAL

PYRIDOSTIGMINE BROMIDE

<u>AA</u>	MILLA PHARMS	<u>60MG/5ML</u>	<u>A212405 001</u>	Apr 19, 2022
<u>AA</u>	MSN	<u>60MG/5ML</u>	<u>A216512 001</u>	Sep 21, 2023
<u>AA</u>	NOVITIUM PHARMA	<u>60MG/5ML</u>	<u>A211694 001</u>	Mar 08, 2019
<u>AA</u>	RISING	<u>60MG/5ML</u>	<u>A208797 001</u>	Jan 09, 2020

TABLET; ORAL

MESTINON

<u>AB</u>	<u>+</u> !	BAUSCH	<u>60MG</u>	<u>N009829 002</u>
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PYRIDOSTIGMINE BROMIDE

<u>AB</u>	EYWA	<u>30MG</u>	<u>A211181 002</u>	May 14, 2019
<u>AB</u>		<u>60MG</u>	<u>A211181 001</u>	Jul 20, 2018
<u>AB</u>	IMPAX LABS	<u>30MG</u>	<u>A040502 002</u>	Jun 28, 2022
<u>AB</u>		<u>60MG</u>	<u>A040502 001</u>	Apr 24, 2003
<u>AB</u>	ZYDUS PHARMS	<u>60MG</u>	<u>A205650 001</u>	Jun 22, 2015

TABLET, EXTENDED RELEASE; ORAL

MESTINON

<u>AB</u>	<u>+</u> !	BAUSCH	<u>180MG</u>	<u>N011665 001</u>
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PYRIDOSTIGMINE BROMIDE

<u>AB</u>	ALVOGEN	<u>180MG</u>	<u>A204737 001</u>	Jun 26, 2015
<u>AB</u>	IMPAX LABS INC	<u>180MG</u>	<u>A203184 001</u>	Sep 15, 2015
<u>AB</u>	RISING	<u>180MG</u>	<u>A205464 001</u>	Aug 15, 2017

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

PYRIDOXINE HYDROCHLORIDE

<u>+</u> !	FRESENIUS KABI USA	100MG/ML	A080618	001
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PYRIMETHAMINE

TABLET; ORAL

DARAPRIM

<u>AB</u>	<u>+</u> !	TILDE SCIENCES	<u>25MG</u>	<u>N008578 001</u>
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PYRIMETHAMINE

<u>AB</u>	ALVOGEN	<u>25MG</u>	<u>A211271 001</u>	Jul 27, 2021
<u>AB</u>	AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A216983 001</u>	Oct 25, 2022
<u>AB</u>	CEROVENE INC	<u>25MG</u>	<u>A207127 001</u>	Feb 28, 2020
<u>AB</u>	TEVA PHARMS	<u>25MG</u>	<u>A215506 001</u>	Aug 13, 2021

QUAZEPAM

TABLET; ORAL

DORAL

<u>+</u> !	GALT PHARMS	15MG	N018708	001	Dec 27, 1985
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QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 25MG BASE</u>	<u>A202152 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202152 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202152 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A202152 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202152 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A202152 006</u>	Mar 27, 2012
<u>AB</u>	ALKEM LABS LTD	<u>EQ 25MG BASE</u>	<u>A201504 001</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201504 002</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201504 003</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A201504 004</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A201504 005</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A201504 006</u>	Feb 12, 2013
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A201504 007</u>	Feb 12, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 25MG BASE</u>	<u>A091388 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A091388 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091388 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091388 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A091388 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A091388 006</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A091388 007</u>	Mar 27, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A077380 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077380 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077380 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A077380 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A077380 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077380 006</u>	Mar 27, 2012

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A077380 007</u>	Mar 27, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 25MG BASE</u>	<u>A204316 001</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204316 002</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204316 003</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204316 004</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204316 005</u>	Jun 16, 2022
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204316 006</u>	Jun 16, 2022
<u>AB</u>	HIKMA	<u>EQ 25MG BASE</u>	<u>A090120 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090749 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090749 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090749 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090749 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090749 005</u>	Mar 27, 2012
<u>AB</u>	LUPIN LTD	<u>EQ 25MG BASE</u>	<u>A201109 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201109 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201109 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A201109 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A201109 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A201109 006</u>	Mar 27, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A203359 001</u>	May 17, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A203359 002</u>	May 17, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A203359 003</u>	May 17, 2016
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A203359 004</u>	May 17, 2016
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A203359 005</u>	May 17, 2016
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A203359 006</u>	May 17, 2016
<u>AB</u>	PRINSTON INC	<u>EQ 25MG BASE</u>	<u>A206954 001</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A206954 002</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A206954 003</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A206954 004</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206954 005</u>	Aug 24, 2022
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A206954 006</u>	Aug 24, 2022
<u>AB</u>	SANDOZ	<u>EQ 25MG BASE</u>	<u>A078679 001</u>	Dec 14, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078679 002</u>	Dec 14, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078679 003</u>	Dec 14, 2012
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078679 004</u>	Dec 14, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A078679 005</u>	Dec 14, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A078679 006</u>	Dec 14, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A078679 007</u>	Dec 14, 2012
<u>AB</u>	SUN PHARM	<u>EQ 25MG BASE</u>	<u>A201190 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201190 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201190 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A201190 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A201190 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A201190 006</u>	Mar 27, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 25MG BASE</u>	<u>A077745 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077745 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077745 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A077745 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A077745 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077745 006</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A077745 007</u>	Mar 27, 2012
<u>AB</u>	UNICHEM	<u>EQ 25MG BASE</u>	<u>A202674 001</u>	Mar 08, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202674 002</u>	Mar 08, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202674 003</u>	Mar 08, 2016
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A202674 004</u>	Mar 08, 2016
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202674 005</u>	Mar 08, 2016
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A202674 006</u>	Mar 08, 2016
<u>AB</u>	ZENNOVA	<u>EQ 25MG BASE</u>	<u>A090960 001</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090960 002</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090960 003</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090960 004</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090960 005</u>	Mar 27, 2012
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090960 006</u>	Mar 27, 2012
<u>SEROQUEL</u>				
<u>AB</u>	+!	<u>EQ 25MG BASE</u>	<u>N020639 001</u>	Sep 26, 1997
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N020639 007</u>	Oct 04, 2005
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N020639 002</u>	Sep 26, 1997
<u>AB</u>	+	<u>EQ 200MG BASE</u>	<u>N020639 003</u>	Sep 26, 1997
<u>AB</u>	+	<u>EQ 300MG BASE</u>	<u>N020639 005</u>	Jul 26, 2000

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET; ORAL

SEROQUEL

<u>AB</u>	+	<u>EQ 400MG BASE</u>	<u>N020639</u>	<u>006</u>	Oct 04, 2005
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TABLET, EXTENDED RELEASE; ORAL

QUETIAPINE FUMARATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 50MG BASE</u>	<u>A206252</u>	<u>001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090681</u>	<u>001</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090681</u>	<u>002</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090681</u>	<u>003</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090681</u>	<u>004</u>	Nov 01, 2016
<u>AB</u>	ALIGNSCIENCE PHARMA	<u>EQ 150MG BASE</u>	<u>A209497</u>	<u>001</u>	Sep 28, 2018
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A209497</u>	<u>002</u>	Sep 28, 2018
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 50MG BASE</u>	<u>A207655</u>	<u>001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A207655</u>	<u>002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A207655</u>	<u>003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A207655</u>	<u>004</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A207655</u>	<u>005</u>	Nov 29, 2017
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>EQ 50MG BASE</u>	<u>A202939</u>	<u>001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A202939</u>	<u>002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A202939</u>	<u>003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202939</u>	<u>004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A202939</u>	<u>005</u>	May 09, 2017
<u>AB</u>	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204203</u>	<u>001</u>	May 17, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A204203</u>	<u>002</u>	May 17, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204203</u>	<u>003</u>	May 17, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204203</u>	<u>004</u>	May 17, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204203</u>	<u>005</u>	May 17, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A204253</u>	<u>001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204253</u>	<u>002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204253</u>	<u>003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204253</u>	<u>004</u>	Nov 29, 2017
<u>AB</u>	NOVAST LABS	<u>EQ 50MG BASE</u>	<u>A208947</u>	<u>001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208947</u>	<u>002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A208947</u>	<u>003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A208947</u>	<u>004</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A208947</u>	<u>005</u>	Nov 29, 2017
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A090482</u>	<u>001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090482</u>	<u>002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090482</u>	<u>003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090482</u>	<u>004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090482</u>	<u>005</u>	May 09, 2017
<u>AB</u>	PHARMADAX INC	<u>EQ 50MG BASE</u>	<u>A206260</u>	<u>001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A206260</u>	<u>002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A206260</u>	<u>003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206260</u>	<u>004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A206260</u>	<u>005</u>	May 09, 2017
<u>AB</u>	PRINSTON INC	<u>EQ 50MG BASE</u>	<u>A208781</u>	<u>001</u>	Apr 26, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208781</u>	<u>002</u>	Apr 26, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A208781</u>	<u>003</u>	Apr 26, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A208781</u>	<u>004</u>	Apr 26, 2022
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 50MG BASE</u>	<u>A209635</u>	<u>005</u>	Nov 16, 2018
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209635</u>	<u>001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A209635</u>	<u>002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A209635</u>	<u>003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A209635</u>	<u>004</u>	Nov 29, 2017
<u>AB</u>	UNICHEM	<u>EQ 50MG BASE</u>	<u>A215478</u>	<u>001</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A215478</u>	<u>002</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A215478</u>	<u>003</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A215478</u>	<u>004</u>	Aug 15, 2022
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A215478</u>	<u>005</u>	Aug 15, 2022
<u>SEROQUEL XR</u>					
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N022047</u>	<u>001</u>	May 17, 2007
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N022047</u>	<u>005</u>	Aug 11, 2008
<u>AB</u>	+	<u>EQ 200MG BASE</u>	<u>N022047</u>	<u>002</u>	May 17, 2007
<u>AB</u>	+	<u>EQ 300MG BASE</u>	<u>N022047</u>	<u>003</u>	May 17, 2007
<u>AB</u>	+	<u>EQ 400MG BASE</u>	<u>N022047</u>	<u>004</u>	May 17, 2007

PRESCRIPTION DRUG PRODUCT LIST

QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

QUINAPRIL HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A202725 001</u>	Apr 29, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202725 002</u>	Apr 29, 2013
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A202725 003</u>	Apr 29, 2013
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202725 004</u>	Apr 29, 2013
<u>AB</u>	INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A078457 001</u>	Aug 24, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078457 002</u>	Aug 24, 2007
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078457 003</u>	Aug 24, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078457 004</u>	Aug 24, 2007
<u>AB</u>	LUPIN	<u>EQ 5MG BASE</u>	<u>A077690 001</u>	Jun 20, 2006
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077690 002</u>	Jun 20, 2006
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077690 003</u>	Jun 20, 2006
<u>AB</u>	!	<u>EQ 40MG BASE</u>	<u>A077690 004</u>	Jun 20, 2006
<u>AB</u>	PRINSTON INC	<u>EQ 5MG BASE</u>	<u>A205823 001</u>	Sep 15, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205823 002</u>	Sep 15, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205823 003</u>	Sep 15, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205823 004</u>	Sep 15, 2016

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE GLUCONATE

<u>AB</u>	EYWA	<u>324MG</u>	<u>A212589 001</u>	Sep 17, 2021
<u>AB</u>	! SUN PHARM INDUSTRIES	<u>324MG</u>	<u>A089338 001</u>	Feb 11, 1987

QUINIDINE SULFATE

TABLET;ORAL

QUINIDINE SULFATE

!	EPIC PHARMA LLC	200MG	A088072 002	
!		300MG	A088072 001	Sep 26, 1983

QUININE SULFATE

CAPSULE;ORAL

QUALAQUIN

<u>AB</u>	+! SUN PHARM INDUSTRIES	<u>324MG</u>	<u>N021799 001</u>	Aug 12, 2005
<u>AB</u>	AMNEAL PHARMS	<u>324MG</u>	<u>A203729 001</u>	Jul 15, 2015
<u>AB</u>	LUPIN LTD	<u>324MG</u>	<u>A203112 001</u>	Apr 24, 2015
<u>AB</u>	NOVAST LABS	<u>324MG</u>	<u>A204372 001</u>	Jul 22, 2015
<u>AB</u>	TEVA PHARMS	<u>324MG</u>	<u>A091661 001</u>	Sep 28, 2012

QUIZARTINIB DIHYDROCHLORIDE

TABLET;ORAL

VANFLYTA

+	DAIICHI SANKYO INC	EQ 17.7MG BASE	N216993 001	Jul 20, 2023
+	!	EQ 26.5MG BASE	N216993 002	Jul 20, 2023

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

<u>AB</u>	+! WOODWARD	<u>20MG</u>	<u>N020973 002</u>	Aug 19, 1999
<u>AB</u>	ALKEM LABS LTD	<u>20MG</u>	<u>A208644 001</u>	Apr 24, 2018
<u>AB</u>	AMNEAL PHARMS	<u>20MG</u>	<u>A204179 001</u>	Jul 31, 2015
<u>AB</u>	AUROBINDO PHARMA	<u>20MG</u>	<u>A205761 001</u>	Feb 17, 2017
<u>AB</u>	DR REDDYS	<u>20MG</u>	<u>A076824 001</u>	Nov 08, 2013
<u>AB</u>	LANNETT CO INC	<u>20MG</u>	<u>A090678 001</u>	Nov 08, 2013
<u>AB</u>	LUPIN LTD	<u>20MG</u>	<u>A078964 001</u>	Nov 08, 2013
<u>AB</u>	RUBICON	<u>20MG</u>	<u>A204237 001</u>	Nov 18, 2015
<u>AB</u>	TORRENT	<u>20MG</u>	<u>A202376 001</u>	Nov 08, 2013

RADIUM RA-223 DICHLORIDE

SOLUTION;INTRAVENOUS

XOFIGO

+	! BAYER HLTHCARE	162mCi/6ML (27mCi/ML)	N203971 001	May 15, 2013
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PRESCRIPTION DRUG PRODUCT LIST

RALOXIFENE HYDROCHLORIDE

TABLET;ORAL

EVISTA

AB	+ !	LILLY	60MG	N020815	001	Dec 09, 1997
<u>RALOXIFENE HYDROCHLORIDE</u>						
AB		AMNEAL PHARMS	60MG	A208206	001	Apr 08, 2016
AB		AUROBINDO PHARMA	60MG	A204310	001	Aug 28, 2015
AB		CADILA PHARMS LTD	60MG	A211324	001	Aug 18, 2020
AB		GLENMARK PHARMS LTD	60MG	A204491	001	Mar 22, 2016
AB		INVAGEN PHARMS	60MG	A090842	001	Sep 24, 2014
AB		SCIEGEN PHARMS INC	60MG	A206384	001	Oct 12, 2016
AB		TEVA PHARMS USA	60MG	A078193	001	Mar 04, 2014
AB		WATSON LABS INC	60MG	A200825	001	Jan 21, 2015

RALTEGRAVIR POTASSIUM

POWDER;ORAL

ISENTRESS

+ !	MSD SUB MERCK	EQ 100MG BASE/PACKET	N205786	001	Dec 20, 2013
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TABLET;ORAL

ISENTRESS

+ !	MSD SUB MERCK	EQ 400MG BASE	N022145	001	Oct 12, 2007
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ISENTRESS HD

+ !	MSD SUB MERCK	EQ 600MG BASE	N022145	002	May 26, 2017
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TABLET, CHEWABLE;ORAL

ISENTRESS

+	MSD SUB MERCK	EQ 25MG BASE	N203045	001	Dec 21, 2011
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+ !		EQ 100MG BASE	N203045	002	Dec 21, 2011
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RAMELTEON

TABLET;ORAL

RAMELTEON

AB		ACTAVIS LABS FL INC	8MG	A091610	001	Aug 19, 2015
AB		AUROBINDO PHARMA LTD	8MG	A215972	001	Jul 10, 2023
AB		DR REDDYS LABS SA	8MG	A091693	001	Jul 26, 2013
AB		GRANULES	8MG	A213186	001	Aug 21, 2020
AB		I3 PHARMS	8MG	A212650	001	Apr 10, 2020
AB		INNOGENIX	8MG	A215435	001	Aug 24, 2022
AB		MICRO LABS	8MG	A215243	001	Feb 09, 2023
AB		UPSHER SMITH LABS	8MG	A213815	001	Oct 26, 2020
AB		XIROMED	8MG	A216209	001	Nov 25, 2022
AB		ZYDUS PHARMS	8MG	A211567	001	Jul 22, 2019
<u>ROZEREM</u>						
AB	+ !	TAKEDA PHARMS USA	8MG	N021782	001	Jul 22, 2005

RAMIPRIL

CAPSULE;ORAL

ALTACE

AB	+	KING PHARMS LLC	1.25MG	N019901	001	Jan 28, 1991
AB	+		2.5MG	N019901	002	Jan 28, 1991
AB	+		5MG	N019901	003	Jan 28, 1991
AB	+ !		10MG	N019901	004	Jan 28, 1991

RAMIPRIL

AB		ACCORD HLTHCARE	1.25MG	A202392	001	Apr 15, 2014
AB			2.5MG	A202392	002	Apr 15, 2014
AB			5MG	A202392	003	Apr 15, 2014
AB			10MG	A202392	004	Apr 15, 2014
AB		AUROBINDO PHARMA LTD	1.25MG	A091604	001	Jun 08, 2011
AB			2.5MG	A091604	002	Jun 08, 2011
AB			5MG	A091604	003	Jun 08, 2011
AB			10MG	A091604	004	Jun 08, 2011
AB		CHARTWELL MOLECULAR	1.25MG	A078745	001	Jun 18, 2008
AB			2.5MG	A078745	002	Jun 18, 2008
AB			5MG	A078745	003	Jun 18, 2008
AB			10MG	A078745	004	Jun 18, 2008
AB		COREPHARMA	1.25MG	A079116	001	Jun 20, 2008
AB			2.5MG	A079116	002	Jun 20, 2008
AB			5MG	A079116	003	Jun 20, 2008
AB			10MG	A079116	004	Jun 20, 2008
AB		DR REDDYS LABS LTD	1.25MG	A078191	001	Jun 18, 2008
AB			2.5MG	A078191	002	Jun 18, 2008
AB			5MG	A078191	003	Jun 18, 2008
AB			10MG	A078191	004	Jun 18, 2008

PRESCRIPTION DRUG PRODUCT LIST

RAMIPRIL

CAPSULE; ORAL

RAMIPRIL

<u>AB</u>	HIKMA	<u>1.25MG</u>	<u>A077900 001</u>	Jun 18, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A077900 002</u>	Jun 18, 2008
<u>AB</u>		<u>5MG</u>	<u>A077900 003</u>	Jun 18, 2008
<u>AB</u>		<u>10MG</u>	<u>A077900 004</u>	Jun 18, 2008
<u>AB</u>	LUPIN	<u>1.25MG</u>	<u>A077626 001</u>	Jun 09, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A077626 002</u>	Jun 09, 2008
<u>AB</u>		<u>5MG</u>	<u>A077626 003</u>	Jun 09, 2008
<u>AB</u>		<u>10MG</u>	<u>A077626 004</u>	Jun 09, 2008
<u>AB</u>	WATSON LABS	<u>1.25MG</u>	<u>A076549 001</u>	Oct 24, 2005
<u>AB</u>		<u>2.5MG</u>	<u>A076549 002</u>	Oct 24, 2005
<u>AB</u>		<u>10MG</u>	<u>A076549 004</u>	Oct 24, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>1.25MG</u>	<u>A078832 001</u>	Sep 02, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A078832 002</u>	Sep 02, 2008
<u>AB</u>		<u>5MG</u>	<u>A078832 003</u>	Sep 02, 2008
<u>AB</u>		<u>10MG</u>	<u>A078832 004</u>	Sep 02, 2008

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A075742 001</u>	Nov 29, 2000
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075742 002</u>	Nov 29, 2000
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074655 001</u>	Oct 22, 1997
<u>AB</u>	!	<u>EQ 300MG BASE</u>	<u>A074655 002</u>	Oct 22, 1997

SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

<u>AA</u>	LANNETT CO INC	<u>EQ 15MG BASE/ML</u>	<u>A091288 001</u>	Dec 09, 2010
<u>AA</u>	!	<u>EQ 15MG BASE/ML</u>	<u>A077405 001</u>	Sep 21, 2007

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 150MG BASE</u>	<u>A074680 001</u>	Sep 12, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074680 002</u>	Sep 12, 1997
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 150MG BASE</u>	<u>A076705 001</u>	Jul 27, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A076705 002</u>	Jul 27, 2005
<u>AB</u>	GLENMARK PHARMS INC	<u>EQ 150MG BASE</u>	<u>A078542 001</u>	Nov 19, 2008
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A078542 002</u>	Nov 19, 2008
<u>AB</u>	PAR PHARM	<u>EQ 150MG BASE</u>	<u>A075180 001</u>	Jan 28, 1999
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075180 002</u>	Jan 28, 1999
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074467 001</u>	Aug 29, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074467 002</u>	Aug 29, 1997
<u>AB</u>	VKT PHARMA	<u>EQ 150MG BASE</u>	<u>A211289 001</u>	Jan 31, 2019
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A211289 002</u>	Jan 31, 2019

RANOLAZINE

GRANULES, EXTENDED RELEASE; ORAL

ASPRUZYO SPRINKLE

+	SPIL	500MG	N216018 001	Feb 28, 2022
+	!	1GM	N216018 002	Feb 28, 2022

TABLET, EXTENDED RELEASE; ORAL

RANOLAZINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>500MG</u>	<u>A208862 001</u>	May 28, 2019
<u>AB</u>		<u>1GM</u>	<u>A208862 002</u>	May 28, 2019
<u>AB</u>	AJANTA PHARMA LTD	<u>500MG</u>	<u>A210054 001</u>	May 28, 2019
<u>AB</u>		<u>1GM</u>	<u>A210054 002</u>	May 28, 2019
<u>AB</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A209953 001</u>	Nov 30, 2020
<u>AB</u>		<u>1GM</u>	<u>A209953 002</u>	Nov 30, 2020
<u>AB</u>	AUROBINDO PHARMA	<u>500MG</u>	<u>A209081 001</u>	Dec 23, 2022
<u>AB</u>		<u>1GM</u>	<u>A209081 002</u>	Dec 23, 2022
<u>AB</u>	CADILA	<u>500MG</u>	<u>A210188 001</u>	Aug 19, 2019
<u>AB</u>		<u>1GM</u>	<u>A210188 002</u>	Aug 19, 2019
<u>AB</u>	GLENMARK PHARMS LTD	<u>500MG</u>	<u>A211082 001</u>	Jul 05, 2019
<u>AB</u>		<u>1GM</u>	<u>A211082 002</u>	Jul 05, 2019
<u>AB</u>	HETERO LABS LTD III	<u>500MG</u>	<u>A212788 001</u>	May 05, 2022
<u>AB</u>		<u>1GM</u>	<u>A212788 002</u>	May 05, 2022
<u>AB</u>	I3 PHARMS	<u>500MG</u>	<u>A213517 001</u>	Apr 27, 2022
<u>AB</u>		<u>1GM</u>	<u>A213517 002</u>	Apr 27, 2022
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A201046 001</u>	Jul 29, 2013
<u>AB</u>		<u>1GM</u>	<u>A201046 002</u>	Jul 29, 2013
<u>AB</u>	MANKIND PHARMA	<u>500MG</u>	<u>A212284 001</u>	Feb 12, 2020
<u>AB</u>		<u>1GM</u>	<u>A212284 002</u>	Feb 12, 2020
<u>AB</u>	MICRO LABS	<u>500MG</u>	<u>A211745 001</u>	Feb 27, 2020

PRESCRIPTION DRUG PRODUCT LIST

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANOLAZINE

<u>AB</u>		<u>1GM</u>	<u>A211745</u>	<u>002</u>	Feb 27, 2020
<u>AB</u>	NOVAST LABS	<u>500MG</u>	<u>A210668</u>	<u>001</u>	Sep 27, 2023
<u>AB</u>		<u>1GM</u>	<u>A210668</u>	<u>002</u>	Sep 27, 2023
<u>AB</u>	PIRAMAL HLTHCARE UK	<u>1GM</u>	<u>A213085</u>	<u>002</u>	Jul 25, 2023
<u>AB</u>		<u>500MG</u>	<u>A213085</u>	<u>001</u>	Jul 25, 2023
<u>AB</u>	PRAXGEN	<u>500MG</u>	<u>A212781</u>	<u>001</u>	Mar 23, 2020
<u>AB</u>		<u>1GM</u>	<u>A212781</u>	<u>002</u>	Mar 23, 2020
<u>AB</u>	RISING	<u>500MG</u>	<u>A212889</u>	<u>001</u>	Jan 28, 2021
<u>AB</u>		<u>1GM</u>	<u>A212889</u>	<u>002</u>	Jan 28, 2021
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A211829</u>	<u>001</u>	Jun 04, 2019
<u>AB</u>		<u>1GM</u>	<u>A211829</u>	<u>002</u>	Jun 04, 2019
<u>AB</u>	SUN PHARM	<u>500MG</u>	<u>A211707</u>	<u>001</u>	May 28, 2019
<u>AB</u>	!	<u>1GM</u>	<u>A211707</u>	<u>002</u>	May 28, 2019
<u>AB</u>	SUNSHINE	<u>500MG</u>	<u>A211865</u>	<u>001</u>	Mar 23, 2020
<u>AB</u>		<u>1GM</u>	<u>A211865</u>	<u>002</u>	Mar 23, 2020
<u>AB</u>	UNICHEM	<u>500MG</u>	<u>A213083</u>	<u>001</u>	Mar 16, 2023
<u>AB</u>		<u>1GM</u>	<u>A213083</u>	<u>002</u>	Mar 16, 2023
<u>AB</u>	VKT PHARMA	<u>500MG</u>	<u>A214035</u>	<u>001</u>	Jan 19, 2022
<u>AB</u>		<u>1GM</u>	<u>A214035</u>	<u>002</u>	Jan 19, 2022

RASAGILINE MESYLATE

TABLET;ORAL

AZILECT

<u>AB</u>	+	TEVA	<u>EQ 0.5MG BASE</u>	<u>N021641</u>	<u>001</u>	May 16, 2006
<u>AB</u>	+	!	<u>EQ 1MG BASE</u>	<u>N021641</u>	<u>002</u>	May 16, 2006

RASAGILINE MESYLATE

<u>AB</u>		ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A201889</u>	<u>001</u>	Oct 30, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201889</u>	<u>002</u>	Oct 30, 2017
<u>AB</u>		AUROBINDO PHARMA USA	<u>EQ 0.5MG BASE</u>	<u>A201971</u>	<u>001</u>	May 15, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201971</u>	<u>002</u>	May 15, 2017
<u>AB</u>		CHARTWELL RX	<u>EQ 0.5MG BASE</u>	<u>A201892</u>	<u>001</u>	Jul 27, 2018
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201892</u>	<u>002</u>	Jul 27, 2018
<u>AB</u>		INDOCO	<u>EQ 0.5MG BASE</u>	<u>A206153</u>	<u>001</u>	Oct 04, 2019
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206153</u>	<u>002</u>	Oct 04, 2019
<u>AB</u>		MICRO LABS	<u>EQ 0.5MG BASE</u>	<u>A207004</u>	<u>001</u>	Mar 29, 2019
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A207004</u>	<u>002</u>	Mar 29, 2019
<u>AB</u>		ORBION PHARMS	<u>EQ 0.5MG BASE</u>	<u>A201970</u>	<u>001</u>	Mar 15, 2016
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A201970</u>	<u>002</u>	Mar 15, 2016

REGADENOSON

SOLUTION;INTRAVENOUS

LEXISCAN

<u>AP</u>	+	!	ASTELLAS	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>N022161</u>	<u>001</u>	Apr 10, 2008
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REGADENOSON

<u>AP</u>		ACCORD HLTHCARE	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A213236</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>		APOTEX	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A207604</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>		BAXTER HLTHCARE CORP	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A217455</u>	<u>001</u>	May 23, 2023
<u>AP</u>		DR REDDYS	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A213210</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>		EUGIA PHARMA	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A216437</u>	<u>001</u>	Oct 26, 2022
<u>AP</u>		GLAND PHARMA LTD	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A207320</u>	<u>001</u>	Jul 12, 2022
<u>AP</u>		HIKMA	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A215827</u>	<u>001</u>	Feb 02, 2023
<u>AP</u>		HOSPIRA	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A214349</u>	<u>001</u>	Aug 31, 2022
<u>AP</u>		IMS LTD	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A214252</u>	<u>001</u>	May 23, 2022
<u>AP</u>		MEITHEAL	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A212806</u>	<u>001</u>	Apr 11, 2022
<u>AP</u>		MYLAN	<u>0.4MG/5ML (0.08MG/ML)</u>	<u>A213856</u>	<u>001</u>	Apr 04, 2023

REGORAFENIB

TABLET;ORAL

STIVARGA

	+	!	BAYER HLTHCARE	40MG	N203085	001	Sep 27, 2012
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RELUGOLIX

TABLET;ORAL

ORGOVYX

	+	!	SUMITOMO PHARMA	120MG	N214621	001	Dec 18, 2020
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PRESCRIPTION DRUG PRODUCT LIST

REMDESIVIR

POWDER; INTRAVENOUS

VEKLURY

+! GILEAD SCIENCES INC 100MG/VIAL

N214787 001 Oct 22, 2020

SOLUTION; INTRAVENOUS

VEKLURY

+! GILEAD SCIENCES INC 100MG/20ML (5MG/ML)

N214787 002 Oct 22, 2020

REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

REMIFENTANIL HYDROCHLORIDEAP FRESENIUS KABI USAEQ 1MG BASE/VIALA206223 001 Jan 16, 2018APEQ 2MG BASE/VIALA206223 002 Jan 16, 2018APEQ 5MG BASE/VIALA206223 003 Jan 16, 2018AP HIKMAEQ 1MG BASE/VIALA210594 001 Oct 13, 2020APEQ 2MG BASE/VIALA210594 002 Oct 13, 2020APEQ 5MG BASE/VIALA210594 003 Oct 13, 2020ULTIVAAP + MYLAN INSTITUTIONALEQ 1MG BASE/VIALN020630 001 Jul 12, 1996AP +EQ 2MG BASE/VIALN020630 002 Jul 12, 1996AP +!EQ 5MG BASE/VIALN020630 003 Jul 12, 1996REMIMAZOLAM BESYLATE

POWDER; INTRAVENOUS

BYFAVO

+! ACACIA

EQ 20MG BASE/VIAL

N212295 001 Oct 06, 2020

REPAGLINIDE

TABLET; ORAL

REPAGLINIDEAB AUROBINDO PHARMA LTD0.5MGA203820 001 Jan 22, 2014AB1MGA203820 002 Jan 22, 2014AB !2MGA203820 003 Jan 22, 2014AB CHARTWELL RX0.5MGA078555 001 Nov 22, 2013AB1MGA078555 002 Jan 22, 2014AB2MGA078555 003 Jan 22, 2014AB MACLEODS PHARMS LTD0.5MGA207209 001 Mar 22, 2023AB1MGA207209 002 Mar 22, 2023AB2MGA207209 003 Mar 22, 2023AB PADAGIS US0.5MGA201189 001 Jul 17, 2013AB1MGA201189 002 Jan 22, 2014AB2MGA201189 003 Jan 22, 2014AB SUN PHARM INDS INC1MGA077571 002 Jul 11, 2013AB2MGA077571 003 Jul 11, 2013REPOTRECTINIB

CAPSULE; ORAL

AUGTYRO

+! BRISTOL

40MG

N218213 001 Nov 15, 2023

RETAPAMULIN

OINTMENT; TOPICAL

ALTABAX

+! ALMIRALL

1%

N022055 001 Apr 12, 2007

REVEFENACIN

SOLUTION; INHALATION

YUPELRI

+! MYLAN IRELAND LTD

175MCG/3ML

N210598 001 Nov 09, 2018

REZAFUNGIN ACETATE

POWDER; INTRAVENOUS

REZZAYO

+! CIDARA THERAPS

EQ 200MG BASE/VIAL

N217417 001 Mar 22, 2023

RIBAVIRIN

CAPSULE; ORAL

RIBAVIRINAB ! AUROBINDO PHARMA200MGA079117 001 Sep 17, 2009AB ZYDUS PHARMS USA200MGA077224 001 Oct 28, 2005

FOR SOLUTION; INHALATION

RIBAVIRINAN NAVINTA LLC6GM/VIALA207366 001 Oct 06, 2016VIRAZOLEAN +! BAUSCH6GM/VIALN018859 001 Dec 31, 1985

PRESCRIPTION DRUG PRODUCT LIST

RIBAVIRIN

TABLET; ORAL

RIBAVIRIN

AB	AUROBINDO PHARMA	200MG	A079111 001	Sep 17, 2009
AB	SANDOZ	200MG	A077743 001	Oct 03, 2006
AB	ZYDUS PHARMS USA	200MG	A077094 001	Dec 05, 2005

RIBOCICLIB SUCCINATE

TABLET; ORAL

KISQALI

+	NOVARTIS	EQ 200MG BASE	N209092 001	Mar 13, 2017
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RIBOFLAVIN 5'-PHOSPHATE

SOLUTION/DROPS; OPHTHALMIC

PHOTREXA

+	GLAUKOS	0.146%	N203324 001	Apr 15, 2016
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PHOTREXA VISCOUS IN DEXTRAN 20%

+	GLAUKOS	0.146%	N203324 002	Apr 15, 2016
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RIFABUTIN

CAPSULE; ORAL

MYCOBUTIN

AB	+	PFIZER	150MG	N050689 001	Dec 23, 1992
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RIFABUTIN

AB	LUPIN LTD	150MG	A090033 001	Feb 24, 2014
AB	NOVITIUM PHARMA	150MG	A215041 001	Dec 17, 2021

RIFAMPIN

CAPSULE; ORAL

RIFAMPIN

AB	CHARTWELL MOLECULAR	150MG	A065390 001	Mar 28, 2008
AB		300MG	A065390 002	Mar 28, 2008
AB	EPIC PHARMA LLC	150MG	A064150 002	Jan 02, 1998
AB		300MG	A064150 001	May 28, 1997
AB	HIKMA	150MG	A065028 001	Mar 14, 2001
AB		300MG	A065028 002	Mar 14, 2001
AB	LUPIN PHARMS	150MG	A090034 001	Aug 21, 2013
AB	!	300MG	A090034 002	Aug 21, 2013

RIMACTANE

AB	OXFORD PHARMS	300MG	N050429 001	
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INJECTABLE; INJECTION

RIFADIN

AP	+	SANOFI AVENTIS US	600MG/VIAL	N050627 001	May 25, 1989
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RIFAMPIN

AP	EPIC PHARMA LLC	600MG/VIAL	A065502 001	Sep 21, 2010
AP	FRESENIUS KABI USA	600MG/VIAL	A091181 001	Aug 21, 2014
AP	HIKMA	600MG/VIAL	A064217 001	Oct 29, 1999
AP	HIKMA PHARMS	600MG/VIAL	A205039 001	Mar 03, 2016
AP	MYLAN LABS LTD	600MG/VIAL	A065421 001	May 22, 2008

RIFAMYCIN SODIUM

TABLET, DELAYED RELEASE; ORAL

AEMCOLO

+	REDHILL	EQ 194MG BASE	N210910 001	Nov 16, 2018
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RIFAPENTINE

TABLET; ORAL

PRIFTIN

+	SANOFI AVENTIS US	150MG	N021024 001	Jun 22, 1998
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RIFAXIMIN

TABLET; ORAL

XIFAXAN

+	SALIX PHARMS	200MG	N021361 001	May 25, 2004
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+		550MG	N021361 002	Mar 24, 2010
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RILPIVIRINE HYDROCHLORIDE

TABLET; ORAL

EDURANT

+	JANSSEN PRODS	EQ 25MG BASE	N202022 001	May 20, 2011
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PRESCRIPTION DRUG PRODUCT LIST

RILUZOLE

FILM;ORAL

EXSERVAN

+! AQUESTIVE

50MG

N212640 001 Nov 22, 2019

SUSPENSION;ORAL

TIGLUTIK KIT

+! ITALFARMACO SA

50MG/10ML

N209080 001 Sep 05, 2018

TABLET;ORAL

RILUTEK**AB** +! COVIS**50MG****N020599 001** Dec 12, 1995RILUZOLE**AB** ALKEM LABS LTD**50MG****A204048 001** Mar 30, 2016**AB** GLENMARK PHARMS LTD**50MG****A091394 001** Jun 18, 2013**AB** IMPAX LABS**50MG****A076173 001** Jan 29, 2003**AB** KENTON**50MG****A206045 001** Dec 09, 2019**AB** MYLAN PHARMS INC**50MG****A203042 001** Jul 01, 2013**AB** SUN PHARM INDS LTD**50MG****A091417 001** Jun 18, 2013RIMANTADINE HYDROCHLORIDE

TABLET;ORAL

RIMANTADINE HYDROCHLORIDE

! IMPAX LABS

100MG

A076132 001 Aug 30, 2002

RIMEGEPANT SULFATE

TABLET, ORALLY DISINTEGRATING;ORAL

NURTEC ODT

+! PFIZER

EQ 75MG BASE

N212728 001 Feb 27, 2020

RIOCIGUAT

TABLET;ORAL

ADEMPAS**AB** + BAYER HLTHCARE**0.5MG****N204819 001** Oct 08, 2013**AB** +**1MG****N204819 002** Oct 08, 2013**AB** +**1.5MG****N204819 003** Oct 08, 2013**AB** +**2MG****N204819 004** Oct 08, 2013**AB** +!**2.5MG****N204819 005** Oct 08, 2013RIOCIGUAT**AB** MSN**0.5MG****A211135 001** Sep 01, 2022**AB****1MG****A211135 002** Sep 01, 2022**AB****1.5MG****A211135 003** Sep 01, 2022**AB****2MG****A211135 004** Sep 01, 2022**AB****2.5MG****A211135 005** Sep 01, 2022RIPRETINIB

TABLET;ORAL

QINLOCK

+! DECIPHERA PHARMS

50MG

N213973 001 May 15, 2020

RISDIPLAM

FOR SOLUTION;ORAL

EVRYSDI

+! GENENTECH INC

0.75MG/ML

N213535 001 Aug 07, 2020

RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL**AB** +! APIL**35MG****N020835 003** May 25, 2002**AB** +!**150MG****N020835 005** Apr 22, 2008RISEDRONATE SODIUM**AB** APOTEX**35MG****A090877 001** Nov 30, 2015**AB****75MG****A090877 002** Jun 10, 2014**AB****150MG****A090877 003** Jun 10, 2014**AB** AUROBINDO PHARMA**150MG****A206768 001** Oct 21, 2016**AB** AUROBINDO PHARMA**5MG****A200296 001** Nov 30, 2015

LTD

AB**30MG****A200296 002** Nov 30, 2015**AB****35MG****A200296 003** Nov 30, 2015**AB** MACLEODS PHARMS LTD**5MG****A203533 001** Dec 09, 2015**AB****30MG****A203533 002** Dec 09, 2015**AB****35MG****A203533 003** Nov 29, 2016**AB** ORBION PHARMS**30MG****A205280 001** May 13, 2019**AB****35MG****A205280 002** May 13, 2019**AB****5MG****A090886 001** Nov 30, 2015**AB****30MG****A090886 002** Nov 30, 2015**AB****35MG****A090886 003** Nov 30, 2015**AB****75MG****A090886 004** Jun 10, 2014

PRESCRIPTION DRUG PRODUCT LIST

RISEDRONATE SODIUM

TABLET;ORAL

RISEDRONATE SODIUM

<u>AB</u>		<u>150MG</u>	<u>A090886</u>	<u>005</u>	Jun 10, 2014
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A077132</u>	<u>001</u>	Oct 05, 2007
<u>AB</u>		<u>30MG</u>	<u>A077132</u>	<u>002</u>	Oct 05, 2007
<u>AB</u>		<u>35MG</u>	<u>A077132</u>	<u>003</u>	Oct 05, 2007
<u>AB</u>		<u>150MG</u>	<u>A079215</u>	<u>001</u>	Jun 13, 2014

TABLET, DELAYED RELEASE;ORAL

ATELVIA

<u>AB</u>	+! APIL	<u>35MG</u>	<u>N022560</u>	<u>001</u>	Oct 08, 2010
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RISEDRONATE SODIUM

<u>AB</u>	SUN PHARM	<u>35MG</u>	<u>A203925</u>	<u>001</u>	Jul 09, 2019
<u>AB</u>	TEVA PHARMS USA	<u>35MG</u>	<u>A203217</u>	<u>001</u>	May 18, 2015

RISPERIDONE

FOR SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

RYKINDO

+!	SHANDONG LUYE	12.5MG	N212849	001	Jan 13, 2023
+!		25MG	N212849	002	Jan 13, 2023
+!		37.5MG	N212849	003	Jan 13, 2023
+!		50MG	N212849	004	Jan 13, 2023

FOR SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

PERSERIS KIT

+	INDIVIOR	90MG	N210655	001	Jul 27, 2018
+!		120MG	N210655	002	Jul 27, 2018

INJECTABLE;INTRAMUSCULAR

RISPERDAL CONSTA

<u>AP</u>	+ JANSSEN PHARMS	<u>12.5MG/VIAL</u>	<u>N021346</u>	<u>004</u>	Apr 12, 2007
<u>AP</u>	+!	<u>25MG/VIAL</u>	<u>N021346</u>	<u>001</u>	Oct 29, 2003
<u>AP</u>	+	<u>37.5MG/VIAL</u>	<u>N021346</u>	<u>002</u>	Oct 29, 2003
<u>AP</u>	+	<u>50MG/VIAL</u>	<u>N021346</u>	<u>003</u>	Oct 29, 2003

RISPERIDONE

<u>AP</u>	TEVA PHARMS USA INC	<u>12.5MG/VIAL</u>	<u>A214068</u>	<u>001</u>	Dec 05, 2023
<u>AP</u>		<u>25MG/VIAL</u>	<u>A214068</u>	<u>002</u>	Dec 05, 2023
<u>AP</u>		<u>37.5MG/VIAL</u>	<u>A214068</u>	<u>003</u>	Dec 05, 2023
<u>AP</u>		<u>50MG/VIAL</u>	<u>A214068</u>	<u>004</u>	Dec 05, 2023

SOLUTION;ORAL

RISPERDAL

<u>AA</u>	+! JANSSEN PHARMS	<u>1MG/ML</u>	<u>N020588</u>	<u>001</u>	Jun 10, 1996
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RISPERIDONE

<u>AA</u>	AMNEAL PHARMS	<u>1MG/ML</u>	<u>A091384</u>	<u>001</u>	May 25, 2011
<u>AA</u>	AUROBINDO PHARMA LTD	<u>1MG/ML</u>	<u>A078452</u>	<u>001</u>	Sep 04, 2009
<u>AA</u>	CHARTWELL MOLECULAR	<u>1MG/ML</u>	<u>A202386</u>	<u>001</u>	Jan 12, 2015
<u>AA</u>	HIKMA	<u>1MG/ML</u>	<u>A076904</u>	<u>001</u>	Jul 29, 2009
<u>AA</u>	LANNETT CO INC	<u>1MG/ML</u>	<u>A079158</u>	<u>001</u>	Dec 03, 2010
<u>AA</u>	PHARM ASSOC	<u>1MG/ML</u>	<u>A077719</u>	<u>001</u>	Jul 29, 2009
<u>AA</u>	TARO	<u>1MG/ML</u>	<u>A090347</u>	<u>001</u>	Feb 07, 2011
<u>AA</u>	TRIS PHARMA INC	<u>1MG/ML</u>	<u>A079059</u>	<u>001</u>	Dec 12, 2012

SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

UZEDY

+	TEVA	50MG/0.14ML (50MG/0.14ML)	N213586	001	Apr 28, 2023
+		75MG/0.21ML (75MG/0.21ML)	N213586	002	Apr 28, 2023
+		100MG/0.28ML (100MG/0.28ML)	N213586	003	Apr 28, 2023
+		125MG/0.35ML (125MG/0.35ML)	N213586	004	Apr 28, 2023
+		150MG/0.42ML (150MG/0.42ML)	N213586	005	Apr 28, 2023
+		200MG/0.56ML (200MG/0.56ML)	N213586	006	Apr 28, 2023
+!		250MG/0.7ML (250MG/0.7ML)	N213586	007	Apr 28, 2023

TABLET;ORAL

RISPERDAL

<u>AB</u>	+ JANSSEN PHARMS	<u>0.25MG</u>	<u>N020272</u>	<u>008</u>	May 10, 1999
<u>AB</u>	+	<u>0.5MG</u>	<u>N020272</u>	<u>007</u>	Jan 27, 1999
<u>AB</u>	+!	<u>1MG</u>	<u>N020272</u>	<u>001</u>	Dec 29, 1993
<u>AB</u>	+	<u>2MG</u>	<u>N020272</u>	<u>002</u>	Dec 29, 1993
<u>AB</u>	+	<u>3MG</u>	<u>N020272</u>	<u>003</u>	Dec 29, 1993
<u>AB</u>	+	<u>4MG</u>	<u>N020272</u>	<u>004</u>	Dec 29, 1993

RISPERIDONE

<u>AB</u>	AJANTA PHARMA LTD	<u>0.25MG</u>	<u>A201003</u>	<u>001</u>	Aug 24, 2011
<u>AB</u>		<u>0.5MG</u>	<u>A201003</u>	<u>002</u>	Aug 24, 2011
<u>AB</u>		<u>1MG</u>	<u>A201003</u>	<u>003</u>	Aug 24, 2011
<u>AB</u>		<u>2MG</u>	<u>A201003</u>	<u>004</u>	Aug 24, 2011
<u>AB</u>		<u>3MG</u>	<u>A201003</u>	<u>005</u>	Aug 24, 2011

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET; ORAL

RISPERIDONE

<u>AB</u>		<u>4MG</u>	<u>A201003 006</u>	Aug 24, 2011
<u>AB</u>	AMNEAL	<u>0.25MG</u>	<u>A078071 001</u>	Jun 17, 2009
<u>AB</u>		<u>0.5MG</u>	<u>A078071 002</u>	Jun 17, 2009
<u>AB</u>		<u>1MG</u>	<u>A078071 003</u>	Jun 17, 2009
<u>AB</u>		<u>2MG</u>	<u>A078071 004</u>	Jun 17, 2009
<u>AB</u>		<u>3MG</u>	<u>A078071 005</u>	Jun 17, 2009
<u>AB</u>		<u>4MG</u>	<u>A078071 006</u>	Jun 17, 2009
<u>AB</u>	APOTEX INC	<u>0.25MG</u>	<u>A077953 001</u>	Sep 15, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A077953 002</u>	Sep 15, 2008
<u>AB</u>		<u>1MG</u>	<u>A077953 003</u>	Sep 15, 2008
<u>AB</u>		<u>2MG</u>	<u>A077953 004</u>	Sep 15, 2008
<u>AB</u>		<u>3MG</u>	<u>A077953 005</u>	Sep 15, 2008
<u>AB</u>		<u>4MG</u>	<u>A077953 006</u>	Sep 15, 2008
<u>AB</u>	CELLTRION	<u>0.25MG</u>	<u>A078871 001</u>	Oct 09, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078871 002</u>	Oct 09, 2008
<u>AB</u>		<u>1MG</u>	<u>A078871 003</u>	Oct 09, 2008
<u>AB</u>		<u>2MG</u>	<u>A078871 004</u>	Oct 09, 2008
<u>AB</u>		<u>3MG</u>	<u>A078871 005</u>	Oct 09, 2008
<u>AB</u>		<u>4MG</u>	<u>A078871 006</u>	Oct 09, 2008
<u>AB</u>	CHARTWELL MOLECULAR	<u>0.25MG</u>	<u>A077543 001</u>	May 18, 2011
<u>AB</u>		<u>0.5MG</u>	<u>A077543 002</u>	May 18, 2011
<u>AB</u>		<u>1MG</u>	<u>A077543 003</u>	May 18, 2011
<u>AB</u>		<u>2MG</u>	<u>A077543 004</u>	May 18, 2011
<u>AB</u>		<u>3MG</u>	<u>A077543 005</u>	May 18, 2011
<u>AB</u>		<u>4MG</u>	<u>A077543 006</u>	May 18, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>0.25MG</u>	<u>A076879 001</u>	Oct 24, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A076879 002</u>	Oct 24, 2008
<u>AB</u>		<u>1MG</u>	<u>A076879 003</u>	Oct 24, 2008
<u>AB</u>		<u>2MG</u>	<u>A076879 004</u>	Oct 24, 2008
<u>AB</u>		<u>3MG</u>	<u>A076879 005</u>	Oct 24, 2008
<u>AB</u>		<u>4MG</u>	<u>A076879 006</u>	Oct 24, 2008
<u>AB</u>	PRINSTON INC	<u>0.25MG</u>	<u>A077493 001</u>	Nov 29, 2011
<u>AB</u>		<u>0.5MG</u>	<u>A077493 002</u>	Nov 29, 2011
<u>AB</u>		<u>1MG</u>	<u>A077493 003</u>	Nov 29, 2011
<u>AB</u>		<u>2MG</u>	<u>A077493 004</u>	Nov 29, 2011
<u>AB</u>		<u>3MG</u>	<u>A077493 005</u>	Nov 29, 2011
<u>AB</u>		<u>4MG</u>	<u>A077493 006</u>	Nov 29, 2011
<u>AB</u>	RISING	<u>0.25MG</u>	<u>A078269 001</u>	Oct 08, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078269 002</u>	Oct 08, 2008
<u>AB</u>		<u>1MG</u>	<u>A078269 003</u>	Oct 08, 2008
<u>AB</u>		<u>2MG</u>	<u>A078269 004</u>	Oct 08, 2008
<u>AB</u>		<u>3MG</u>	<u>A078269 005</u>	Oct 08, 2008
<u>AB</u>		<u>4MG</u>	<u>A078269 006</u>	Oct 08, 2008
<u>AB</u>	SANDOZ	<u>0.25MG</u>	<u>A078528 001</u>	Oct 16, 2009
<u>AB</u>		<u>0.5MG</u>	<u>A078528 002</u>	Oct 16, 2009
<u>AB</u>		<u>1MG</u>	<u>A078528 003</u>	Oct 16, 2009
<u>AB</u>		<u>2MG</u>	<u>A078528 004</u>	Oct 16, 2009
<u>AB</u>		<u>3MG</u>	<u>A078528 005</u>	Oct 16, 2009
<u>AB</u>		<u>4MG</u>	<u>A078528 006</u>	Oct 16, 2009
<u>AB</u>	TORRENT PHARMS	<u>0.25MG</u>	<u>A079088 001</u>	Oct 30, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A079088 002</u>	Oct 30, 2008
<u>AB</u>		<u>1MG</u>	<u>A079088 003</u>	Oct 30, 2008
<u>AB</u>		<u>2MG</u>	<u>A079088 004</u>	Oct 30, 2008
<u>AB</u>		<u>3MG</u>	<u>A079088 005</u>	Oct 30, 2008
<u>AB</u>		<u>4MG</u>	<u>A079088 006</u>	Oct 30, 2008
<u>AB</u>	WESTMINSTER PHARMS	<u>0.25MG</u>	<u>A078707 001</u>	Dec 29, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078707 002</u>	Dec 29, 2008
<u>AB</u>		<u>1MG</u>	<u>A078707 003</u>	Dec 29, 2008
<u>AB</u>		<u>2MG</u>	<u>A078707 004</u>	Dec 29, 2008
<u>AB</u>		<u>3MG</u>	<u>A078707 005</u>	Dec 29, 2008
<u>AB</u>		<u>4MG</u>	<u>A078707 006</u>	Dec 29, 2008
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.25MG</u>	<u>A078040 001</u>	Oct 16, 2008
<u>AB</u>		<u>0.5MG</u>	<u>A078040 002</u>	Oct 16, 2008
<u>AB</u>		<u>1MG</u>	<u>A078040 003</u>	Oct 16, 2008
<u>AB</u>		<u>2MG</u>	<u>A078040 004</u>	Oct 16, 2008
<u>AB</u>		<u>3MG</u>	<u>A078040 005</u>	Oct 16, 2008
<u>AB</u>		<u>4MG</u>	<u>A078040 006</u>	Oct 16, 2008

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET, ORALLY DISINTEGRATING;ORAL

RISPERIDONE

<u>AB</u>	DR REDDYS LABS LTD	<u>0.5MG</u>	<u>A077328 001</u>	Feb 24, 2009
<u>AB</u>		<u>1MG</u>	<u>A077328 002</u>	Oct 05, 2009
<u>AB</u>		<u>2MG</u>	<u>A077328 003</u>	Feb 24, 2009
<u>AB</u>		<u>3MG</u>	<u>A077328 004</u>	Nov 30, 2009
<u>AB</u>		<u>4MG</u>	<u>A077328 005</u>	Nov 30, 2009
<u>AB</u>	JUBILANT GENERICS	<u>0.5MG</u>	<u>A090839 001</u>	Nov 04, 2011
<u>AB</u>	!	<u>1MG</u>	<u>A090839 002</u>	Nov 04, 2011
<u>AB</u>		<u>2MG</u>	<u>A090839 003</u>	Nov 04, 2011
<u>AB</u>		<u>3MG</u>	<u>A090839 004</u>	Nov 04, 2011
<u>AB</u>		<u>4MG</u>	<u>A090839 005</u>	Nov 04, 2011
<u>AB</u>	PAR PHARM	<u>0.5MG</u>	<u>A077494 002</u>	Apr 30, 2009
<u>AB</u>		<u>1MG</u>	<u>A077494 003</u>	Oct 26, 2009
<u>AB</u>		<u>2MG</u>	<u>A077494 004</u>	Apr 30, 2009
<u>AB</u>		<u>3MG</u>	<u>A077494 005</u>	Apr 30, 2009
<u>AB</u>		<u>4MG</u>	<u>A077494 006</u>	Apr 30, 2009
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A078116 001</u>	Dec 22, 2009
<u>AB</u>		<u>1MG</u>	<u>A078116 002</u>	Dec 22, 2009
<u>AB</u>		<u>2MG</u>	<u>A078116 003</u>	Dec 22, 2009
<u>AB</u>		<u>3MG</u>	<u>A078116 004</u>	Dec 22, 2009
<u>AB</u>		<u>4MG</u>	<u>A078116 005</u>	Dec 22, 2009
<u>AB</u>	SUN PHARM INDS LTD	<u>0.5MG</u>	<u>A077542 001</u>	Aug 06, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078464 001</u>	Apr 08, 2013
<u>AB</u>		<u>1MG</u>	<u>A077542 002</u>	Aug 06, 2010
<u>AB</u>		<u>1MG</u>	<u>A078464 002</u>	Apr 08, 2013
<u>AB</u>		<u>2MG</u>	<u>A077542 003</u>	Aug 06, 2010
<u>AB</u>		<u>2MG</u>	<u>A078464 003</u>	Apr 08, 2013
<u>AB</u>		<u>3MG</u>	<u>A078464 004</u>	Apr 08, 2013
<u>AB</u>		<u>3MG</u>	<u>A078474 001</u>	Aug 06, 2010
<u>AB</u>		<u>4MG</u>	<u>A078464 005</u>	Apr 08, 2013
<u>AB</u>		<u>4MG</u>	<u>A078474 002</u>	Aug 06, 2010
<u>AB</u>	ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A078516 001</u>	May 01, 2009
<u>AB</u>		<u>2MG</u>	<u>A078516 003</u>	May 01, 2009
	PAR PHARM	0.25MG	A077494 001	Apr 30, 2009

RITLACITINIB TOSYLATE

CAPSULE;ORAL

LITFULO

+! PFIZER

EQ 50MG BASE

N215830 001 Jun 23, 2023

RITONAVIR

POWDER;ORAL

NORVIR

+! ABBVIE

100MG/PACKET

N209512 001 Jun 07, 2017

TABLET;ORAL

NORVIRAB +! ABBVIE 100MG N022417 001 Feb 10, 2010RITONAVIRAB AMNEAL 100MG A208890 001 Sep 17, 2018AB AUROBINDO PHARMA LTD 100MG A206614 001 Sep 17, 2018AB CIPLA 100MG A202573 001 Jan 15, 2015AB 100MG A203759 001 May 10, 2022AB HETERO LABS LTD III 100MG A204587 001 Sep 17, 2018RIVAROXABAN

FOR SUSPENSION;ORAL

XARELTO

+! JANSSEN PHARMS

1MG/ML

N215859 001 Dec 20, 2021

TABLET;ORAL

XARELTO

+ JANSSEN PHARMS

2.5MG

N022406 004 Oct 11, 2018

+

10MG

N022406 001 Jul 01, 2011

+

15MG

N022406 002 Nov 04, 2011

+!

20MG

N022406 003 Nov 04, 2011

RIVASTIGMINE

FILM, EXTENDED RELEASE;TRANSDERMAL

EXELONAB + SANDOZ 4.6MG/24HR N022083 001 Jul 06, 2007AB +! 9.5MG/24HR N022083 002 Jul 06, 2007AB + 13.3MG/24HR N022083 005 Aug 31, 2012

PRESCRIPTION DRUG PRODUCT LIST

RIVASTIGMINE

FILM, EXTENDED RELEASE;TRANSDERMAL

RIVASTIGMINE

<u>AB</u>	ALVOGEN	<u>4.6MG/24HR</u>	<u>A204403 001</u>	Sep 03, 2015
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A204403 002</u>	Sep 03, 2015
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A204403 003</u>	Aug 31, 2015
<u>AB</u>	AMNEAL PHARMS	<u>4.6MG/24HR</u>	<u>A207308 001</u>	Jan 08, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A207308 002</u>	Jan 08, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A207308 003</u>	Jan 08, 2019
<u>AB</u>	BRECKENRIDGE	<u>4.6MG/24HR</u>	<u>A209063 001</u>	Nov 26, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A209063 002</u>	Nov 26, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A209063 003</u>	Nov 26, 2019
<u>AB</u>	MYLAN TECHNOLOGIES	<u>4.6MG/24HR</u>	<u>A205622 001</u>	Jun 20, 2018
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A205622 002</u>	Jun 20, 2018
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A205622 003</u>	Jun 20, 2018
<u>AB</u>	ZYDUS PHARMS	<u>4.6MG/24HR</u>	<u>A206318 001</u>	Mar 04, 2019
<u>AB</u>		<u>9.5MG/24HR</u>	<u>A206318 002</u>	Mar 04, 2019
<u>AB</u>		<u>13.3MG/24HR</u>	<u>A206318 003</u>	Mar 04, 2019

RIVASTIGMINE TARTRATE

CAPSULE;ORAL

RIVASTIGMINE TARTRATE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A091689 001</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A091689 002</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A091689 003</u>	Jun 12, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A091689 004</u>	Jun 12, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 1.5MG BASE</u>	<u>A204572 001</u>	Mar 25, 2016
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A204572 002</u>	Mar 25, 2016
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A204572 003</u>	Mar 25, 2016
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A204572 004</u>	Mar 25, 2016
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A203844 001</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A203844 002</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A203844 003</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A203844 004</u>	Feb 13, 2017
<u>AB</u>	CHARTWELL RX	<u>EQ 1.5MG BASE</u>	<u>A207797 001</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A207797 002</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A207797 003</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A207797 004</u>	Sep 28, 2017
<u>AB</u>	! DR REDDYS LABS INC	<u>EQ 1.5MG BASE</u>	<u>A077130 001</u>	Oct 31, 2007
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077130 002</u>	Oct 31, 2007
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077130 003</u>	Oct 31, 2007
<u>AB</u>	!	<u>EQ 6MG BASE</u>	<u>A077130 004</u>	Oct 31, 2007
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A203148 001</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A203148 002</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A203148 003</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A203148 004</u>	Aug 22, 2014
<u>AB</u>	ORBION PHARMS	<u>EQ 1.5MG BASE</u>	<u>A090879 001</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A090879 002</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A090879 003</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A090879 004</u>	Jun 10, 2015
<u>AB</u>	SUN PHARM	<u>EQ 1.5MG BASE</u>	<u>A077131 001</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077131 002</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077131 003</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A077131 004</u>	Oct 22, 2007
<u>AB</u>	WATSON LABS	<u>EQ 1.5MG BASE</u>	<u>A077129 001</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077129 002</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077129 003</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A077129 004</u>	Jan 08, 2008

RIZATRIPTAN BENZOATE

FILM;ORAL

RIZAFILM

+! GENSCO

EQ 10MG BASE

N205394 001 Apr 14, 2023

TABLET;ORAL

MAXALT

<u>AB</u>	+! ORGANON LLC	<u>EQ 10MG BASE</u>	<u>N020864 002</u>	Jun 29, 1998
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RIZATRIPTAN BENZOATE

<u>AB</u>	ALKEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A203269 001</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203269 002</u>	Feb 18, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A202490 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202490 002</u>	Dec 31, 2012
<u>AB</u>	CHARTWELL RX	<u>EQ 5MG BASE</u>	<u>A077526 001</u>	Mar 26, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077526 002</u>	Mar 26, 2013

PRESCRIPTION DRUG PRODUCT LIST

RIZATRIPTAN BENZOATE

TABLET; ORAL

RIZATRIPTAN BENZOATE

<u>AB</u>	CREEKWOOD PHARMS	<u>EQ 5MG BASE</u>	<u>A202047 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202047 002</u>	Dec 31, 2012
<u>AB</u>	GLENMARK GENERICS	<u>EQ 5MG BASE</u>	<u>A201967 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201967 002</u>	Dec 31, 2012
<u>AB</u>	INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A204339 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204339 002</u>	Jul 01, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A203147 001</u>	Feb 11, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203147 002</u>	Feb 11, 2014
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 5MG BASE</u>	<u>A201993 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201993 002</u>	Dec 31, 2012
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A200482 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A200482 002</u>	Dec 31, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 5MG BASE</u>	<u>A077263 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077263 002</u>	Dec 31, 2012

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

<u>AB</u>	+! ORGANON	<u>EQ 10MG BASE</u>	<u>N020865 002</u>	Jun 29, 1998
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RIZATRIPTAN BENZOATE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A203062 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203062 002</u>	Jul 01, 2013
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A201914 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201914 002</u>	Jul 01, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A203146 001</u>	Sep 19, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203146 002</u>	Sep 19, 2014
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A203478 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203478 002</u>	Jul 01, 2013
<u>AB</u>	PANACEA	<u>EQ 5MG BASE</u>	<u>A204722 001</u>	Jan 11, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204722 002</u>	Jan 11, 2017

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ROCURONIUM BROMIDE

<u>AP</u>	CAPLIN	<u>50MG/5ML (10MG/ML)</u>	<u>A216234 001</u>	Mar 02, 2023
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A216234 002</u>	Mar 02, 2023
<u>AP</u>	EUGIA PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A206206 001</u>	Apr 12, 2017
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A206206 002</u>	Apr 12, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/5ML (10MG/ML)</u>	<u>A078651 001</u>	Dec 29, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078651 002</u>	Dec 29, 2008
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A205656 001</u>	Apr 04, 2018
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A205656 002</u>	Apr 04, 2018
<u>AP</u>	HIKMA	<u>50MG/5ML (10MG/ML)</u>	<u>A217034 001</u>	May 24, 2023
<u>AP</u>	HOSPIRA	<u>50MG/5ML (10MG/ML)</u>	<u>A078519 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078519 002</u>	Nov 26, 2008
<u>AP</u>	MEITHEAL	<u>50MG/5ML (10MG/ML)</u>	<u>A213453 001</u>	May 08, 2023
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A213453 002</u>	May 08, 2023
<u>AP</u>	MYLAN INSTITUTIONAL	<u>50MG/5ML (10MG/ML)</u>	<u>A079199 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A079199 002</u>	Nov 26, 2008
<u>AP</u>	PIRAMAL CRITICAL	<u>50MG/5ML (10MG/ML)</u>	<u>A210437 001</u>	Aug 13, 2019
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A210437 002</u>	Aug 13, 2019
<u>AP</u>	PRINSTON INC	<u>50MG/5ML (10MG/ML)</u>	<u>A212668 001</u>	May 26, 2022
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A212668 002</u>	May 26, 2022
<u>AP</u>	SAGENT PHARMS INC	<u>50MG/5ML (10MG/ML)</u>	<u>A091458 001</u>	Jul 28, 2010
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A091458 002</u>	Jul 28, 2010
<u>AP</u>	! SANDOZ	<u>50MG/5ML (10MG/ML)</u>	<u>A079195 001</u>	Dec 05, 2008
<u>AP</u>	!	<u>100MG/10ML (10MG/ML)</u>	<u>A079195 002</u>	Dec 05, 2008
<u>AP</u>	TAMARANG	<u>50MG/5ML (10MG/ML)</u>	<u>A091115 001</u>	Aug 27, 2012
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A091115 002</u>	Aug 27, 2012
<u>AP</u>	WEST WARD PHARM CORP	<u>50MG/5ML (10MG/ML)</u>	<u>A204679 001</u>	Feb 28, 2017
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A204679 002</u>	Feb 28, 2017

ROFLUMILAST

CREAM; TOPICAL

ZORYVE

+! ARCUTIS

0.3%

N215985 001 Jul 29, 2022

FOAM; TOPICAL

ZORYVE

+! ARCUTIS

0.3%

N217242 001 Dec 15, 2023

PRESCRIPTION DRUG PRODUCT LIST

ROFLUMILAST

TABLET; ORAL

DALIRESP

<u>AB</u>	<u>+</u>	ASTRAZENECA	<u>250MCG</u>	<u>N022522</u>	<u>002</u>	Jan 23, 2018
<u>AB</u>	<u>+</u>	!	<u>500MCG</u>	<u>N022522</u>	<u>001</u>	Feb 28, 2011

ROFLUMILAST

<u>AB</u>		ALKEM LABS LTD	<u>250MCG</u>	<u>A212490</u>	<u>001</u>	Apr 18, 2023
<u>AB</u>			<u>500MCG</u>	<u>A212490</u>	<u>002</u>	Apr 18, 2023
<u>AB</u>		AUROBINDO PHARMA LTD	<u>500MCG</u>	<u>A213298</u>	<u>001</u>	Apr 17, 2023
<u>AB</u>		BRECKENRIDGE	<u>250MCG</u>	<u>A208236</u>	<u>002</u>	Oct 19, 2023
<u>AB</u>		HETERO LABS LTD III	<u>250MCG</u>	<u>A208213</u>	<u>002</u>	Apr 18, 2023
<u>AB</u>			<u>500MCG</u>	<u>A208213</u>	<u>001</u>	Nov 23, 2018
<u>AB</u>		MICRO LABS	<u>250MCG</u>	<u>A208180</u>	<u>002</u>	Apr 18, 2023
<u>AB</u>			<u>500MCG</u>	<u>A208180</u>	<u>001</u>	Mar 22, 2019
<u>AB</u>		MSN	<u>250MCG</u>	<u>A208256</u>	<u>001</u>	Sep 07, 2022
<u>AB</u>			<u>500MCG</u>	<u>A208256</u>	<u>002</u>	Sep 07, 2022
<u>AB</u>		MYLAN	<u>500MCG</u>	<u>A208257</u>	<u>001</u>	Jul 13, 2018
<u>AB</u>		PRINSTON INC	<u>500MCG</u>	<u>A208299</u>	<u>001</u>	May 10, 2022
<u>AB</u>		STRIDES PHARMA	<u>250MCG</u>	<u>A208247</u>	<u>002</u>	Jun 20, 2023
<u>AB</u>			<u>500MCG</u>	<u>A208247</u>	<u>001</u>	Mar 30, 2020
<u>AB</u>		ZYDUS PHARMS	<u>250MCG</u>	<u>A208303</u>	<u>002</u>	Apr 18, 2023
<u>AB</u>			<u>500MCG</u>	<u>A208303</u>	<u>001</u>	Feb 10, 2022

ROLAPITANT HYDROCHLORIDE

TABLET; ORAL

VARUBI

<u>+</u>	TERSERA	EQ 90MG BASE		N206500	001	Sep 01, 2015
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ROMIDEPSIN

POWDER; INTRAVENOUS

ISTODAX

<u>AP</u>	<u>+</u>	BRISTOL-MYERS	<u>10MG/VIAL</u>	<u>N022393</u>	<u>001</u>	Nov 05, 2009
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ROMIDEPSIN

<u>AP</u>		FRESENIUS KABI USA	<u>10MG/VIAL</u>	<u>A206254</u>	<u>001</u>	Oct 12, 2021
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ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>EQ 0.25MG BASE</u>	<u>A204022</u>	<u>001</u>	Feb 28, 2017
<u>AB</u>			<u>EQ 0.5MG BASE</u>	<u>A204022</u>	<u>002</u>	Feb 28, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A204022</u>	<u>003</u>	Feb 28, 2017
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A204022</u>	<u>004</u>	Feb 28, 2017
<u>AB</u>			<u>EQ 3MG BASE</u>	<u>A204022</u>	<u>005</u>	Feb 28, 2017
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A204022</u>	<u>006</u>	Feb 28, 2017
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A204022</u>	<u>007</u>	Feb 28, 2017
<u>AB</u>		ALEMBIC LTD	<u>EQ 0.25MG BASE</u>	<u>A090429</u>	<u>001</u>	Mar 24, 2010
<u>AB</u>			<u>EQ 0.5MG BASE</u>	<u>A090429</u>	<u>002</u>	Mar 24, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090429</u>	<u>003</u>	Mar 24, 2010
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A090429</u>	<u>004</u>	Mar 24, 2010
<u>AB</u>			<u>EQ 3MG BASE</u>	<u>A090429</u>	<u>005</u>	Mar 24, 2010
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A090429</u>	<u>006</u>	Mar 24, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090429</u>	<u>007</u>	Mar 24, 2010
<u>AB</u>		CADILA	<u>EQ 0.25MG BASE</u>	<u>A090411</u>	<u>001</u>	Jun 01, 2009
<u>AB</u>			<u>EQ 0.5MG BASE</u>	<u>A090411</u>	<u>002</u>	Jun 01, 2009
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090411</u>	<u>003</u>	Jun 01, 2009
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A090411</u>	<u>004</u>	Jun 01, 2009
<u>AB</u>			<u>EQ 3MG BASE</u>	<u>A090411</u>	<u>005</u>	Jun 01, 2009
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A090411</u>	<u>006</u>	Jun 01, 2009
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090411</u>	<u>007</u>	Jun 01, 2009
<u>AB</u>		CHARTWELL RX	<u>EQ 0.25MG BASE</u>	<u>A079050</u>	<u>001</u>	May 29, 2008
<u>AB</u>			<u>EQ 0.5MG BASE</u>	<u>A079050</u>	<u>002</u>	May 29, 2008
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A079050</u>	<u>003</u>	May 29, 2008
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A079050</u>	<u>004</u>	May 29, 2008
<u>AB</u>			<u>EQ 3MG BASE</u>	<u>A079050</u>	<u>005</u>	May 29, 2008
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A079050</u>	<u>006</u>	May 29, 2008
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A079050</u>	<u>007</u>	May 29, 2008
<u>AB</u>		GLENMARK GENERICS	<u>EQ 0.25MG BASE</u>	<u>A090135</u>	<u>001</u>	Feb 25, 2010
<u>AB</u>			<u>EQ 0.5MG BASE</u>	<u>A090135</u>	<u>002</u>	Feb 25, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090135</u>	<u>003</u>	Feb 25, 2010
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A090135</u>	<u>004</u>	Feb 25, 2010
<u>AB</u>			<u>EQ 3MG BASE</u>	<u>A090135</u>	<u>005</u>	Feb 25, 2010
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A090135</u>	<u>006</u>	Feb 25, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090135</u>	<u>007</u>	Feb 25, 2010

PRESCRIPTION DRUG PRODUCT LIST

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>	MLV	<u>EQ 0.25MG BASE</u>	<u>A079165 001</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A079165 002</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A079165 003</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A079165 004</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A079165 005</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A079165 006</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A079165 007</u>	Feb 07, 2012
<u>AB</u>	ORBION PHARMS	<u>EQ 0.25MG BASE</u>	<u>A079229 001</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A079229 002</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A079229 003</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A079229 004</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A079229 005</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A079229 006</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A079229 007</u>	Nov 28, 2012
<u>AB</u>	! PRINSTON INC	<u>EQ 0.25MG BASE</u>	<u>A078110 001</u>	May 05, 2008
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A078110 002</u>	May 05, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A078110 003</u>	May 05, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A078110 004</u>	May 05, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A078110 005</u>	May 05, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A078110 006</u>	May 05, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078110 007</u>	Jul 11, 2008

TABLET, EXTENDED RELEASE; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 2MG BASE</u>	<u>A090869 001</u>	May 17, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A090869 002</u>	May 17, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A090869 003</u>	May 17, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A090869 004</u>	May 17, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A090869 005</u>	May 17, 2012
<u>AB</u>	ALEMBIC	<u>EQ 2MG BASE</u>	<u>A202786 001</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A202786 002</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A202786 003</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A202786 004</u>	Apr 22, 2013
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A202786 005</u>	Apr 22, 2013
<u>AB</u>	CHARTWELL RX	<u>EQ 2MG BASE</u>	<u>A091395 001</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A091395 002</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A091395 003</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A091395 004</u>	Aug 27, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A091395 005</u>	Aug 27, 2012
<u>AB</u>	! DR REDDYS LABS LTD	<u>EQ 2MG BASE</u>	<u>A201576 001</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201576 002</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201576 003</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A201576 004</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A201576 005</u>	Jun 06, 2012
<u>AB</u>	PRINSTON INC	<u>EQ 2MG BASE</u>	<u>A204413 001</u>	May 11, 2022
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A204413 002</u>	May 11, 2022
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A204413 003</u>	May 11, 2022
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A204413 004</u>	May 11, 2022
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A204413 005</u>	May 11, 2022
<u>AB</u>	SANDOZ INC	<u>EQ 2MG BASE</u>	<u>A201047 001</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201047 003</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201047 004</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A201047 005</u>	Jun 06, 2012
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A201047 006</u>	Jun 06, 2012

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

NAROPIN

<u>AP</u>	+	FRESENIUS KABI USA	<u>20MG/10ML (2MG/ML)</u>	<u>N020533 001</u>	May 01, 1998
<u>AP</u>	+		<u>40MG/20ML (2MG/ML)</u>	<u>N020533 002</u>	Sep 24, 1996
<u>AP</u>	+		<u>100MG/20ML (5MG/ML)</u>	<u>N020533 003</u>	May 01, 1998
<u>AP</u>	+		<u>100MG/10ML (10MG/ML)</u>	<u>N020533 005</u>	Sep 24, 1996
<u>AP</u>	+		<u>150MG/20ML (7.5MG/ML)</u>	<u>N020533 004</u>	Sep 24, 1996
<u>AP</u>	+		<u>150MG/30ML (5MG/ML)</u>	<u>N020533 008</u>	Sep 24, 1996
<u>AP</u>	+		<u>200MG/100ML (2MG/ML)</u>	<u>N020533 006</u>	Sep 24, 1996
<u>AP</u>	+		<u>200MG/20ML (10MG/ML)</u>	<u>N020533 011</u>	Sep 24, 1996
<u>AP</u>	+		<u>400MG/200ML (2MG/ML)</u>	<u>N020533 007</u>	Sep 24, 1996
<u>AP</u>	+		<u>500MG/100ML (5MG/ML)</u>	<u>N020533 009</u>	Jan 04, 2011
<u>AP</u>	+		<u>1GM/200ML (5MG/ML)</u>	<u>N020533 010</u>	Jan 04, 2011

PRESCRIPTION DRUG PRODUCT LIST

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

ROPIVACAINE HYDROCHLORIDE

AP	AMNEAL	<u>200MG/100ML (2MG/ML)</u>	<u>A216605 001</u>	Mar 08, 2023
AP		<u>400MG/200ML (2MG/ML)</u>	<u>A216605 002</u>	Mar 08, 2023
AP		<u>500MG/100ML (5MG/ML)</u>	<u>A216605 003</u>	Mar 07, 2023
AP		<u>1GM/200ML (5MG/ML)</u>	<u>A216605 004</u>	Mar 07, 2023
AP	CAPLIN	<u>40MG/20ML (2MG/ML)</u>	<u>A212808 001</u>	Apr 09, 2020
AP		<u>100MG/20ML (5MG/ML)</u>	<u>A212808 002</u>	Apr 09, 2020
AP		<u>150MG/30ML (5MG/ML)</u>	<u>A212808 003</u>	Apr 09, 2020
AP		<u>200MG/20ML (10MG/ML)</u>	<u>A212808 004</u>	Apr 09, 2020
AP	EUGIA PHARMA	<u>40MG/20ML (2MG/ML)</u>	<u>A205612 001</u>	Jul 13, 2016
AP		<u>100MG/20ML (5MG/ML)</u>	<u>A205612 003</u>	Jul 13, 2016
AP		<u>100MG/10ML (10MG/ML)</u>	<u>A205612 006</u>	Jul 13, 2016
AP		<u>150MG/30ML (5MG/ML)</u>	<u>A205612 004</u>	Jul 13, 2016
AP		<u>150MG/20ML (7.5MG/ML)</u>	<u>A205612 005</u>	Jul 13, 2016
AP		<u>200MG/100ML (2MG/ML)</u>	<u>A205612 002</u>	Jul 13, 2016
AP		<u>200MG/20ML (10MG/ML)</u>	<u>A205612 007</u>	Jul 13, 2016
AP	GLAND PHARMA LTD	<u>200MG/100ML (2MG/ML)</u>	<u>A210102 001</u>	Aug 18, 2022
AP		<u>400MG/200ML (2MG/ML)</u>	<u>A210102 002</u>	Aug 18, 2022
AP	HIKMA	<u>40MG/20ML (2MG/ML)</u>	<u>A214074 001</u>	Jul 20, 2020
AP		<u>150MG/30ML (5MG/ML)</u>	<u>A214074 002</u>	Jul 20, 2020
AP		<u>150MG/20ML (7.5MG/ML)</u>	<u>A214074 003</u>	Jul 20, 2020
AP		<u>200MG/20ML (10MG/ML)</u>	<u>A214074 004</u>	Jul 20, 2020
AP	HOSPIRA	<u>20MG/10ML (2MG/ML)</u>	<u>A090194 001</u>	Sep 23, 2014
AP		<u>40MG/20ML (2MG/ML)</u>	<u>A090194 005</u>	Sep 23, 2014
AP		<u>100MG/10ML (10MG/ML)</u>	<u>A090194 004</u>	Sep 23, 2014
AP		<u>150MG/30ML (5MG/ML)</u>	<u>A090194 002</u>	Sep 23, 2014
AP		<u>150MG/20ML (7.5MG/ML)</u>	<u>A090194 003</u>	Sep 23, 2014
AP		<u>200MG/20ML (10MG/ML)</u>	<u>A090194 006</u>	Sep 23, 2014
AP	INFORLIFE	<u>200MG/100ML (2MG/ML)</u>	<u>A206166 001</u>	Jun 11, 2018
AP		<u>400MG/200ML (2MG/ML)</u>	<u>A206166 002</u>	Jun 11, 2018
AP		<u>500MG/100ML (5MG/ML)</u>	<u>A206166 003</u>	Jun 11, 2018
AP		<u>1GM/200ML (5MG/ML)</u>	<u>A206166 004</u>	Jun 11, 2018
AP	MYLAN LABS LTD	<u>200MG/100ML (2MG/ML)</u>	<u>A206091 001</u>	Oct 26, 2023
AP		<u>400MG/200ML (2MG/ML)</u>	<u>A206091 002</u>	Oct 26, 2023
AP	NAVINTA LLC	<u>150MG/30ML (5MG/ML)</u>	<u>A078601 002</u>	Jul 17, 2014
AP		<u>200MG/20ML (10MG/ML)</u>	<u>A078601 003</u>	Jul 17, 2014
AP	RISING	<u>200MG/100ML (2MG/ML)</u>	<u>A204636 001</u>	Mar 16, 2018
AP		<u>400MG/200ML (2MG/ML)</u>	<u>A204636 002</u>	Mar 16, 2018
AP		<u>150MG/30ML (5MG/ML)</u>	<u>A203955 001</u>	Apr 11, 2016
AP	SOMERSET THERAPS LLC	<u>20MG/10ML (2MG/ML)</u>	<u>A207636 001</u>	Jun 15, 2018
AP		<u>40MG/20ML (2MG/ML)</u>	<u>A207636 002</u>	Jun 15, 2018
AP		<u>100MG/20ML (5MG/ML)</u>	<u>A207636 003</u>	Jun 15, 2018
AP		<u>100MG/10ML (10MG/ML)</u>	<u>A207636 006</u>	Jun 15, 2018
AP		<u>150MG/30ML (5MG/ML)</u>	<u>A207636 004</u>	Jun 15, 2018
AP		<u>150MG/20ML (7.5MG/ML)</u>	<u>A207636 005</u>	Jun 15, 2018
AP		<u>200MG/20ML (10MG/ML)</u>	<u>A207636 007</u>	Jun 15, 2018

ROSUVASTATIN CALCIUM

TABLET; ORAL

CRESTOR

AB	+ IPR	<u>EQ 5MG BASE</u>	<u>N021366 002</u>	Aug 12, 2003
AB	+	<u>EQ 10MG BASE</u>	<u>N021366 003</u>	Aug 12, 2003
AB	+	<u>EQ 20MG BASE</u>	<u>N021366 004</u>	Aug 12, 2003
AB	+!	<u>EQ 40MG BASE</u>	<u>N021366 005</u>	Aug 12, 2003

ROSUVASTATIN CALCIUM

AB	ACCORD HLTHCARE	<u>EQ 5MG BASE</u>	<u>A206434 001</u>	Oct 31, 2016
AB		<u>EQ 10MG BASE</u>	<u>A206434 002</u>	Oct 31, 2016
AB		<u>EQ 20MG BASE</u>	<u>A206434 003</u>	Oct 31, 2016
AB		<u>EQ 40MG BASE</u>	<u>A206434 004</u>	Oct 31, 2016
AB	ALKEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A206465 001</u>	Mar 21, 2017
AB		<u>EQ 10MG BASE</u>	<u>A206465 002</u>	Mar 21, 2017
AB		<u>EQ 20MG BASE</u>	<u>A206465 003</u>	Mar 21, 2017
AB		<u>EQ 40MG BASE</u>	<u>A206465 004</u>	Mar 21, 2017
AB	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A079170 001</u>	Jul 19, 2016
AB		<u>EQ 10MG BASE</u>	<u>A079170 002</u>	Jul 19, 2016
AB		<u>EQ 20MG BASE</u>	<u>A079170 003</u>	Jul 19, 2016
AB		<u>EQ 40MG BASE</u>	<u>A079170 004</u>	Jul 19, 2016
AB	BIOCON PHARMA	<u>EQ 5MG BASE</u>	<u>A207752 001</u>	Oct 31, 2016
AB		<u>EQ 10MG BASE</u>	<u>A207752 002</u>	Oct 31, 2016

PRESCRIPTION DRUG PRODUCT LIST

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207752 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207752 004</u>	Oct 31, 2016
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A207453 001</u>	Nov 23, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207453 002</u>	Nov 23, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207453 003</u>	Nov 23, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207453 004</u>	Nov 23, 2016
<u>AB</u>	CHANGZHOU PHARM	<u>EQ 5MG BASE</u>	<u>A207408 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207408 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207408 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207408 004</u>	Oct 31, 2016
<u>AB</u>	CHARTWELL RX	<u>EQ 5MG BASE</u>	<u>A079168 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079168 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079168 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079168 004</u>	Jul 19, 2016
<u>AB</u>	GLENMARK PHARMS	<u>EQ 5MG BASE</u>	<u>A079172 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079172 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079172 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079172 004</u>	Jul 19, 2016
<u>AB</u>	HETERO LABS LTD V	<u>EQ 5MG BASE</u>	<u>A207616 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207616 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207616 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207616 004</u>	Oct 31, 2016
<u>AB</u>	LUPIN	<u>EQ 5MG BASE</u>	<u>A205587 001</u>	Jul 31, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205587 002</u>	Jul 31, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205587 003</u>	Jul 31, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205587 004</u>	Jul 31, 2017
<u>AB</u>	MSN	<u>EQ 5MG BASE</u>	<u>A208898 001</u>	Nov 22, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A208898 002</u>	Nov 22, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A208898 003</u>	Nov 22, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208898 004</u>	Nov 22, 2017
<u>AB</u>	RENATA	<u>EQ 5MG BASE</u>	<u>A207062 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207062 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207062 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207062 004</u>	Oct 31, 2016
<u>AB</u>	SANDOZ	<u>EQ 5MG BASE</u>	<u>A079171 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079171 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079171 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079171 004</u>	Jul 19, 2016
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 5MG BASE</u>	<u>A206381 001</u>	Apr 24, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A206381 002</u>	Apr 24, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A206381 003</u>	Apr 24, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A206381 004</u>	Apr 24, 2019
<u>AB</u>	SHANDONG	<u>EQ 5MG BASE</u>	<u>A207375 001</u>	May 07, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207375 002</u>	May 07, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207375 003</u>	May 07, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207375 004</u>	May 07, 2019
<u>AB</u>	SUN PHARM	<u>EQ 5MG BASE</u>	<u>A079169 001</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079169 002</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079169 003</u>	Jul 19, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079169 004</u>	Jul 19, 2016
<u>AB</u>	TORRENT	<u>EQ 5MG BASE</u>	<u>A201619 001</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201619 002</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A201619 003</u>	Oct 31, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A201619 004</u>	Oct 31, 2016
<u>AB</u>	UMEDICA	<u>EQ 5MG BASE</u>	<u>A207626 001</u>	Apr 09, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207626 002</u>	Apr 09, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A207626 003</u>	Apr 09, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A207626 004</u>	Apr 09, 2019
<u>AB</u>	WATSON LABS INC	<u>EQ 5MG BASE</u>	<u>A079167 001</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079167 002</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A079167 003</u>	Apr 29, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A079167 004</u>	Apr 29, 2016
<u>AB</u>	ZHEJIANG YONGTAI	<u>EQ 5MG BASE</u>	<u>A212059 001</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A212059 002</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A212059 003</u>	Nov 04, 2019
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A212059 004</u>	Nov 04, 2019

PRESCRIPTION DRUG PRODUCT LIST

ROTIGOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NEUPRO

+	UCB INC	1MG/24HR	N021829	004	Apr 02, 2012
+	!	2MG/24HR	N021829	001	May 09, 2007
+		3MG/24HR	N021829	005	Apr 02, 2012
+		4MG/24HR	N021829	002	May 09, 2007
+		6MG/24HR	N021829	003	May 09, 2007
+		8MG/24HR	N021829	006	Apr 02, 2012

RUBIDIUM CHLORIDE RB-82

INJECTABLE; INJECTION

CARDIOGEN-82

BRACCO

N/A

N019414 001 Dec 29, 1989

SOLUTION; INTRAVENOUS

RUBY-FILL

JUBILANT

N/A

N202153 001 Sep 30, 2016

RUCAPARIB CAMSYLATE

TABLET; ORAL

RUBRACA

+	ZR PHARMA	EQ 200MG BASE	N209115	001	Dec 19, 2016
+		EQ 250MG BASE	N209115	003	May 01, 2017
+	!	EQ 300MG BASE	N209115	002	Dec 19, 2016

RUFINAMIDE

SUSPENSION; ORAL

BANZEL**AB** +! EISAI INC **40MG/ML** **N201367 001** Mar 03, 2011RUFINAMIDE

AB	ALKEM LABS LTD	40MG/ML	A213410	001	Feb 23, 2021
AB	AUROBINDO PHARMA	40MG/ML	A216549	001	Oct 26, 2022
AB	BIONPHARMA	40MG/ML	A211388	001	Apr 23, 2019
AB	CHARTWELL RX	40MG/ML	A214009	001	Nov 07, 2022
AB	HETERO LABS LTD III	40MG/ML	A216841	001	Dec 05, 2023
AB	HIKMA	40MG/ML	A207363	001	Apr 23, 2019
AB	LUPIN LTD	40MG/ML	A213457	001	Dec 18, 2020

TABLET; ORAL

BANZEL**AB** + EISAI INC **200MG** **N021911 002** Nov 14, 2008**AB** +! **400MG** **N021911 003** Nov 14, 2008RUFINAMIDE

AB	AUROBINDO PHARMA	200MG	A217230	001	Jun 16, 2023
AB		400MG	A217230	002	Jun 16, 2023
AB	GLENMARK PHARMS LTD	200MG	A205075	001	May 16, 2016
AB		400MG	A205075	002	May 16, 2016
AB	HETERO LABS LTD III	200MG	A204993	001	May 11, 2021
AB		400MG	A204993	002	May 11, 2021
AB	HIKMA	200MG	A204988	001	May 16, 2016
AB		400MG	A204988	002	May 16, 2016
AB	LUPIN LTD	200MG	A204964	002	Aug 17, 2022
AB		400MG	A204964	003	Aug 17, 2022
AB	MICRO LABS	200MG	A216688	001	May 15, 2023
AB		400MG	A216688	002	May 15, 2023
AB	MYLAN	200MG	A205095	001	May 16, 2016
AB		400MG	A205095	002	May 16, 2016
	LUPIN LTD	100MG	A204964	001	Aug 17, 2022

RUXOLITINIB PHOSPHATE

CREAM; TOPICAL

OPZELURA

+! INCYTE CORP EQ 1.5% BASE N215309 001 Sep 21, 2021

TABLET; ORAL

JAKAFI

+	INCYTE CORP	EQ 5MG BASE	N202192	001	Nov 16, 2011
+		EQ 10MG BASE	N202192	002	Nov 16, 2011
+		EQ 15MG BASE	N202192	003	Nov 16, 2011
+		EQ 20MG BASE	N202192	004	Nov 16, 2011
+	!	EQ 25MG BASE	N202192	005	Nov 16, 2011

PRESCRIPTION DRUG PRODUCT LIST

SACUBITRIL; VALSARTAN

TABLET; ORAL

ENTRESTO

+	NOVARTIS PHARMS CORP	24MG;26MG	N207620 001	Jul 07, 2015
+		49MG;51MG	N207620 002	Jul 07, 2015
+	!	97MG;103MG	N207620 003	Jul 07, 2015

SAFINAMIDE MESYLATE

TABLET; ORAL

XADAGO

+	MDD US	EQ 50MG BASE	N207145 001	Mar 21, 2017
+	!	EQ 100MG BASE	N207145 002	Mar 21, 2017

SALMETEROL XINAFOATE

POWDER; INHALATION

SEREVENT

+	GLAXOSMITHKLINE	EQ 0.05MG BASE/INH	N020692 001	Sep 19, 1997
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SAPROPTERIN DIHYDROCHLORIDE

POWDER; ORAL

KUVAN

<u>AB</u>	+	BIOMARIN PHARM	<u>100MG/PACKET</u>	<u>N205065 001</u>	Dec 19, 2013
<u>AB</u>	+		<u>500MG/PACKET</u>	<u>N205065 002</u>	Oct 27, 2015

SAPROPTERIN DIHYDROCHLORIDE

<u>AB</u>		ANNORA PHARMA	<u>100MG/PACKET</u>	<u>A215420 001</u>	Aug 18, 2022
<u>AB</u>			<u>500MG/PACKET</u>	<u>A215420 002</u>	Aug 18, 2022
<u>AB</u>		DR REDDYS	<u>100MG/PACKET</u>	<u>A209452 001</u>	Mar 30, 2021
<u>AB</u>			<u>500MG/PACKET</u>	<u>A215798 001</u>	May 13, 2022
<u>AB</u>		PAR PHARM INC	<u>100MG/PACKET</u>	<u>A207207 001</u>	Aug 20, 2019
<u>AB</u>			<u>500MG/PACKET</u>	<u>A210027 001</u>	Aug 20, 2019

TABLET; ORAL

KUVAN

<u>AB</u>	+	BIOMARIN PHARM	<u>100MG</u>	<u>N022181 001</u>	Dec 13, 2007
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SAPROPTERIN DIHYDROCHLORIDE

<u>AB</u>		ANNORA PHARMA	<u>100MG</u>	<u>A215534 001</u>	Aug 23, 2022
<u>AB</u>		DR REDDYS	<u>100MG</u>	<u>A207685 001</u>	Jun 15, 2021
<u>AB</u>		PAR PHARM INC	<u>100MG</u>	<u>A207200 001</u>	May 10, 2019

SARECYCLINE HYDROCHLORIDE

TABLET; ORAL

SEYSARA

+	ALMIRALL	EQ 60MG BASE	N209521 001	Oct 01, 2018
+		EQ 100MG BASE	N209521 002	Oct 01, 2018
+	!	EQ 150MG BASE	N209521 003	Oct 01, 2018

SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

SAXAGLIPTIN

<u>AB</u>		AMNEAL	<u>EQ 2.5MG BASE</u>	<u>A205941 001</u>	Jul 31, 2023
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A205941 002</u>	Jul 31, 2023
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 2.5MG BASE</u>	<u>A205972 001</u>	Jul 31, 2023
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A205972 002</u>	Jul 31, 2023
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A205994 001</u>	Jul 31, 2023
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A205994 002</u>	Jul 31, 2023
<u>AB</u>		MYLAN	<u>EQ 2.5MG BASE</u>	<u>A205980 001</u>	Jul 31, 2023
<u>AB</u>	!		<u>EQ 5MG BASE</u>	<u>A205980 002</u>	Jul 31, 2023
<u>AB</u>		SUN PHARM	<u>EQ 2.5MG BASE</u>	<u>A206078 001</u>	Jul 31, 2023
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A206078 002</u>	Jul 31, 2023

SCOPOLAMINE

SYSTEM; TRANSDERMAL

SCOPOLAMINE

<u>AB</u>		ACTAVIS LABS UT INC	<u>1MG/72HR</u>	<u>A208769 001</u>	Jan 10, 2022
<u>AB</u>		MYLAN TECHNOLOGIES	<u>1MG/72HR</u>	<u>A203753 001</u>	Jun 19, 2019
<u>AB</u>		PADAGIS US	<u>1MG/72HR</u>	<u>A078830 001</u>	Jan 30, 2015
<u>AB</u>		RICONPHARMA LLC	<u>1MG/72HR</u>	<u>A212342 001</u>	Nov 24, 2020

TRANSDERM SCOP

<u>AB</u>	+	BAXTER HLTHCARE CORP	<u>1MG/72HR</u>	<u>N017874 001</u>	
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PRESCRIPTION DRUG PRODUCT LIST

SECNIDAZOLE

GRANULE; ORAL

SOLOSEC

+! LUPIN

2GM/PACKET

N209363 001 Sep 15, 2017

SECRETIN SYNTHETIC HUMAN

FOR SOLUTION; INTRAVENOUS

CHIRHOSTIM

+! CHIRHOCLIN

16MCG/VIAL

N021256 001 Apr 09, 2004

+

40MCG/VIAL

N021256 002 Jun 21, 2007

SELEGILINE

FILM, EXTENDED RELEASE; TRANSDERMAL

EMSAM

+! SOMERSET

6MG/24HR

N021336 001 Feb 27, 2006

+

9MG/24HR

N021336 002 Feb 27, 2006

+

12MG/24HR

N021336 003 Feb 27, 2006

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

SELEGILINE HYDROCHLORIDEAB ! APOTEX5MGA075321 001 Dec 04, 1998AB NOVITIUM PHARMA5MGA075352 001 Nov 30, 1998AB RISING5MGA206803 001 Apr 02, 2019

TABLET; ORAL

SELEGILINE HYDROCHLORIDEAB ! APOTEX INC5MGA074871 001 Jun 06, 1997AB I3 PHARMS5MGA074672 001 Apr 01, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

ZELAPAR

+! BAUSCH

1.25MG

N021479 001 Jun 14, 2006

SELENIOS ACID

SOLUTION; INTRAVENOUS

SELENIOS ACID

+! AM REGENT

EQ 12MCG SELENIUM/2ML (EQ 6MCG
SELENIUM/ML)

N209379 003 Aug 30, 2021

+!

EQ 60MCG SELENIUM/ML (EQ 60MCG
SELENIUM/ML)

N209379 002 Jan 25, 2021

+!

EQ 600MCG SELENIUM/10ML (EQ 60MCG
SELENIUM/ML)

N209379 001 Apr 30, 2019

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

SELENIUM SULFIDE

! PADAGIS US

2.5%

A089996 001 Jan 10, 1991

SELEXIPAG

POWDER; INTRAVENOUS

UPTRAVI

+! ACTELION

1.8MG/VIAL

N214275 001 Jul 29, 2021

TABLET; ORAL

SELEXIPAGAB ALEMBIC0.2MGA214414 001 Oct 11, 2023AB0.4MGA214414 002 Oct 11, 2023AB0.6MGA214414 003 Oct 11, 2023AB0.8MGA214414 004 Oct 11, 2023AB1.4MGA214414 005 Oct 11, 2023AB1.6MGA214414 006 Oct 11, 2023UPTRAVIAB + ACTELION0.2MGN207947 001 Dec 21, 2015AB +!0.4MGN207947 002 Dec 21, 2015AB +0.6MGN207947 003 Dec 21, 2015AB +0.8MGN207947 004 Dec 21, 2015AB +1.4MGN207947 007 Dec 21, 2015AB +1.6MGN207947 008 Dec 21, 2015

+

1MG

N207947 005 Dec 21, 2015

+

1.2MG

N207947 006 Dec 21, 2015

SELINEXOR

TABLET; ORAL

XPOVIO

+ KARYOPHARM THERAPS

20MG

N212306 001 Jul 03, 2019

+

40MG

N212306 002 Apr 15, 2021

+

50MG

N212306 003 Apr 15, 2021

+!

60MG

N212306 004 Apr 15, 2021

PRESCRIPTION DRUG PRODUCT LIST

SELPERCATINIB

CAPSULE; ORAL

RETEVMO

+	LOXO ONCOL ELI LILLY	40MG	N213246 001	May 08, 2020
+	!	80MG	N213246 002	May 08, 2020

SELUMETINIB SULFATE

CAPSULE; ORAL

KOSELUGO

+	ASTRAZENECA	EQ 10MG BASE	N213756 001	Apr 10, 2020
+	!	EQ 25MG BASE	N213756 002	Apr 10, 2020

SEMAGLUTIDE

SOLUTION; SUBCUTANEOUS

OZEMPIC

+	!	NOVO	2MG/1.5ML (1.34MG/ML)	N209637 001	Dec 05, 2017
+	!		2MG/3ML (0.68MG/ML)	N209637 004	Oct 06, 2022
+	!		4MG/3ML (1.34MG/ML)	N209637 002	Apr 09, 2019
+	!		8MG/3ML (2.68MG/ML)	N209637 003	Mar 28, 2022

WEGOVY

+	!	NOVO	0.25MG/0.5ML (0.25MG/0.5ML)	N215256 001	Jun 04, 2021
+	!		0.5MG/0.5ML (0.5MG/0.5ML)	N215256 002	Jun 04, 2021
+	!		1MG/0.5ML (1MG/0.5ML)	N215256 003	Jun 04, 2021
+	!		1.7MG/0.75ML (1.7MG/0.75ML)	N215256 004	Jun 04, 2021
+	!		2.4MG/0.75ML (2.4MG/0.75ML)	N215256 005	Jun 04, 2021

TABLET; ORAL

RYBELSUS

+	!	NOVO	3MG	N213051 001	Sep 20, 2019
+	!		7MG	N213051 002	Sep 20, 2019
+	!		14MG	N213051 003	Sep 20, 2019

SERTACONAZOLE NITRATE

CREAM; TOPICAL

ERTACZO

+	!	LACER PHARMA	2%	N021385 001	Dec 10, 2003
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SERTRALINE HYDROCHLORIDE

CAPSULE; ORAL

SERTRALINE HYDROCHLORIDE

+	!	ALMATICA	EQ 150MG BASE	N215133 001	Oct 04, 2021
+	!		EQ 200MG BASE	N215133 002	Oct 04, 2021

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

AA	ACI	EQ 20MG BASE/ML	A076934 001	Jun 30, 2006
AA	AUROBINDO PHARMA	EQ 20MG BASE/ML	A078861 001	Oct 31, 2008

ZOLOFT

AA	+	!	VIATRIS	EQ 20MG BASE/ML	N020990 001	Dec 07, 1999
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TABLET; ORAL

SERTRALINE HYDROCHLORIDE

AB	ACCORD HLTHCARE	EQ 25MG BASE	A202825 001	Nov 07, 2014
AB		EQ 50MG BASE	A202825 002	Nov 07, 2014
AB		EQ 100MG BASE	A202825 003	Nov 07, 2014
AB	ACI	EQ 25MG BASE	A076881 001	Feb 06, 2007
AB		EQ 50MG BASE	A076881 002	Feb 06, 2007
AB		EQ 100MG BASE	A076881 003	Feb 06, 2007
AB	ASCENT PHARMS INC	EQ 25MG BASE	A214790 001	May 03, 2021
AB		EQ 50MG BASE	A214790 002	May 03, 2021
AB		EQ 100MG BASE	A214790 003	May 03, 2021
AB	AUROBINDO PHARMA	EQ 25MG BASE	A077206 001	Feb 06, 2007
AB		EQ 50MG BASE	A077206 002	Feb 06, 2007
AB		EQ 100MG BASE	A077206 003	Feb 06, 2007
AB	GRANULES	EQ 25MG BASE	A078403 001	Jan 08, 2008
AB		EQ 50MG BASE	A078403 002	Jan 08, 2008
AB		EQ 100MG BASE	A078403 003	Jan 08, 2008
AB	HERITAGE PHARMA	EQ 25MG BASE	A076465 001	Aug 11, 2006
AB	AVET	EQ 50MG BASE	A076465 002	Aug 11, 2006
AB		EQ 100MG BASE	A076465 003	Aug 11, 2006
AB	INVAGEN PHARMS	EQ 25MG BASE	A077397 001	Feb 06, 2007
AB		EQ 50MG BASE	A077397 002	Feb 06, 2007
AB		EQ 100MG BASE	A077397 003	Feb 06, 2007
AB	LUPIN	EQ 25MG BASE	A077670 001	Feb 06, 2007
AB		EQ 50MG BASE	A077670 002	Feb 06, 2007
AB		EQ 100MG BASE	A077670 003	Feb 06, 2007

PRESCRIPTION DRUG PRODUCT LIST

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

<u>AB</u>	OXFORD PHARMS	<u>EQ 25MG BASE</u>	<u>A078175 001</u>	Jul 21, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078175 002</u>	Jul 21, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078175 003</u>	Jul 21, 2010
<u>AB</u>	REYOUNG	<u>EQ 25MG BASE</u>	<u>A078677 001</u>	Mar 04, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078677 002</u>	Mar 04, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078677 003</u>	Mar 04, 2009
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 25MG BASE</u>	<u>A076442 001</u>	Apr 30, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076442 002</u>	Apr 30, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076442 003</u>	Apr 30, 2007
<u>AB</u>	VIWIT PHARM	<u>EQ 25MG BASE</u>	<u>A076882 001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076882 002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076882 003</u>	Feb 06, 2007
<u>ZOLOFT</u>				
<u>AB</u>	+ VIATRIS	<u>EQ 25MG BASE</u>	<u>N019839 005</u>	Mar 06, 1996
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N019839 001</u>	Dec 30, 1991
<u>AB</u>	+!	<u>EQ 100MG BASE</u>	<u>N019839 002</u>	Dec 30, 1991

SETMELANOTIDE ACETATE

SOLUTION; SUBCUTANEOUS

IMCIVREE

+! RHYTHM

EQ 10MG BASE/ML (EQ 10MG BASE/ML)

N213793 001 Nov 25, 2020

SEVELAMER CARBONATE

FOR SUSPENSION; ORAL

REVELA

<u>AB</u>	+ GENZYME	<u>800MG/PACKET</u>	<u>N022318 001</u>	Aug 12, 2009
<u>AB</u>	+!	<u>2.4GM/PACKET</u>	<u>N022318 002</u>	Feb 18, 2009

SEVELAMER CARBONATE

<u>AB</u>	AUROBINDO PHARMA	<u>800MG/PACKET</u>	<u>A207624 001</u>	Jun 13, 2017
<u>AB</u>		<u>2.4GM/PACKET</u>	<u>A207624 002</u>	Jun 13, 2017
<u>AB</u>	BIONPHARMA	<u>800MG/PACKET</u>	<u>A209374 001</u>	May 17, 2021
<u>AB</u>	CHARTWELL RX	<u>800MG/PACKET</u>	<u>A217319 001</u>	Mar 02, 2023
<u>AB</u>		<u>2.4GM/PACKET</u>	<u>A217319 002</u>	Mar 02, 2023
<u>AB</u>	DR REDDYS	<u>800MG/PACKET</u>	<u>A210464 001</u>	Oct 25, 2018
<u>AB</u>		<u>2.4GM/PACKET</u>	<u>A210464 002</u>	Oct 25, 2018
<u>AB</u>	IMPAX	<u>800MG/PACKET</u>	<u>A211316 001</u>	Nov 20, 2020
<u>AB</u>		<u>2.4GM/PACKET</u>	<u>A211316 002</u>	Nov 20, 2020
<u>AB</u>	STRIDES PHARMA	<u>800MG/PACKET</u>	<u>A211917 001</u>	Sep 08, 2023
<u>AB</u>		<u>2.4GM/PACKET</u>	<u>A211917 002</u>	Sep 08, 2023

TABLET; ORAL

REVELA

<u>AB</u>	+! SANOFI	<u>800MG</u>	<u>N022127 001</u>	Oct 19, 2007
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SEVELAMER CARBONATE

<u>AB</u>	AMNEAL PHARMS CO	<u>800MG</u>	<u>A207288 001</u>	Nov 28, 2017
<u>AB</u>	ANXIN	<u>800MG</u>	<u>A212970 001</u>	Apr 09, 2020
<u>AB</u>	ARTHUR GRP	<u>800MG</u>	<u>A200959 001</u>	Mar 20, 2018
<u>AB</u>	AUROBINDO PHARMA	<u>800MG</u>	<u>A207179 001</u>	Jul 17, 2017
<u>AB</u>	DR REDDYS	<u>800MG</u>	<u>A206094 001</u>	Sep 29, 2017
<u>AB</u>	INVAGEN PHARMS	<u>800MG</u>	<u>A203860 001</u>	Oct 26, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>800MG</u>	<u>A206100 001</u>	Apr 19, 2023
<u>AB</u>	MICRO LABS	<u>800MG</u>	<u>A215537 001</u>	Feb 07, 2022
<u>AB</u>	RISING	<u>800MG</u>	<u>A204451 001</u>	Nov 29, 2018
<u>AB</u>	ZYDUS PHARMS	<u>800MG</u>	<u>A207759 001</u>	Aug 11, 2020

SEVELAMER HYDROCHLORIDE

TABLET; ORAL

RENAGEL

<u>AB</u>	+ GENZYME	<u>400MG</u>	<u>N021179 001</u>	Jul 12, 2000
<u>AB</u>	+!	<u>800MG</u>	<u>N021179 002</u>	Jul 12, 2000

SEVELAMER HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>400MG</u>	<u>A212599 001</u>	Jul 11, 2023
<u>AB</u>		<u>800MG</u>	<u>A212599 002</u>	Jul 11, 2023
<u>AB</u>	GLENMARK PHARMS LTD	<u>400MG</u>	<u>A204724 001</u>	Feb 08, 2019
<u>AB</u>		<u>800MG</u>	<u>A204724 002</u>	Feb 08, 2019
<u>AB</u>	LUPIN LTD	<u>400MG</u>	<u>A213145 001</u>	Jun 16, 2021
<u>AB</u>		<u>800MG</u>	<u>A213145 002</u>	Jun 16, 2021
<u>AB</u>	MACLEODS PHARMS LTD	<u>400MG</u>	<u>A206883 001</u>	May 26, 2023
<u>AB</u>		<u>800MG</u>	<u>A206883 002</u>	May 26, 2023

PRESCRIPTION DRUG PRODUCT LIST

SEVOFLURANE

LIQUID; INHALATION

SEVOFLURANE

<u>AN</u>	BAXTER HLTHCARE	<u>100%</u>	<u>A075895</u>	<u>001</u>	Jul 02, 2002
<u>AN</u>	HALOCARBON PRODS	<u>100%</u>	<u>A078650</u>	<u>001</u>	Nov 19, 2007
<u>AN</u>	SHANDONG	<u>100%</u>	<u>A214382</u>	<u>001</u>	Aug 18, 2023
<u>AN</u>	SHANGHAI HENGRUI	<u>100%</u>	<u>A203793</u>	<u>001</u>	Nov 03, 2015

SOJOURN

<u>AN</u>	PIRAMAL CRITICAL	<u>100%</u>	<u>A077867</u>	<u>001</u>	May 02, 2007
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ULTANE

<u>AN</u>	+! ABBVIE	<u>100%</u>	<u>N020478</u>	<u>001</u>	Jun 07, 1995
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SILDENAFIL CITRATE

FOR SUSPENSION; ORAL

REVATIO

<u>AB</u>	+! VIATRIS	<u>EQ 10MG BASE/ML</u>	<u>N203109</u>	<u>001</u>	Aug 30, 2012
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SILDENAFIL CITRATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 10MG BASE/ML</u>	<u>A212883</u>	<u>001</u>	Nov 27, 2019
<u>AB</u>	ALKEM LABS LTD	<u>EQ 10MG BASE/ML</u>	<u>A212440</u>	<u>001</u>	Nov 27, 2019
<u>AB</u>	AMNEAL PHARMS	<u>EQ 10MG BASE/ML</u>	<u>A211092</u>	<u>001</u>	Nov 27, 2019
<u>AB</u>	APPCO	<u>EQ 10MG BASE/ML</u>	<u>A213814</u>	<u>001</u>	Apr 29, 2021
<u>AB</u>	APTAPHARMA INC	<u>EQ 10MG BASE/ML</u>	<u>A213041</u>	<u>001</u>	Aug 06, 2020
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE/ML</u>	<u>A214773</u>	<u>001</u>	Dec 23, 2022
<u>AB</u>	GRANULES	<u>EQ 10MG BASE/ML</u>	<u>A214556</u>	<u>001</u>	Dec 01, 2023
<u>AB</u>	HETERO LABS LTD V	<u>EQ 10MG BASE/ML</u>	<u>A213014</u>	<u>001</u>	May 11, 2021
<u>AB</u>	LUPIN LTD	<u>EQ 10MG BASE/ML</u>	<u>A211638</u>	<u>001</u>	Mar 23, 2022
<u>AB</u>	MSN	<u>EQ 10MG BASE/ML</u>	<u>A214641</u>	<u>001</u>	Feb 08, 2022
<u>AB</u>	NOVITIUM PHARMA	<u>EQ 10MG BASE/ML</u>	<u>A212260</u>	<u>001</u>	May 31, 2019
<u>AB</u>	TARO	<u>EQ 10MG BASE/ML</u>	<u>A215522</u>	<u>001</u>	Nov 16, 2021
<u>AB</u>	TEVA PHARMS USA	<u>EQ 10MG BASE/ML</u>	<u>A211534</u>	<u>001</u>	May 29, 2020
<u>AB</u>	ZYDUS	<u>EQ 10MG BASE/ML</u>	<u>A215708</u>	<u>001</u>	Sep 29, 2022

SOLUTION; INTRAVENOUS

REVATIO

<u>AP</u>	+! VIATRIS	<u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u>	<u>N022473</u>	<u>001</u>	Nov 18, 2009
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SILDENAFIL CITRATE

<u>AP</u>	EUGIA PHARMA	<u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u>	<u>A203988</u>	<u>001</u>	Apr 01, 2015
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SUSPENSION; ORAL

LIQREV

	+! CMP DEV LLC	<u>EQ 10MG BASE/ML</u>	<u>N214952</u>	<u>001</u>	Apr 28, 2023
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TABLET; ORAL

REVATIO

<u>AB</u>	+! VIATRIS	<u>EQ 20MG BASE</u>	<u>N021845</u>	<u>001</u>	Jun 03, 2005
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SILDENAFIL CITRATE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A210394</u>	<u>001</u>	May 04, 2018
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A206401</u>	<u>001</u>	Oct 12, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A206401</u>	<u>002</u>	Oct 12, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A206401</u>	<u>003</u>	Oct 12, 2018
<u>AB</u>	AMNEAL PHARMS	<u>EQ 20MG BASE</u>	<u>A202025</u>	<u>001</u>	Feb 28, 2013
<u>AB</u>	AMNEAL PHARMS NY	<u>EQ 25MG BASE</u>	<u>A202023</u>	<u>001</u>	Jun 27, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202023</u>	<u>002</u>	Jun 27, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202023</u>	<u>003</u>	Jun 27, 2018
<u>AB</u>	APPCO	<u>EQ 25MG BASE</u>	<u>A207178</u>	<u>001</u>	Mar 02, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A207178</u>	<u>002</u>	Mar 02, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A207178</u>	<u>003</u>	Mar 02, 2020
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A203963</u>	<u>001</u>	Nov 18, 2015
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A203962</u>	<u>001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A203962</u>	<u>002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A203962</u>	<u>003</u>	Jun 11, 2018
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A218045</u>	<u>001</u>	Nov 28, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A218045</u>	<u>002</u>	Nov 28, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A218045</u>	<u>003</u>	Nov 28, 2023
<u>AB</u>	CHARTWELL RX	<u>EQ 20MG BASE</u>	<u>A202598</u>	<u>001</u>	Nov 06, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 20MG BASE</u>	<u>A203623</u>	<u>001</u>	Nov 26, 2014
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A202659</u>	<u>001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202659</u>	<u>002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202659</u>	<u>003</u>	Jun 11, 2018
<u>AB</u>	LUPIN LTD	<u>EQ 25MG BASE</u>	<u>A212051</u>	<u>001</u>	Mar 22, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A212051</u>	<u>002</u>	Mar 22, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A212051</u>	<u>003</u>	Mar 22, 2019
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A203814</u>	<u>001</u>	Dec 17, 2013
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A202255</u>	<u>001</u>	Apr 19, 2023

PRESCRIPTION DRUG PRODUCT LIST

SILDENAFIL CITRATE

TABLET;ORAL

SILDENAFIL CITRATE

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202255 002</u>	Apr 19, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202255 003</u>	Apr 19, 2023
<u>AB</u>	MYLAN	<u>EQ 25MG BASE</u>	<u>A201171 001</u>	Mar 25, 2019
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A201171 002</u>	Mar 25, 2019
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A201171 003</u>	Mar 25, 2019
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A201150 001</u>	Nov 09, 2012
<u>AB</u>	NOVITIUM PHARMA	<u>EQ 25MG BASE</u>	<u>A216383 001</u>	Aug 29, 2023
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A216383 002</u>	Aug 29, 2023
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A216383 003</u>	Aug 29, 2023
<u>AB</u>	REYOUNG	<u>EQ 25MG BASE</u>	<u>A208494 001</u>	Jun 12, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A208494 002</u>	Jun 12, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A208494 003</u>	Jun 12, 2020
<u>AB</u>	RUBICON	<u>EQ 20MG BASE</u>	<u>A204883 001</u>	Jun 20, 2016
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204882 001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A204882 002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204882 003</u>	Jun 11, 2018
<u>AB</u>	SUNSHINE	<u>EQ 25MG BASE</u>	<u>A213032 001</u>	Jun 11, 2020
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A213032 002</u>	Jun 11, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A213032 003</u>	Jun 11, 2020
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A077342 001</u>	Mar 09, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077342 002</u>	Mar 09, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077342 003</u>	Mar 09, 2016
<u>AB</u>	TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078380 001</u>	Jan 07, 2013
<u>AB</u>	TORRENT	<u>EQ 25MG BASE</u>	<u>A091448 001</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A091448 002</u>	Jun 11, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091448 003</u>	Jun 11, 2018
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A091479 001</u>	Nov 06, 2012
<u>AB</u>	UMEDICA	<u>EQ 25MG BASE</u>	<u>A209302 003</u>	Jun 15, 2021
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A209302 001</u>	Aug 25, 2020
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209302 002</u>	Aug 25, 2020
<u>AB</u>	WATSON LABS INC	<u>EQ 20MG BASE</u>	<u>A202503 001</u>	Nov 06, 2012

VIAGRA

<u>AB</u>	+	VIATRIS	<u>EQ 25MG BASE</u>	<u>N020895 001</u>	Mar 27, 1998
<u>AB</u>	+		<u>EQ 50MG BASE</u>	<u>N020895 002</u>	Mar 27, 1998
<u>AB</u>	+	!	<u>EQ 100MG BASE</u>	<u>N020895 003</u>	Mar 27, 1998

SILODOSIN

CAPSULE;ORAL

RAPAFLO

<u>AB</u>	+	!	ALLERGAN	<u>4MG</u>	<u>N022206 001</u>	Oct 08, 2008
<u>AB</u>	+			<u>8MG</u>	<u>N022206 002</u>	Oct 08, 2008

SILODOSIN

<u>AB</u>		AJANTA PHARMA LTD	<u>4MG</u>	<u>A211060 001</u>	Dec 03, 2018
<u>AB</u>			<u>8MG</u>	<u>A211060 002</u>	Dec 03, 2018
<u>AB</u>		AMNEAL PHARMS CO	<u>4MG</u>	<u>A209745 001</u>	Dec 03, 2018
<u>AB</u>			<u>8MG</u>	<u>A209745 002</u>	Dec 03, 2018
<u>AB</u>		AUROBINDO PHARMA LTD	<u>4MG</u>	<u>A210626 001</u>	Dec 10, 2018
<u>AB</u>			<u>8MG</u>	<u>A210626 002</u>	Dec 10, 2018
<u>AB</u>		CREEKWOOD PHARMS	<u>4MG</u>	<u>A213230 001</u>	Jan 03, 2022
<u>AB</u>			<u>8MG</u>	<u>A213230 002</u>	Jan 03, 2022
<u>AB</u>		HETERO LABS LTD V	<u>4MG</u>	<u>A204793 001</u>	Feb 14, 2020
<u>AB</u>			<u>8MG</u>	<u>A204793 002</u>	Feb 14, 2020
<u>AB</u>		LUPIN LTD	<u>4MG</u>	<u>A206541 001</u>	Dec 03, 2018
<u>AB</u>			<u>8MG</u>	<u>A206541 002</u>	Dec 03, 2018
<u>AB</u>		MACLEODS PHARMS LTD	<u>4MG</u>	<u>A211166 001</u>	Dec 03, 2018
<u>AB</u>			<u>8MG</u>	<u>A211166 002</u>	Dec 03, 2018
<u>AB</u>		MSN	<u>4MG</u>	<u>A210687 001</u>	Dec 03, 2018
<u>AB</u>			<u>8MG</u>	<u>A210687 002</u>	Dec 03, 2018
<u>AB</u>		PRINSTON INC	<u>4MG</u>	<u>A209029 001</u>	Jan 04, 2022
<u>AB</u>			<u>8MG</u>	<u>A209029 002</u>	Jan 04, 2022
<u>AB</u>		SANDOZ	<u>4MG</u>	<u>A204726 001</u>	Mar 31, 2017
<u>AB</u>			<u>8MG</u>	<u>A204726 002</u>	Mar 31, 2017

SILVER SULFADIAZINE

CREAM;TOPICAL

SILVADENE

<u>AB</u>	+	!	KING PHARMS LLC	<u>1%</u>	<u>N017381 001</u>	
<u>AB</u>			DR REDDYS LA	<u>1%</u>	<u>N018578 001</u>	Feb 25, 1982

PRESCRIPTION DRUG PRODUCT LIST

SILVER SULFADIAZINE

CREAM; TOPICAL

THERMAZENE

<u>AB</u>	THEPHARMANETWORK LLC	<u>1%</u>	<u>N018810</u>	<u>001</u>	Dec 23, 1985
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SIMVASTATIN

SUSPENSION; ORAL

FLOLIPID

	+ TCG FLUENT PHARMA	20MG/5ML	N206679	001	Apr 21, 2016
	+!	40MG/5ML	N206679	002	Apr 21, 2016

TABLET; ORAL

SIMVASTATIN

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A078155</u>	<u>005</u>	Apr 05, 2013
<u>AB</u>		<u>10MG</u>	<u>A078155</u>	<u>002</u>	Feb 26, 2008
<u>AB</u>		<u>20MG</u>	<u>A078155</u>	<u>003</u>	Feb 26, 2008
<u>AB</u>		<u>40MG</u>	<u>A078155</u>	<u>004</u>	Feb 26, 2008
<u>AB</u>		<u>80MG</u>	<u>A078155</u>	<u>001</u>	Feb 26, 2008
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A077691</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A077691</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077691</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077691</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077691</u>	<u>005</u>	Dec 20, 2006
<u>AB</u>	BIOCON PHARMA	<u>5MG</u>	<u>A078034</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A078034</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A078034</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A078034</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A078034</u>	<u>005</u>	Dec 20, 2006
<u>AB</u>	DR REDDYS LABS INC	<u>5MG</u>	<u>A077752</u>	<u>005</u>	Jan 23, 2008
<u>AB</u>		<u>10MG</u>	<u>A077752</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077752</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077752</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077752</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>	HETERO LABS LTD III	<u>5MG</u>	<u>A200895</u>	<u>001</u>	Nov 25, 2014
<u>AB</u>		<u>10MG</u>	<u>A200895</u>	<u>002</u>	Nov 25, 2014
<u>AB</u>		<u>20MG</u>	<u>A200895</u>	<u>003</u>	Nov 25, 2014
<u>AB</u>		<u>40MG</u>	<u>A200895</u>	<u>004</u>	Nov 25, 2014
<u>AB</u>		<u>80MG</u>	<u>A200895</u>	<u>005</u>	Nov 25, 2014
<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A206557</u>	<u>001</u>	Nov 13, 2017
<u>AB</u>		<u>20MG</u>	<u>A206557</u>	<u>002</u>	Nov 13, 2017
<u>AB</u>		<u>40MG</u>	<u>A206557</u>	<u>003</u>	Nov 13, 2017
<u>AB</u>		<u>80MG</u>	<u>A206557</u>	<u>004</u>	Nov 13, 2017
<u>AB</u>	LUPIN	<u>5MG</u>	<u>A078103</u>	<u>005</u>	Apr 14, 2009
<u>AB</u>		<u>10MG</u>	<u>A078103</u>	<u>001</u>	May 11, 2007
<u>AB</u>		<u>20MG</u>	<u>A078103</u>	<u>002</u>	May 11, 2007
<u>AB</u>		<u>40MG</u>	<u>A078103</u>	<u>003</u>	May 11, 2007
<u>AB</u>	!	<u>80MG</u>	<u>A078103</u>	<u>004</u>	May 11, 2007
<u>AB</u>	MICRO LABS	<u>5MG</u>	<u>A090383</u>	<u>001</u>	Sep 16, 2011
<u>AB</u>		<u>10MG</u>	<u>A090383</u>	<u>002</u>	Sep 16, 2011
<u>AB</u>		<u>20MG</u>	<u>A090383</u>	<u>003</u>	Sep 16, 2011
<u>AB</u>		<u>40MG</u>	<u>A090383</u>	<u>004</u>	Sep 16, 2011
<u>AB</u>		<u>80MG</u>	<u>A090383</u>	<u>005</u>	Sep 16, 2011
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A078735</u>	<u>001</u>	Aug 30, 2010
<u>AB</u>		<u>10MG</u>	<u>A078735</u>	<u>002</u>	Aug 30, 2010
<u>AB</u>		<u>20MG</u>	<u>A078735</u>	<u>003</u>	Aug 30, 2010
<u>AB</u>		<u>40MG</u>	<u>A078735</u>	<u>004</u>	Aug 30, 2010
<u>AB</u>		<u>80MG</u>	<u>A078735</u>	<u>005</u>	Aug 30, 2010
<u>AB</u>	ZYDUS PHARMS USA	<u>5MG</u>	<u>A077837</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>10MG</u>	<u>A077837</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>		<u>20MG</u>	<u>A077837</u>	<u>003</u>	Dec 20, 2006
<u>AB</u>		<u>40MG</u>	<u>A077837</u>	<u>004</u>	Dec 20, 2006
<u>AB</u>		<u>80MG</u>	<u>A077837</u>	<u>005</u>	Dec 20, 2006
<u>ZOCOR</u>					
<u>AB</u>	+ ORGANON	<u>5MG</u>	<u>N019766</u>	<u>001</u>	Dec 23, 1991
<u>AB</u>	+	<u>10MG</u>	<u>N019766</u>	<u>002</u>	Dec 23, 1991
<u>AB</u>	+	<u>20MG</u>	<u>N019766</u>	<u>003</u>	Dec 23, 1991
<u>AB</u>	+	<u>40MG</u>	<u>N019766</u>	<u>004</u>	Dec 23, 1991

PRESCRIPTION DRUG PRODUCT LIST

SINCALIDE

INJECTABLE; INJECTION

KINEVAC

+! BRACCO 0.005MG/VIAL N017697 001

POWDER; INTRAVENOUS

SINCALIDE

+! MAIA PHARMS INC 0.005MG/VIAL N210850 001 Nov 22, 2022

SINECATECHINS

OINTMENT; TOPICAL

VEREGEN

+! ANI PHARMS 15% N021902 001 Oct 31, 2006

SIPONIMOD

TABLET; ORAL

MAYZENT

+ NOVARTIS EQ 0.25MG BASE N209884 001 Mar 26, 2019

+ EQ 1MG BASE N209884 003 Aug 24, 2021

+! EQ 2MG BASE N209884 002 Mar 26, 2019

SIROLIMUS

GEL; TOPICAL

HYFTOR

+! NOBELPHARMA 0.2% N213478 001 Mar 22, 2022

POWDER; INTRAVENOUS

FYARRO

+! AADI 100MG/VIAL N213312 001 Nov 22, 2021

SOLUTION; ORAL

RAPAMUNE**AA** +! PF PRISM CV 1MG/ML **N021083 001** Sep 15, 1999SIROLIMUS**AA** AMNEAL 1MG/ML **A211212 001** Oct 18, 2019**AA** APOTEX 1MG/ML **A211406 001** Oct 22, 2019**AA** MSN 1MG/ML **A216728 001** Jan 19, 2023**AA** NOVITIUM PHARMA 1MG/ML **A211040 001** Jan 28, 2019

TABLET; ORAL

RAPAMUNE**AB** + PF PRISM CV 0.5MG **N021110 004** Jan 25, 2010**AB** + 1MG **N021110 001** Aug 25, 2000**AB** +! 2MG **N021110 002** Aug 22, 2002SIROLIMUS**AB** ALKEM LABS LTD 0.5MG **A214753 001** Mar 12, 2021**AB** 1MG **A214753 002** Mar 12, 2021**AB** 2MG **A214753 003** Mar 12, 2021**AB** DR REDDYS 1MG **A201578 001** Oct 27, 2014**AB** 2MG **A201578 002** Oct 27, 2014**AB** GLENMARK PHARMS LTD 0.5MG **A208691 001** Oct 16, 2020**AB** 1MG **A208691 002** Oct 16, 2020**AB** 2MG **A208691 003** Oct 16, 2020**AB** ZYDUS PHARMS 1MG **A201676 001** Feb 15, 2023**AB** 2MG **A201676 002** Feb 15, 2023**AB** 0.5MG **A201676 003** Jan 08, 2014SITAGLIPTIN

TABLET; ORAL

ZITUVIO

+ ZYDUS 25MG N211566 001 Oct 18, 2023

+ 50MG N211566 002 Oct 18, 2023

+! 100MG N211566 003 Oct 18, 2023

SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUVIA

+ MERCK SHARP DOHME EQ 25MG BASE N021995 001 Oct 16, 2006

+ EQ 50MG BASE N021995 002 Oct 16, 2006

+! EQ 100MG BASE N021995 003 Oct 16, 2006

SODIUM ACETATE

INJECTABLE; INJECTION

SODIUM ACETATE**AP** FRESENIUS KABI USA 4MEQ/ML **A206687 001** Oct 30, 2017**AP** +! HOSPIRA 2MEQ/ML **N018893 001** May 04, 1983**AP** MILLA PHARMS 2MEQ/ML **A214805 001** May 04, 2021**AP** 4MEQ/ML **A214805 002** Aug 25, 2023

PRESCRIPTION DRUG PRODUCT LIST

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; INTRAVENOUS

AMMONUL

<u>AP</u>	<u>+!</u>	<u>BAUSCH</u>	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>N020645 001</u>	Feb 17, 2005
<u>SODIUM PHENYLACETATE AND SODIUM BENZOATE</u>					
<u>AP</u>		<u>AILEX PHARMS LLC</u>	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>A207096 001</u>	Feb 24, 2016
<u>AP</u>		<u>MAIA PHARMS INC</u>	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>A208521 001</u>	May 08, 2017
<u>AP</u>	<u>+!</u>		<u>10%;10% (2GM/20ML;2GM/20ML)</u>	<u>N215025 001</u>	Jun 10, 2021
<u>AP</u>		<u>NAVINTA LLC</u>	<u>10%;10% (5GM/50ML;5GM/50ML)</u>	<u>A205880 001</u>	Aug 04, 2016
<u>AP</u>			<u>10%;10% (2GM/20ML; 2GM/20ML)</u>	<u>A217526 001</u>	Jul 14, 2023

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE

<u>AP</u>	<u>!</u>	<u>EXELA PHARMA</u>	<u>0.5MEQ/ML</u>	<u>A211091 001</u>	Jun 20, 2019
<u>AP</u>	<u>!</u>		<u>0.9MEQ/ML</u>	<u>A211091 002</u>	Jun 20, 2019
<u>AP</u>	<u>!</u>		<u>1MEQ/ML</u>	<u>A211091 003</u>	Jun 20, 2019
<u>AP</u>		<u>HOSPIRA</u>	<u>0.5MEQ/ML</u>	<u>A202679 001</u>	Mar 07, 2017
<u>AP</u>	<u>!</u>		<u>0.5MEQ/ML</u>	<u>A202981 001</u>	Mar 04, 2016
<u>AP</u>			<u>0.9MEQ/ML</u>	<u>A202494 001</u>	Mar 06, 2017
<u>AP</u>			<u>1MEQ/ML</u>	<u>A202432 001</u>	Sep 26, 2017
<u>AP</u>			<u>1MEQ/ML</u>	<u>A202494 002</u>	Mar 06, 2017
<u>AP</u>		<u>INTL MEDICATION SYS</u>	<u>1MEQ/ML</u>	<u>A203449 001</u>	Sep 19, 2017
<u>AP</u>		<u>OMNIVIVUM PHARMS</u>	<u>1MEQ/ML</u>	<u>A216364 001</u>	Jun 15, 2023
<u>AP</u>		<u>STERISCIENCE</u>	<u>1MEQ/ML</u>	<u>A217594 001</u>	Jun 28, 2023
		<u>HOSPIRA</u>	<u>1MEQ/ML</u>	<u>A202495 001</u>	Mar 06, 2017

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u>	<u>B BRAUN</u>	<u>450MG/100ML</u>	<u>N019635 001</u>	Mar 09, 1988
<u>AP</u>	<u>+!</u>	<u>BAXTER HLTHCARE</u>	<u>450MG/100ML</u>	<u>N018016 001</u>	
<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>450MG/100ML</u>	<u>A208122 001</u>	Jul 23, 2018
<u>AP</u>		<u>HOSPIRA</u>	<u>450MG/100ML</u>	<u>N019759 001</u>	Jun 08, 1988
<u>AP</u>	<u>+!</u>	<u>ICU MEDICAL INC</u>	<u>450MG/100ML</u>	<u>N018090 001</u>	

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	<u>+!</u>	<u>B BRAUN</u>	<u>900MG/100ML</u>	<u>N017464 001</u>	
<u>AP</u>	<u>+!</u>		<u>900MG/100ML</u>	<u>N019635 002</u>	Mar 09, 1988
<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>900MG/100ML</u>	<u>A207310 001</u>	Sep 19, 2017
<u>AP</u>		<u>FRESENIUS MEDCL</u>	<u>900MG/100ML</u>	<u>A078177 001</u>	Apr 12, 2007
<u>AP</u>		<u>HAEMONETICS</u>	<u>900MG/100ML</u>	<u>A076316 001</u>	Oct 27, 2004
<u>AP</u>	<u>+!</u>	<u>HOSPIRA</u>	<u>900MG/100ML</u>	<u>N019480 001</u>	Sep 17, 1985
<u>AP</u>	<u>+!</u>	<u>ICU MEDICAL INC</u>	<u>900MG/100ML</u>	<u>N016366 001</u>	
<u>AP</u>		<u>LABORATORIOS GRIFOLS</u>	<u>900MG/100ML</u>	<u>A207956 001</u>	May 25, 2017

SODIUM CHLORIDE 23.4%

<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>234MG/ML</u>	<u>A212248 001</u>	Apr 28, 2021
<u>AP</u>			<u>234MG/ML</u>	<u>A217796 001</u>	Jul 11, 2023

SODIUM CHLORIDE 3% IN PLASTIC CONTAINER

<u>AP</u>		<u>B BRAUN</u>	<u>3GM/100ML</u>	<u>N019635 003</u>	Mar 09, 1988
<u>AP</u>	<u>+!</u>	<u>BAXTER HLTHCARE</u>	<u>3GM/100ML</u>	<u>N019022 001</u>	Nov 01, 1983
<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>3GM/100ML</u>	<u>A209476 001</u>	Mar 13, 2019

SODIUM CHLORIDE 5% IN PLASTIC CONTAINER

<u>AP</u>		<u>B BRAUN</u>	<u>5GM/100ML</u>	<u>N019635 004</u>	Mar 09, 1988
<u>AP</u>	<u>+</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML</u>	<u>N019022 002</u>	Nov 01, 1983

SODIUM CHLORIDE 0.9%

<u>+!</u>	<u>MEDEFIL INC</u>	<u>90MG/10ML (9MG/ML)</u>	<u>N202832 006</u>	Jan 06, 2012
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SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>+</u>	<u>LIEBEL-FLARSHEIM</u>	<u>1.125GM/125ML (9MG/ML)</u>	<u>N021569 002</u>	Jul 27, 2006
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SOLUTION; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>		<u>FRESENIUS KABI USA</u>	<u>90MG/10ML (9MG/ML)</u>	<u>A088911 002</u>	May 17, 1985
<u>AP</u>			<u>270MG/30ML (9MG/ML)</u>	<u>A088911 001</u>	Feb 07, 1985
<u>AP</u>	<u>+!</u>	<u>HOSPIRA</u>	<u>90MG/10ML (9MG/ML)</u>	<u>N018800 001</u>	Oct 29, 1982
<u>AP</u>	<u>+!</u>		<u>180MG/20ML (9MG/ML)</u>	<u>N018800 002</u>	Oct 29, 1982
<u>AP</u>	<u>+!</u>		<u>270MG/30ML (9MG/ML)</u>	<u>N018800 003</u>	Oct 29, 1982

SODIUM CHLORIDE 0.9%

<u>AP</u>		<u>HIKMA</u>	<u>90MG/10ML (9MG/ML)</u>	<u>A201833 002</u>	Jan 07, 2015
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A201850 001</u>	Jan 20, 2012
<u>AP</u>		<u>NEXUS PHARMS</u>	<u>90MG/10ML (9MG/ML)</u>	<u>A217535 001</u>	Aug 23, 2023
<u>AP</u>			<u>180MG/20ML (9MG/ML)</u>	<u>A217535 002</u>	Aug 23, 2023
<u>AP</u>		<u>SPECTRA MDCL DEVICES</u>	<u>45MG/5ML (9MG/ML)</u>	<u>A206171 001</u>	Jul 21, 2017
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A206171 002</u>	Jul 21, 2017

PRESCRIPTION DRUG PRODUCT LIST

SODIUM CHLORIDE

SOLUTION; INJECTION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP	+!	BAXTER HLTHCARE	450MG/50ML (9MG/ML)	N016677 003	
AP	+		450MG/50ML (9MG/ML)	N020178 002	Dec 07, 1992
AP	+!		900MG/100ML	N016677 001	
AP	+!		900MG/100ML	N020178 001	Dec 07, 1992
AP	!	FRESENIUS KABI USA	90MG/10ML (9MG/ML)	A088912 001	Jan 10, 1985
AP	!		180MG/20ML (9MG/ML)	A088912 002	Jan 10, 1985
AP	!		270MG/30ML (9MG/ML)	A088912 003	Jan 10, 1985
AP	+!	HOSPIRA	90MG/10ML (9MG/ML)	N018803 001	Oct 29, 1982
AP	+!		180MG/20ML (9MG/ML)	N018803 002	Oct 29, 1982
AP	+!		450MG/50ML (9MG/ML)	N018803 003	Oct 29, 1982
AP	+		450MG/50ML (9MG/ML)	N019465 002	Jul 15, 1985
AP	+!		900MG/100ML	N019465 001	Jul 15, 1985
AP		NEPHRON	450MG/50ML (9MG/ML)	A211968 001	Apr 23, 2020
AP			900MG/100ML	A211968 002	Apr 23, 2020
AP	!	TARO	45MG/5ML (9MG/ML)	A077407 001	Aug 11, 2006
AP	!		90MG/10ML (9MG/ML)	A077407 002	Aug 11, 2006

SODIUM CHLORIDE 0.9%

+	B BRAUN	90MG/10ML (9MG/ML)	N019635 005	Aug 08, 2016
	HIKMA	18MG/2ML (9MG/ML)	A201833 001	Sep 24, 2013

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+!	BAXTER HLTHCARE	225MG/25ML (9MG/ML)	N016677 004	Oct 30, 1985
!	FRESENIUS KABI USA	18MG/2ML (9MG/ML)	A088912 004	Nov 29, 1985

SOLUTION; INTRAVENOUS

SODIUM CHLORIDE 14.6%

AP		FRESENIUS KABI USA	100MEQ/40ML (2.5MEQ/ML)	A212070 002	Apr 28, 2021
AP	+!	HOSPIRA	100MEQ/40ML (2.5MEQ/ML)	N018897 002	Jul 20, 1984

SODIUM CHLORIDE 23.4%

AP		FRESENIUS KABI USA	400MEQ/100ML (4MEQ/ML)	A212070 003	Feb 14, 2022
AP	+!	HOSPIRA	400MEQ/100ML (4MEQ/ML)	N018897 003	Jun 18, 2020

SODIUM CHLORIDE 14.6%

	FRESENIUS KABI USA	50MEQ/20ML (2.5MEQ/ML)	A212070 001	Apr 28, 2021
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SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AT	+!	B BRAUN	900MG/100ML	N016733 001	
AT		BAXTER HLTHCARE	900MG/100ML	N017427 001	
AT			900MG/100ML	N017867 001	
AT	+	ICU MEDICAL INC	900MG/100ML	N017514 001	
AT			900MG/100ML	N018314 001	

SOLUTION FOR SLUSH; IRRIGATION

SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER

+!	BAXTER HLTHCARE	900MG/100ML	N019319 002	May 17, 1985
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SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

SODIUM FLUORIDE F-18

AP		3D IMAGING DRUG	10-200mCi/ML	A203777 001	Oct 19, 2015
AP		BAMF	10-200mCi/ML	A216126 001	Dec 06, 2022
AP		BIOMEDCL RES FDN	10-200mCi/ML	A204351 001	Jan 09, 2015
AP		CARDINAL HEALTH 414	10-200mCi/ML	A203780 001	Jul 30, 2015
AP		ESSENTIAL ISOTOPES	10-200mCi/ML	A204541 001	Oct 29, 2014
AP		HOT SHOTS NM LLC	10-200mCi/ML	A204530 001	Jul 29, 2015
AP		JUBILANT DRAXIMAGE	10-200mCi/ML	A203968 001	Oct 23, 2015
AP		KREITCHMAN PET CTR	10-200mCi/ML	A203936 001	May 19, 2016
AP		MIDWEST MEDCL	10-200mCi/ML	A204440 001	Nov 17, 2015
AP		MIPS CRF	10-200mCi/ML	A204517 001	Jul 21, 2015
AP		NCM USA BRONX LLC	10-200mCi/ML	A204513 001	Nov 28, 2014
AP		NUKEMED	10-200mCi/ML	A203912 001	Apr 22, 2015
AP		PETNET	10-200mCi/ML	A203890 001	Sep 28, 2015
AP		PRECISION NUCLEAR	10-200mCi/ML	A204542 001	Feb 27, 2015
AP		SOFIE	10-200mCi/ML	A203592 001	Aug 18, 2015
AP	!		10-200mCi/ML	A203544 001	Dec 26, 2012
AP		UNIV UTAH CYCLOTRON	10-200mCi/ML	A204497 001	Apr 20, 2015
!	MCPRF	10-91.5mCi/ML	A203605 001	Jun 28, 2013	
	THE FEINSTEIN INST	20-600mCi/ML	A204328 001	Nov 19, 2014	

PRESCRIPTION DRUG PRODUCT LIST

SODIUM IODIDE I-123

CAPSULE;ORAL

SODIUM IODIDE I 123

AA	+ !	CARDINAL HEALTH 418	100uCi	N018671 001	May 27, 1982
AA	+ !		200uCi	N018671 002	May 27, 1982
AA		CURIUM	100uCi	A071909 001	Feb 28, 1989
AA			200uCi	A071910 001	Feb 28, 1989

SODIUM IODIDE I-131

CAPSULE;ORAL

SODIUM IODIDE I 131

+ JUBILANT

0.009-0.1mCi

N021305 006 May 19, 2005

SOLUTION;ORAL

HICON

AA	+ !	JUBILANT	250-1000mCi	N021305 007	Dec 05, 2011
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SODIUM IODIDE I 131

AA		INTL ISOTOPES	250-1000mCi	A209166 001	Feb 05, 2020
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SODIUM NITRITE

SOLUTION;INTRAVENOUS

SODIUM NITRITE

+! HOPE PHARMS

300MG/10ML (30MG/ML)

N203922 001 Feb 14, 2012

SODIUM NITRITE; SODIUM THIOSULFATE

SOLUTION, SOLUTION;INTRAVENOUS, INTRAVENOUS

NITHIODET

+! HOPE PHARMS

300MG/10ML (30MG/ML), N/A;N/A, 12.5GM/50ML
(250MG/ML)

N201444 001 Jan 14, 2011

SODIUM NITROPRUSSIDE

INJECTABLE;INJECTION

SODIUM NITROPRUSSIDE

AP		AMNEAL	25MG/ML	A209493 001	Nov 07, 2017
AP		BE PHARMS	25MG/ML	A214971 001	Jul 12, 2021
AP		CAPLIN	25MG/ML	A211016 001	Nov 29, 2019
AP		DR REDDYS	25MG/ML	A210114 001	Apr 10, 2019
AP		HAINAN POLY PHARM	25MG/ML	A214199 001	Aug 25, 2020
AP	!	MICRO LABS	25MG/ML	A209352 001	Dec 08, 2017
AP		MYLAN	25MG/ML	A209584 001	Aug 10, 2018
AP		MYLAN LABS LTD	25MG/ML	A210763 001	Apr 17, 2018
AP		NEXUS	25MG/ML	A207499 001	May 25, 2017
AP		SAGENT PHARMS INC	25MG/ML	A207426 001	Dec 08, 2016
AP		SOMERSET THERAPS LLC	25MG/ML	A210882 001	Aug 17, 2018
AP		XIROMED	25MG/ML	A211277 001	Oct 29, 2020

SOLUTION;INTRAVENOUS

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%

AP	+ !	EXELA PHARMA	20MG/100ML (0.2MG/ML)	N209387 003	Jul 13, 2018
AP	+ !		50MG/100ML (0.5MG/ML)	N209387 001	Mar 08, 2017

SODIUM NITROPRUSSIDE

AP		HAINAN POLY	20MG/100ML (0.2MG/ML)	A215846 002	Aug 26, 2022
AP			50MG/100ML (0.5MG/ML)	A215846 003	Aug 26, 2022
			10MG/50ML (0.2MG/ML)	A215846 001	Aug 26, 2022

SODIUM OXYBATE

FOR SUSPENSION, EXTENDED RELEASE;ORAL

LUMRYZ

+! AVADEL CNS

4.5GM/PACKET

N214755 001 May 01, 2023

+

6GM/PACKET

N214755 002 May 01, 2023

+

7.5GM/PACKET

N214755 003 May 01, 2023

+

9GM/PACKET

N214755 004 May 01, 2023

SOLUTION;ORAL

XYREM

+! JAZZ PHARMS

0.5GM/ML

N021196 001 Jul 17, 2002

SODIUM PHENYLBUTYRATE

FOR SUSPENSION;ORAL

OLPRUVA

+ ACER

2GM/PACKET

N214860 001 Dec 22, 2022

+

3GM/PACKET

N214860 002 Dec 22, 2022

+

4GM/PACKET

N214860 003 Dec 22, 2022

+

5GM/PACKET

N214860 004 Dec 22, 2022

+

6GM/PACKET

N214860 005 Dec 22, 2022

+!

6.67GM/PACKET

N214860 006 Dec 22, 2022

PRESCRIPTION DRUG PRODUCT LIST

SODIUM PHENYLBUTYRATE

PELLETS; ORAL

PHEBURANE

+! MEDUNIK

84GM/BOT

N216513 001 Jun 17, 2022

POWDER; ORAL

BUPHENYL**AB** +! HORIZON THERAP US**3GM/TEASPOONFUL****N020573 001** Apr 30, 1996SODIUM PHENYLBUTYRATE**AB** PAR PHARM**3GM/TEASPOONFUL****A203918 001** Jun 15, 2016**AB** SIGMAPHARM LABS LLC**3GM/TEASPOONFUL****A202819 001** Mar 22, 2013

TABLET; ORAL

BUPHENYL**AB** +! HORIZON THERAP US**500MG****N020572 001** May 13, 1996SODIUM PHENYLBUTYRATE**AB** GLENMARK PHARMS LTD**500MG****A216462 001** Nov 01, 2022**AB** PAR PHARM**500MG****A204395 001** Apr 15, 2016SODIUM PHENYLBUTYRATE; TAURUSODIOL

FOR SUSPENSION; ORAL

RELYVRIO

+! AMYLYX

3GM/PACKET; 1GM/PACKET

N216660 001 Sep 29, 2022

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE

! NOVEL LABS INC

0.398GM; 1.102GM

A079247 001 Dec 30, 2011

SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

SODIUM PHOSPHATES**AP** ! FRESENIUS KABI USA**142MG/ML; 276MG/ML****A209997 001** Mar 30, 2022SODIUM PHOSPHATES IN PLASTIC CONTAINER**AP** +! HOSPIRA**142MG/ML; 276MG/ML****N018892 001** May 10, 1983SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KALEXATE**AA** ! KVK TECH**454GM/BOT****A040905 001** Mar 30, 2009KIONEX**AA** ANI PHARMS**454GM/BOT****A040029 001** Feb 06, 1998SODIUM POLYSTYRENE SULFONATE**AA** BELCHER**454GM/BOT****A205727 001** Feb 23, 2016**AA** CHARTWELL RX**454GM/BOT****A206815 001** Feb 18, 2016**AA** CMP PHARMA INC**454GM/BOT****A089910 001** Jan 19, 1989**AA** EPIC PHARMA LLC**453.6GM/BOT****A202333 001** Mar 19, 2014**AA** NUVO PHARMS INC**454GM/BOT****A204071 001** Nov 28, 2014**AA** UPSHER SMITH LABS**453.6GM/BOT****A090313 001** Dec 21, 2011

KALEXATE

KVK TECH

15GM/BOT

A040905 002 Apr 03, 2015

SODIUM POLYSTYRENE SULFONATE

NUVO PHARMS INC

15GM/BOT

A204071 002 Nov 28, 2014

SUSPENSION; ORAL, RECTAL

KIONEX**AA** ANI PHARMS**15GM/60ML****A040028 001** Sep 17, 2007SPS**AA** +! CMP PHARMA INC**15GM/60ML****A087859 001** Dec 08, 1982SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SODIUM TETRADECYL SULFATE**AP** HIKMA**60MG/2ML (30MG/ML)****A209937 001** Dec 09, 2019SOTRADECOL**AP** ! MYLAN INSTITUTIONAL**60MG/2ML (30MG/ML)****A040541 002** Nov 12, 2004

! 20MG/2ML (10MG/ML)

A040541 001 Nov 12, 2004

SODIUM THIOSULFATE

SOLUTION; INTRAVENOUS

PEDMARK

+! FENNEC PHARMS INC

12.5GM/100ML (125MG/ML)

N212937 001 Sep 20, 2022

SODIUM THIOSULFATE

+! HOPE PHARMS

12.5GM/50ML (250MG/ML)

N203923 001 Feb 14, 2012

PRESCRIPTION DRUG PRODUCT LIST

SODIUM ZIRCONIUM CYCLOSILICATE

FOR SUSPENSION; ORAL

LOKELMA

+	ASTRAZENECA	5GM/PACKET	N207078	001	May 18, 2018
+	!	10GM/PACKET	N207078	002	May 18, 2018

SOFOSBUVIR

PELLETS; ORAL

SOVALDI

+	GILEAD SCIENCES INC	150MG/PACKET	N212480	001	Aug 28, 2019
+	!	200MG/PACKET	N212480	002	Aug 28, 2019

TABLET; ORAL

SOVALDI

+	GILEAD SCIENCES INC	200MG	N204671	002	Aug 28, 2019
+	!	400MG	N204671	001	Dec 06, 2013

SOFOSBUVIR; VELPATASVIR

PELLETS; ORAL

EPCLUSA

+	GILEAD SCIENCES INC	150MG; 37.5MG/PACKET	N214187	001	Jun 10, 2021
+	!	200MG; 50MG/PACKET	N214187	002	Jun 10, 2021

TABLET; ORAL

EPCLUSA

+	GILEAD SCIENCES INC	200MG; 50MG	N208341	002	Mar 19, 2020
+	!	400MG; 100MG	N208341	001	Jun 28, 2016

SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR

TABLET; ORAL

VOSEVI

+	GILEAD SCIENCES INC	400MG; 100MG; 100MG	N209195	001	Jul 18, 2017
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SOLIFENACIN SUCCINATE

SUSPENSION; ORAL

VESICARE LS

+	ASTELLAS	1MG/ML	N209529	001	May 26, 2020
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TABLET; ORAL

SOLIFENACIN SUCCINATE

AB	ALEMBIC	5MG	A205575	001	May 20, 2019
AB		10MG	A205575	002	May 20, 2019
AB	ALKEM LABS LTD	5MG	A210224	001	May 20, 2019
AB		10MG	A210224	002	May 20, 2019
AB	ANNORA PHARMA	5MG	A215761	001	Jun 15, 2022
AB		10MG	A215761	002	Jun 15, 2022
AB	AUROBINDO PHARMA	5MG	A206817	001	Dec 27, 2022
AB		10MG	A206817	002	Dec 27, 2022
AB	AUSTARPHARMA	5MG	A210281	001	May 20, 2019
AB		10MG	A210281	002	May 20, 2019
AB	CHARTWELL RX	5MG	A210582	001	May 20, 2019
AB		10MG	A210582	002	May 20, 2019
AB	GLENMARK PHARMS INC	5MG	A209239	001	May 20, 2019
AB		10MG	A209239	002	May 20, 2019
AB	MSN	5MG	A210688	001	May 20, 2019
AB		10MG	A210688	002	May 20, 2019
AB	QILU	5MG	A209333	001	May 20, 2019
AB		10MG	A209333	002	May 20, 2019
AB	RISING	5MG	A211423	001	Dec 11, 2019
AB		10MG	A211423	002	Dec 11, 2019
AB	SCIEGEN PHARMS INC	5MG	A211657	001	May 20, 2019
AB		10MG	A211657	002	May 20, 2019
AB	STRIDES PHARMA	5MG	A212214	001	Sep 26, 2019
AB		10MG	A212214	002	Sep 26, 2019
AB	TEVA PHARMS USA	5MG	A091464	001	Apr 02, 2014
AB		10MG	A091464	002	Apr 02, 2014
AB	UNICHEM	5MG	A211701	001	Aug 27, 2019
AB		10MG	A211701	002	Aug 27, 2019
AB	WATSON LABS INC	5MG	A202551	001	May 20, 2019
AB		10MG	A202551	002	May 20, 2019

VESICARE

AB	+	ASTELLAS	5MG	N021518	001	Nov 19, 2004
AB	+	!	10MG	N021518	002	Nov 19, 2004

PRESCRIPTION DRUG PRODUCT LIST

SOLRIAMFETOL HYDROCHLORIDE

TABLET;ORAL

SUNOSI

+ AXSOME MALTA

EQ 75MG BASE

N211230 001 Jun 17, 2019

+!

EQ 150MG BASE

N211230 002 Jun 17, 2019

SONIDEGIB PHOSPHATE

CAPSULE;ORAL

ODOMZO

+! SUN PHARM

EQ 200MG BASE

N205266 001 Jul 24, 2015

SORAFENIB TOSYLATE

TABLET;ORAL

NEXAVAR**AB** +! BAYER HLTHCARE**EQ 200MG BASE****N021923 001** Dec 20, 2005SORAFENIB TOSYLATE**AB** DR REDDYS**EQ 200MG BASE****A216073 001** Jun 07, 2022**AB** MYLAN**EQ 200MG BASE****A207012 001** Sep 10, 2020**AB** TEVA PHARMS USA INC**EQ 200MG BASE****A209567 001** Nov 12, 2020**AB** TORRENT**EQ 200MG BASE****A217095 001** Apr 12, 2023**AB** YABAO PHARM**EQ 200MG BASE****A209050 001** Nov 09, 2022SORBITOL

SOLUTION;IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE 3GM/100ML

N017863 001

SORBITOL 3.3% IN PLASTIC CONTAINER

B BRAUN 3.3GM/100ML

N016741 001

SOTAGLIFLOZIN

TABLET;ORAL

INPEFA

+ LEXICON PHARMS INC 200MG

N216203 001 May 26, 2023

+! 400MG

N216203 002 May 26, 2023

SOTALOL HYDROCHLORIDE

SOLUTION;INTRAVENOUS

SOTALOL HYDROCHLORIDE

+! ALTATHERA PHARMS 150MG/10ML (15MG/ML)

N022306 001 Jul 02, 2009

LLC

SOLUTION;ORAL

SOTYLIZE

+! AZURITY 5MG/ML (5MG/ML)

N205108 001 Oct 22, 2014

TABLET;ORAL

BETAPACE**AB** + LEGACY PHARMA**80MG****N019865 001** Oct 30, 1992**AB** +**120MG****N019865 005** Apr 20, 1994**AB** +!**160MG****N019865 002** Oct 30, 1992**AB** +**240MG****N019865 003** Oct 30, 1992BETAPACE AF**AB** + LEGACY PHARMA**80MG****N021151 001** Feb 22, 2000**AB** +**120MG****N021151 002** Feb 22, 2000**AB** +!**160MG****N021151 003** Feb 22, 2000SORINE**AB** AIPING PHARM INC**80MG****A075500 001** Apr 27, 2001**AB****120MG****A075500 004** Apr 27, 2001**AB****160MG****A075500 002** Apr 27, 2001**AB****240MG****A075500 003** Apr 27, 2001SOTALOL HYDROCHLORIDE**AB** APOTEX**80MG****A076140 001** Sep 26, 2002**AB****80MG****A076214 001** Aug 27, 2003**AB****120MG****A076140 002** Sep 26, 2002**AB****120MG****A076214 002** Aug 27, 2003**AB****160MG****A076140 003** Sep 26, 2002**AB****160MG****A076214 003** Aug 27, 2003**AB****240MG****A076140 004** Sep 26, 2002**AB**

AUROBINDO PHARMA

80MG**A077616 001** Feb 07, 2007

USA

AB**120MG****A077616 002** Feb 07, 2007**AB****160MG****A077616 003** Feb 07, 2007**AB**

BEXIMCO PHARMS USA

80MG**A207428 001** Oct 21, 2016**AB****80MG****A207429 001** Nov 02, 2018**AB****120MG****A207428 002** Oct 21, 2016**AB****120MG****A207429 002** Nov 02, 2018**AB****160MG****A207428 003** Oct 21, 2016

PRESCRIPTION DRUG PRODUCT LIST

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

<u>AB</u>		<u>160MG</u>	<u>A207429 003</u>	Nov 02, 2018
<u>AB</u>	EPIC PHARMA LLC	<u>80MG</u>	<u>A077070 001</u>	Nov 04, 2005
<u>AB</u>		<u>120MG</u>	<u>A077070 002</u>	Nov 04, 2005
<u>AB</u>		<u>160MG</u>	<u>A077070 003</u>	Nov 04, 2005
<u>AB</u>	OXFORD PHARMS	<u>80MG</u>	<u>A075563 001</u>	Nov 07, 2003
<u>AB</u>		<u>120MG</u>	<u>A075563 002</u>	Nov 07, 2003
<u>AB</u>		<u>160MG</u>	<u>A075563 003</u>	Nov 07, 2003
<u>AB</u>		<u>240MG</u>	<u>A075563 004</u>	Nov 07, 2003
<u>AB</u>	TEVA	<u>80MG</u>	<u>A075429 001</u>	May 01, 2000
<u>AB</u>		<u>120MG</u>	<u>A075429 002</u>	May 01, 2000
<u>AB</u>		<u>160MG</u>	<u>A075429 003</u>	May 01, 2000
<u>AB</u>		<u>240MG</u>	<u>A075429 004</u>	May 01, 2000

SOTORASIB

TABLET; ORAL

LUMAKRAS

+	AMGEN INC	120MG	N214665 001	May 28, 2021
+	!	320MG	N214665 002	Jan 20, 2023

SOYBEAN OIL

INJECTABLE; INJECTION

INTRALIPID 20%

<u>AP</u>	+	FRESENIUS	<u>20%</u>	<u>N018449 001</u>
<u>AP</u>	+	!	<u>20%</u>	<u>N020248 001</u> Aug 07, 1996
<u>AP</u>	+	B BRAUN	<u>20%</u>	<u>N019531 002</u> May 28, 1993
		INTRALIPID 30%		
	+	FRESENIUS	30%	N019942 001 Dec 30, 1993
		NUTRILIPID 10%		
	+	B BRAUN	10%	N019531 001 May 28, 1993

SPARSENTAN

TABLET; ORAL

FILSPARI

+	TRAVERE	200MG	N216403 001	Feb 17, 2023
+	!	400MG	N216403 002	Feb 17, 2023

SPINOSAD

SUSPENSION; TOPICAL

NATROBA

+	PARAPRO LLC	0.9%	N022408 001	Jan 18, 2011
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SPIRONOLACTONE

SUSPENSION; ORAL

CAROSPIR

<u>AB</u>	+	CMP DEV LLC	<u>25MG/5ML</u>	<u>N209478 001</u> Aug 04, 2017
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SPIRONOLACTONE

<u>AB</u>		AMNEAL	<u>25MG/5ML</u>	<u>A215572 001</u> Sep 05, 2023
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TABLET; ORAL

ALDACTONE

<u>AB</u>	+	PFIZER	<u>25MG</u>	<u>N012151 009</u> Dec 30, 1983
<u>AB</u>	+		<u>50MG</u>	<u>N012151 008</u> Dec 30, 1982
<u>AB</u>	+	!	<u>100MG</u>	<u>N012151 010</u> Dec 30, 1983

SPIRONOLACTONE

<u>AB</u>		ACCORD HLTHCARE	<u>25MG</u>	<u>A203512 001</u> Sep 19, 2016
<u>AB</u>			<u>50MG</u>	<u>A203512 002</u> Sep 19, 2016
<u>AB</u>			<u>100MG</u>	<u>A203512 003</u> Sep 19, 2016
<u>AB</u>		AMNEAL PHARMS	<u>25MG</u>	<u>A091426 001</u> Jul 02, 2010
<u>AB</u>			<u>50MG</u>	<u>A091426 002</u> Jul 02, 2010
<u>AB</u>			<u>100MG</u>	<u>A091426 003</u> Jul 02, 2010
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A202187 001</u> Mar 06, 2014
<u>AB</u>			<u>50MG</u>	<u>A202187 002</u> Mar 06, 2014
<u>AB</u>			<u>100MG</u>	<u>A202187 003</u> Mar 06, 2014
<u>AB</u>		JUBILANT GENERICS	<u>25MG</u>	<u>A203253 001</u> Apr 23, 2014
<u>AB</u>			<u>50MG</u>	<u>A203253 002</u> Apr 23, 2014
<u>AB</u>			<u>100MG</u>	<u>A203253 003</u> Apr 23, 2014
<u>AB</u>		MYLAN	<u>25MG</u>	<u>A040424 001</u> Aug 20, 2001
<u>AB</u>			<u>50MG</u>	<u>A040424 002</u> Aug 20, 2001
<u>AB</u>			<u>100MG</u>	<u>A040424 003</u> Aug 20, 2001
<u>AB</u>		OXFORD PHARMS	<u>25MG</u>	<u>A040750 001</u> Aug 29, 2006
<u>AB</u>			<u>50MG</u>	<u>A040750 002</u> Aug 29, 2006
<u>AB</u>			<u>100MG</u>	<u>A040750 003</u> Aug 29, 2006

PRESCRIPTION DRUG PRODUCT LIST

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A089424 001</u>	Jul 23, 1986
<u>AB</u>		<u>50MG</u>	<u>A089424 002</u>	Aug 11, 1999
<u>AB</u>		<u>100MG</u>	<u>A089424 003</u>	Aug 11, 1999
<u>AB</u>	ZYDUS PHARMS	<u>25MG</u>	<u>A205936 001</u>	Jul 18, 2018
<u>AB</u>		<u>50MG</u>	<u>A205936 002</u>	Jul 18, 2018
<u>AB</u>		<u>100MG</u>	<u>A205936 003</u>	Jul 18, 2018

STERILE WATER FOR INJECTION

LIQUID; N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u> ! HOSPIRA	<u>100%</u>	<u>N018802 001</u>	Oct 27, 1982
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STERILE WATER FOR INJECTION

<u>AP</u>	AM REGENT	<u>100% (10ML)</u>	<u>A217341 001</u>	Aug 29, 2023
<u>AP</u>		<u>100% (20ML)</u>	<u>A217341 002</u>	Aug 29, 2023
<u>AP</u>	FRESENIUS KABI USA	<u>100%</u>	<u>A209689 001</u>	Nov 24, 2017
<u>AP</u>	HIKMA	<u>100% (10ML)</u>	<u>A206369 001</u>	Sep 02, 2015
<u>AP</u>		<u>100%</u>	<u>A212735 001</u>	Jan 31, 2023
<u>AP</u>	<u>+</u> ! HOSPIRA	<u>100% (10ML)</u>	<u>N018801 002</u>	Oct 27, 1982
<u>AP</u>	<u>+</u> !	<u>100% (20ML)</u>	<u>N018801 003</u>	Oct 27, 1982
<u>AP</u>	<u>!</u> MEDEFIL INC	<u>100% (5ML)</u>	<u>A211188 004</u>	Dec 02, 2019
<u>AP</u>		<u>100% (10ML)</u>	<u>A211188 005</u>	Dec 02, 2019
<u>AP</u>	NEPHRON	<u>100% (5ML)</u>	<u>A211222 001</u>	Feb 10, 2021
<u>AP</u>	NEXUS	<u>100% (10ML)</u>	<u>A217536 001</u>	Jul 11, 2023
<u>AP</u>		<u>100% (20ML)</u>	<u>A217536 002</u>	Jul 11, 2023

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u> ! B BRAUN	<u>100%</u>	<u>N019633 001</u>	Feb 29, 1988
<u>AP</u>	<u>+</u> ! BAXTER HLTHCARE	<u>100%</u>	<u>N018632 001</u>	Jun 30, 1982
<u>AP</u>	<u>+</u> !	<u>100%</u>	<u>N018632 002</u>	Apr 19, 1988
<u>AP</u>	FRESENIUS KABI USA	<u>100%</u>	<u>A088400 001</u>	Jan 16, 1984
<u>AP</u>	<u>+</u> ! ICU MEDICAL INC	<u>100%</u>	<u>N018233 001</u>	
<u>AP</u>	<u>+</u> !	<u>100%</u>	<u>N019869 001</u>	Dec 26, 1989
<u>AP</u>	TARO	<u>100%</u>	<u>A077393 001</u>	Aug 11, 2006

STERILE WATER FOR INJECTION

<u>+</u> !	HOSPIRA	100% (50ML)	N018801 004	Oct 27, 1982
<u>+</u> !		100% (100ML)	N018801 005	Oct 27, 1982
<u>!</u>	MEDEFIL INC	100% (1ML)	A211188 001	Dec 02, 2019
<u>!</u>		100% (2.5ML)	A211188 002	Dec 02, 2019
<u>!</u>		100% (3ML)	A211188 003	Dec 02, 2019

STERILE WATER FOR IRRIGATION

LIQUID; IRRIGATION

STERILE WATER

<u>AT</u>	<u>+</u> BAXTER HLTHCARE	<u>100%</u>	<u>N017428 001</u>	
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STERILE WATER IN PLASTIC CONTAINER

<u>AT</u>	<u>+</u> B BRAUN	<u>100%</u>	<u>N016734 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N017866 001</u>	
<u>AT</u>	ICU MEDICAL INC	<u>100%</u>	<u>N017513 001</u>	
<u>AT</u>		<u>100%</u>	<u>N018313 001</u>	

STIRIPENTOL

CAPSULE; ORAL

DIACOMIT

<u>+</u>	BIOCODEX SA	250MG	N206709 001	Aug 20, 2018
<u>+</u> !		500MG	N206709 002	Aug 20, 2018

FOR SUSPENSION; ORAL

DIACOMIT

<u>+</u>	BIOCODEX SA	250MG/PACKET	N207223 001	Aug 20, 2018
<u>+</u> !		500MG/PACKET	N207223 002	Aug 20, 2018

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

<u>!</u>	XGEN PHARMS	EQ 1GM BASE/VIAL	A064210 001	Jun 30, 1998
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STREPTOZOCIN

INJECTABLE; INJECTION

ZANOSAR

<u>+</u> !	TEVA PHARMS USA	1GM/VIAL	N050577 001	May 07, 1982
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PRESCRIPTION DRUG PRODUCT LIST

STRONTIUM CHLORIDE SR-89

INJECTABLE; INJECTION

METASTRON

AP	+ !	Q BIOMED	1mCi/ML	N020134	001	Jun 18, 1993
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STRONTIUM CHLORIDE SR-89

AP		Q BIOMED	1mCi/ML	A075941	001	Jan 06, 2003
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SUCCIMER

CAPSULE; ORAL

CHEMET

+ !	RECORDATI RARE	100MG		N019998	002	Jan 30, 1991
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SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

ANECTINE

AP	+ !	SANDOZ	20MG/ML	N008453	002	
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QUELICIN

AP	+ !	HOSPIRA	20MG/ML	N008845	006	
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SUCCINYLCHOLINE CHLORIDE

AP		ACCORD HLTHCARE	20MG/ML	A213705	001	May 20, 2020
AP		AMNEAL	20MG/ML	A211432	001	Nov 16, 2018
AP		AMRING PHARMS	20MG/ML	A210231	001	Jun 04, 2018
AP		ASPIRO	20MG/ML	A213810	001	May 04, 2020
AP		BE PHARMS	20MG/ML	A216003	001	Feb 07, 2022
AP		DEVA HOLDING AS	20MG/ML	A214491	001	Dec 21, 2020
AP		DR REDDYS	20MG/ML	A210698	001	Aug 02, 2019
AP		EUGIA PHARMA	20MG/ML	A217808	001	Oct 16, 2023
AP		FRESENIUS KABI USA	20MG/ML	A211346	001	Nov 20, 2020
AP		GLAND PHARMA LTD	20MG/ML	A214246	001	Jun 16, 2020
AP		HIKMA	20MG/ML	A213229	001	Jun 12, 2020
AP		INDOCO	20MG/ML	A214308	001	May 22, 2020
AP		MEITHEAL	20MG/ML	A214514	001	Oct 19, 2021
AP		MICRO LABS	20MG/ML	A214879	001	Nov 24, 2020
AP		NEXUS	20MG/ML	A213552	001	Oct 27, 2020
AP		SAGENT PHARMS INC	20MG/ML	A215022	001	Mar 29, 2021
AP		SOMERSET THERAPS LLC	20MG/ML	A211589	001	Jan 15, 2020

AP		UMEDICA	20MG/ML	A211625	001	May 19, 2020
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AP	!	ZYDUS PHARMS	20MG/ML	A209467	001	May 04, 2018
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SOLUTION; INTRAMUSCULAR, INTRAVENOUS

SUCCINYLCHOLINE CHLORIDE

+ !	HIKMA	100MG/5ML (20MG/ML)		N215143	001	Aug 20, 2021
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SUCRALFATE

SUSPENSION; ORAL

CARAFATE

AB	+ !	ABBVIE	1GM/10ML	N019183	001	Dec 16, 1993
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SUCRALFATE

AB		AMNEAL	1GM/10ML	A209356	001	Dec 02, 2019
AB		MYLAN	1GM/10ML	A212913	001	Sep 12, 2022
AB		VISTAPHARM	1GM/10ML	A211884	001	Mar 15, 2022

TABLET; ORAL

CARAFATE

AB	+ !	ABBVIE	1GM	N018333	001	
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SUCRALFATE

AB		AMNEAL PHARMS	1GM	A215576	001	Apr 15, 2022
AB		NOSTRUM LABS INC	1GM	A074415	001	Jun 08, 1998
AB		TEVA	1GM	A070848	001	Mar 29, 1996
AB		ZYDUS LIFESCIENCES	1GM	A215705	001	May 03, 2023

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTA PRESERVATIVE FREE

AP	+ !	RISING	EQ 0.05MG BASE/ML	N019050	001	May 04, 1984
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SUFENTANIL CITRATE

AP		HIKMA	EQ 0.05MG BASE/ML	A074413	001	Dec 15, 1995
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AP		HOSPIRA	EQ 0.05MG BASE/ML	A074534	001	Dec 11, 1996
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TABLET; SUBLINGUAL

DSUVIA

+ !	VERTICAL PHARMS	EQ 0.03MG BASE		N209128	001	Nov 02, 2018
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PRESCRIPTION DRUG PRODUCT LIST

SUGAMMADEX SODIUM

SOLUTION; INTRAVENOUS

BRIDION

AP	+	MSD SUB MERCK	<u>EQ 200MG BASE/2ML (EQ 100MG BASE/ML)</u>	<u>N022225 002</u>	Dec 15, 2015
AP	+	!	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N022225 001</u>	Dec 15, 2015

SUGAMMADEX SODIUM

AP		ASPIRO	<u>EQ 200MG BASE/2ML (EQ 100MG BASE/ML)</u>	<u>A214337 001</u>	Jun 09, 2023
AP			<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>A214337 002</u>	Jun 09, 2023
AP		ZYDUS PHARMS	<u>EQ 200MG BASE/2ML (EQ 100MG BASE/ML)</u>	<u>A214290 001</u>	Oct 04, 2023
AP			<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>A214290 002</u>	Oct 04, 2023

SULCONAZOLE NITRATE

CREAM; TOPICAL

EXELDERM

	+	JOURNEY	1%	N018737 001	Feb 28, 1989
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SOLUTION; TOPICAL

EXELDERM

	+	JOURNEY	1%	N018738 001	Aug 30, 1985
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SULFACETAMIDE SODIUM

LOTION; TOPICAL

KLARON

AB	+	BAUSCH	10%	<u>N019931 001</u>	Dec 23, 1996
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SULFACETAMIDE SODIUM

AB		FOUGERA PHARMS	10%	<u>A077015 001</u>	Nov 17, 2006
AB		PADAGIS US	10%	<u>A078649 001</u>	Mar 23, 2009
AB		TARO	10%	<u>A078668 001</u>	May 20, 2009

OINTMENT; OPHTHALMIC

SULFACETAMIDE SODIUM

	!	PADAGIS US	10%	A080029 001	
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SOLUTION/DROPS; OPHTHALMIC

SULFACETAMIDE SODIUM

AT		BAUSCH AND LOMB	10%	<u>A040066 001</u>	Dec 28, 1994
AT		SANDOZ	10%	<u>A089560 001</u>	Oct 18, 1988

SULFADIAZINE

TABLET; ORAL

SULFADIAZINE

	!	EPIC PHARMA LLC	500MG	A040091 001	Jul 29, 1994
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SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AP	!	MYLAN LABS LTD	<u>80MG/ML; 16MG/ML</u>	<u>A206607 001</u>	Aug 30, 2017
AP		SOMERSET	<u>80MG/ML; 16MG/ML</u>	<u>A212231 001</u>	Jun 26, 2019
AP		TEVA PHARMS USA	<u>80MG/ML; 16MG/ML</u>	<u>A073303 001</u>	Oct 31, 1991

SUSPENSION; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB		AUROBINDO PHARMA	<u>200MG/5ML; 40MG/5ML</u>	<u>A091348 001</u>	Jun 08, 2010
AB		CHARTWELL MOLECULAR	<u>200MG/5ML; 40MG/5ML</u>	<u>A077785 001</u>	Jan 24, 2007
AB	!	HIKMA	<u>200MG/5ML; 40MG/5ML</u>	<u>A074650 001</u>	Dec 29, 1997
AB		NOVITIUM PHARMA	<u>200MG/5ML; 40MG/5ML</u>	<u>A214330 001</u>	Feb 08, 2022
AB		PRASCO	<u>200MG/5ML; 40MG/5ML</u>	<u>A077612 001</u>	Nov 13, 2006

SULFATRIM PEDIATRIC

AB		PHARM ASSOC	<u>200MG/5ML; 40MG/5ML</u>	<u>N018615 001</u>	Jan 07, 1983
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TABLET; ORAL

BACTRIM

AB	+	SUN PHARM INDUSTRIES	<u>400MG; 80MG</u>	<u>N017377 001</u>	
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BACTRIM DS

AB	+	!	<u>800MG; 160MG</u>	<u>N017377 002</u>	
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SEPTRA

AB		MONARCH PHARMS	<u>400MG; 80MG</u>	<u>N017376 001</u>	
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SEPTRA DS

AB		MONARCH PHARMS	<u>800MG; 160MG</u>	<u>N017376 002</u>	
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SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB		AMNEAL PHARMS NY	<u>400MG; 80MG</u>	<u>A076899 001</u>	Jan 27, 2005
AB			<u>800MG; 160MG</u>	<u>A076899 002</u>	Jan 27, 2005
AB		AUROBINDO PHARMA	<u>400MG; 80MG</u>	<u>A090624 001</u>	Feb 16, 2010
AB			<u>800MG; 160MG</u>	<u>A090624 002</u>	Feb 16, 2010
AB		CHARTWELL MOLECULES	<u>400MG; 80MG</u>	<u>A078060 002</u>	Jan 25, 2007
AB			<u>800MG; 160MG</u>	<u>A078060 001</u>	Jan 25, 2007
AB		GLENMARK GENERICS	<u>400MG; 80MG</u>	<u>A090828 002</u>	Dec 22, 2010
AB			<u>800MG; 160MG</u>	<u>A090828 001</u>	Dec 22, 2010

PRESCRIPTION DRUG PRODUCT LIST

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB	SUN PHARM INDUSTRIES	400MG; 80MG	A071017 002	Aug 25, 1986
AB		800MG; 160MG	A071017 001	Aug 25, 1986
AB	VISTA PHARMS	400MG; 80MG	A076817 001	Oct 07, 2005
AB		800MG; 160MG	A076817 002	Oct 07, 2005

SULFASALAZINE

TABLET; ORAL

AZULFIDINE

AB	+ ! PFIZER	500MG	N007073 001	
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SULFASALAZINE

AB	CHARTWELL	500MG	A080197 001	
AB	NUVO PHARMS INC	500MG	A040349 001	Jan 11, 2002
AB	WATSON LABS	500MG	A085828 001	

TABLET, DELAYED RELEASE; ORAL

AZULFIDINE EN-TABS

AB	+ ! PFIZER	500MG	N007073 002	Apr 06, 1983
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SULFASALAZINE

AB	NUVO PHARMS INC	500MG	A075339 001	Jan 11, 2002
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SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES

FOR SUSPENSION; INTRAVENOUS

LUMASON

+ !	BRACCO	60.7MG/25MG	N203684 001	Oct 15, 2014
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SULINDAC

TABLET; ORAL

SULINDAC

AB	EPIC PHARMA	150MG	A072710 001	Mar 25, 1991
AB		200MG	A072711 001	Mar 25, 1991
AB	SUN PHARM INDUSTRIES	150MG	A072050 001	Apr 17, 1991
AB		200MG	A072051 001	Apr 17, 1991
AB	WATSON LABS	150MG	A071891 001	Apr 03, 1990
AB	!	200MG	A071795 001	Apr 03, 1990

SUMATRIPTAN

SPRAY; NASAL

IMITREX

AB	+ ! GLAXOSMITHKLINE	5MG/SPRAY	N020626 001	Aug 26, 1997
AB	+ !	20MG/SPRAY	N020626 003	Aug 26, 1997

SUMATRIPTAN

AB	CIPLA	20MG/SPRAY	A214209 001	Feb 22, 2021
AB	FLORIDA	20MG/SPRAY	A208967 001	Feb 17, 2021
AB	LANNETT CO INC	5MG/SPRAY	A204841 001	Feb 19, 2016
AB		20MG/SPRAY	A204841 002	Feb 19, 2016
AB	PADAGIS ISRAEL	5MG/SPRAY	A213465 002	Sep 21, 2020
AB		20MG/SPRAY	A213465 001	Sep 21, 2020

TOSYMRA

+ !	TONIX MEDS	10MG/SPRAY	N210884 001	Jan 25, 2019
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SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX STATDOSE

AB	+ ! GLAXOSMITHKLINE	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N020080 003	Dec 23, 1996
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SUMATRIPTAN SUCCINATE

AB	DR REDDYS	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090495 001	Jan 29, 2014
AB	SUN PHARM	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090358 001	Jun 21, 2011

IMITREX

AP	+ ! GLAXOSMITHKLINE	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N020080 001	Dec 28, 1992
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SUMATRIPTAN SUCCINATE

AP	EUGIA PHARMA	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A202758 001	Apr 23, 2013
AP	FRESENIUS KABI USA	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A079242 001	Mar 02, 2009
AP	HIKMA	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A200183 001	Sep 16, 2013
AP	PAR PHARM	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A077332 001	Oct 09, 2009
AP	SOMERSET	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A213998 001	Jul 13, 2021
AP	WEST-WARD PHARMS INT	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A079123 001	Feb 06, 2009
AP	WOCKHARDT	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A078593 001	Feb 06, 2009

IMITREX STATDOSE

+ !	GLAXOSMITHKLINE	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	N020080 002	Feb 01, 2006
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PRESCRIPTION DRUG PRODUCT LIST

SUMATRIPTAN SUCCINATE

POWDER;NASAL

ONZETRA XSAIL

+! CURRAX

EQ 11MG BASE

N206099 001 Jan 27, 2016

SOLUTION;SUBCUTANEOUS

ZEMBRACE SYMTOUCH

+ TONIX MEDS

EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)

N208223 001 Jan 28, 2016

TABLET;ORAL

IMITREXAB + GLAXOSMITHKLINEEQ 25MG BASEN020132 002 Jun 01, 1995AB +EQ 50MG BASEN020132 003 Jun 01, 1995AB +!EQ 100MG BASEN020132 001 Jun 01, 1995SUMATRIPTAN SUCCINATEAB AUROBINDO PHARMAEQ 25MG BASEA078327 001 Aug 10, 2009ABEQ 50MG BASEA078327 002 Aug 10, 2009ABEQ 100MG BASEA078327 003 Aug 10, 2009AB

COREPHARMA

EQ 25MG BASEA200263 001 Jun 19, 2012ABEQ 50MG BASEA200263 002 Jun 19, 2012ABEQ 100MG BASEA200263 003 Jun 19, 2012AB

DR REDDYS LABS INC

EQ 25MG BASEA076847 001 Aug 10, 2009ABEQ 50MG BASEA076847 002 Aug 10, 2009ABEQ 100MG BASEA076847 003 Aug 10, 2009AB

MYLAN

EQ 25MG BASEA077744 001 Aug 10, 2009ABEQ 50MG BASEA077744 002 Aug 10, 2009ABEQ 100MG BASEA077744 003 Aug 10, 2009AB

ORBION PHARMS

EQ 25MG BASEA078284 001 Aug 10, 2009ABEQ 50MG BASEA078284 002 Aug 10, 2009ABEQ 100MG BASEA078284 003 Aug 10, 2009AB

SUN PHARM INDS

EQ 25MG BASEA078295 001 Aug 10, 2009ABEQ 50MG BASEA078295 002 Aug 10, 2009ABEQ 100MG BASEA078295 003 Aug 10, 2009AB

WATSON LABS

EQ 25MG BASEA076933 001 Aug 10, 2009ABEQ 50MG BASEA076933 002 Aug 10, 2009ABEQ 100MG BASEA076933 003 Aug 10, 2009SUNITINIB MALATE

CAPSULE;ORAL

SUNITINIB MALATEAB DR REDDYSEQ 12.5MG BASEA215843 001 Apr 11, 2022ABEQ 25MG BASEA215843 002 Apr 11, 2022ABEQ 37.5MG BASEA215843 003 Apr 11, 2022ABEQ 50MG BASEA215843 004 Apr 11, 2022AB

MYLAN

EQ 12.5MG BASEA201275 001 Dec 06, 2021ABEQ 25MG BASEA201275 002 Dec 06, 2021ABEQ 37.5MG BASEA201275 003 Dec 06, 2021ABEQ 50MG BASEA201275 004 Dec 06, 2021AB

NATCO PHARMA

EQ 12.5MG BASEA218024 001 Oct 24, 2023ABEQ 25MG BASEA218024 002 Oct 24, 2023ABEQ 37.5MG BASEA218024 003 Oct 24, 2023ABEQ 50MG BASEA218024 004 Oct 24, 2023AB

NOVUGEN

EQ 12.5MG BASEA218012 001 Aug 21, 2023ABEQ 25MG BASEA218012 002 Aug 21, 2023ABEQ 37.5MG BASEA218012 003 Aug 21, 2023ABEQ 50MG BASEA218012 004 Aug 21, 2023AB

SUN PHARM

EQ 12.5MG BASEA213914 001 Aug 16, 2021ABEQ 25MG BASEA213914 002 Aug 16, 2021ABEQ 37.5MG BASEA213914 003 Aug 16, 2021ABEQ 50MG BASEA213914 004 Aug 16, 2021AB

TEVA PHARMS USA

EQ 12.5MG BASEA213803 001 Nov 30, 2021ABEQ 25MG BASEA213803 002 Nov 30, 2021ABEQ 37.5MG BASEA213803 003 Nov 30, 2021ABEQ 50MG BASEA213803 004 Nov 30, 2021SUTENTAB + CPPI CVEQ 12.5MG BASEN021938 001 Jan 26, 2006AB +EQ 25MG BASEN021938 002 Jan 26, 2006AB +EQ 37.5MG BASEN021938 004 Mar 31, 2009AB +!EQ 50MG BASEN021938 003 Jan 26, 2006

PRESCRIPTION DRUG PRODUCT LIST

SUVOREXANT

TABLET; ORAL

BELSOMRA

+	MERCK SHARP DOHME	5MG	N204569	001	Aug 13, 2014
+		10MG	N204569	002	Aug 13, 2014
+		15MG	N204569	003	Aug 13, 2014
+	!	20MG	N204569	004	Aug 13, 2014

TACROLIMUS

CAPSULE; ORAL

PROGRAF

<u>AB</u>	+	ASTELLAS	<u>EQ 0.5MG BASE</u>	<u>N050708</u>	<u>003</u>	Aug 24, 1998
<u>AB</u>	+		<u>EQ 1MG BASE</u>	<u>N050708</u>	<u>001</u>	Apr 08, 1994
<u>AB</u>	+	!	<u>EQ 5MG BASE</u>	<u>N050708</u>	<u>002</u>	Apr 08, 1994

TACROLIMUS

<u>AB</u>		ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A203740</u>	<u>001</u>	Nov 12, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A203740</u>	<u>002</u>	Nov 12, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A203740</u>	<u>003</u>	Nov 12, 2020
<u>AB</u>		BELCHER	<u>EQ 0.5MG BASE</u>	<u>A206651</u>	<u>001</u>	Nov 30, 2017
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206651</u>	<u>002</u>	Nov 30, 2017
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A206651</u>	<u>003</u>	Nov 30, 2017
<u>AB</u>		BIOCON PHARMA	<u>EQ 0.5MG BASE</u>	<u>A212297</u>	<u>001</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A212297</u>	<u>002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A212297</u>	<u>003</u>	Nov 10, 2020
<u>AB</u>		CONCORD BIOTECH LTD	<u>EQ 1MG BASE</u>	<u>A213112</u>	<u>002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A213112</u>	<u>003</u>	Nov 10, 2020
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A090509</u>	<u>001</u>	May 12, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090509</u>	<u>002</u>	May 12, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090509</u>	<u>003</u>	May 12, 2010
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 0.5MG BASE</u>	<u>A206662</u>	<u>001</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A206662</u>	<u>002</u>	Nov 10, 2020
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A206662</u>	<u>003</u>	Nov 10, 2020
<u>AB</u>		HANGZHOU ZHONGMEI	<u>EQ 0.5MG BASE</u>	<u>A210929</u>	<u>001</u>	Apr 17, 2023
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A210929</u>	<u>002</u>	Apr 17, 2023
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A210929</u>	<u>003</u>	Apr 17, 2023
<u>AB</u>		MYLAN	<u>EQ 0.5MG BASE</u>	<u>A090596</u>	<u>001</u>	Sep 17, 2010
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090596</u>	<u>002</u>	Sep 17, 2010
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090596</u>	<u>003</u>	Sep 17, 2010
<u>AB</u>		PANACEA	<u>EQ 0.5MG BASE</u>	<u>A090802</u>	<u>001</u>	Sep 28, 2012
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090802</u>	<u>002</u>	Sep 28, 2012
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090802</u>	<u>003</u>	Sep 28, 2012
<u>AB</u>		SANDOZ	<u>EQ 0.5MG BASE</u>	<u>A065461</u>	<u>001</u>	Aug 10, 2009
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A065461</u>	<u>002</u>	Aug 10, 2009
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A065461</u>	<u>003</u>	Aug 10, 2009
<u>AB</u>		STRIDES PHARMA	<u>EQ 0.5MG BASE</u>	<u>A090687</u>	<u>001</u>	Jul 22, 2014
<u>AB</u>			<u>EQ 1MG BASE</u>	<u>A090687</u>	<u>002</u>	Jul 22, 2014
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A090687</u>	<u>003</u>	Jul 22, 2014
BX		ACCORD HLTHCARE	EQ 0.5MG BASE	A091195	001	Aug 31, 2011
BX			EQ 1MG BASE	A091195	002	Aug 31, 2011
BX			EQ 5MG BASE	A091195	003	Aug 31, 2011

CAPSULE, EXTENDED RELEASE; ORAL

ASTAGRAF XL

+	ASTELLAS	EQ 0.5MG BASE	N204096	001	Jul 19, 2013
+		EQ 1MG BASE	N204096	002	Jul 19, 2013
+	!	EQ 5MG BASE	N204096	003	Jul 19, 2013

FOR SUSPENSION; ORAL

PROGRAF

+	ASTELLAS	EQ 0.2MG BASE/PACKET	N210115	001	May 24, 2018
+	!	EQ 1MG BASE/PACKET	N210115	002	May 24, 2018

INJECTABLE; INJECTION

PROGRAF

+	!	ASTELLAS	EQ 5MG BASE/ML	N050709	001	Apr 08, 1994
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OINTMENT; TOPICAL

PROTOPIC

<u>AB</u>	+	!	LEO PHARMA AS	<u>0.03%</u>	<u>N050777</u>	<u>001</u>	Dec 08, 2000
<u>AB</u>	+	!		<u>0.1%</u>	<u>N050777</u>	<u>002</u>	Dec 08, 2000

TACROLIMUS

<u>AB</u>		ACCORD HLTHCARE	<u>0.03%</u>	<u>A211688</u>	<u>001</u>	Jan 31, 2019
<u>AB</u>			<u>0.1%</u>	<u>A211688</u>	<u>002</u>	Jan 31, 2019
<u>AB</u>		ENCUBE	<u>0.1%</u>	<u>A212387</u>	<u>001</u>	Oct 10, 2023
<u>AB</u>		FOUGERA PHARMS INC	<u>0.03%</u>	<u>A200744</u>	<u>001</u>	Sep 09, 2014
<u>AB</u>			<u>0.1%</u>	<u>A200744</u>	<u>002</u>	Sep 09, 2014
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.03%</u>	<u>A210393</u>	<u>002</u>	Aug 16, 2023

PRESCRIPTION DRUG PRODUCT LIST

TACROLIMUS

OINTMENT; TOPICAL

TACROLIMUS

AB		0.1%	A210393 001	Apr 16, 2018
TABLET, EXTENDED RELEASE; ORAL				
ENVARUSUS XR				
+	VELOXIS PHARMS INC	EQ 0.75MG BASE	N206406 001	Jul 10, 2015
+		EQ 1MG BASE	N206406 002	Jul 10, 2015
+	!	EQ 4MG BASE	N206406 003	Jul 10, 2015

TADALAFIL

SUSPENSION; ORAL

TADLIQ

+! CMP DEV LLC

20MG/5ML

N214522 001 Jun 17, 2022

TABLET; ORAL

CIALIS

AB1	+	LILLY	2.5MG	N021368 004	Jan 07, 2008
AB1	+		5MG	N021368 001	Nov 21, 2003
AB1	+		10MG	N021368 002	Nov 21, 2003
AB1	+	!	20MG	N021368 003	Nov 21, 2003

TADALAFIL

AB1		ACCORD HLTHCARE	2.5MG	A209167 001	Mar 26, 2019
AB1			5MG	A209167 002	Mar 26, 2019
AB1			10MG	A209167 003	Oct 23, 2019
AB1			20MG	A209167 004	Mar 26, 2019
AB1		AJANTA PHARMA LTD	2.5MG	A209654 001	Mar 26, 2019
AB1			5MG	A209654 002	Mar 26, 2019
AB1			10MG	A209654 003	Mar 26, 2019
AB1			20MG	A209654 004	Mar 26, 2019
AB1		ALEMBIC	2.5MG	A204809 001	Mar 26, 2019
AB1			5MG	A204809 002	Mar 26, 2019
AB1			10MG	A204809 003	Mar 26, 2019
AB1			20MG	A204809 004	Mar 26, 2019
AB1		AMNEAL PHARMS CO	2.5MG	A209744 001	Mar 26, 2019
AB1			5MG	A209744 002	Mar 26, 2019
AB1			10MG	A209744 003	Mar 26, 2019
AB1			20MG	A209744 004	Mar 26, 2019
AB1		AUROBINDO PHARMA LTD	2.5MG	A206285 001	Mar 26, 2019
AB1			5MG	A206285 002	Mar 26, 2019
AB1			10MG	A206285 003	Mar 26, 2019
AB1			20MG	A206285 004	Mar 26, 2019
AB1		AUSTARPHARMA	2.5MG	A208824 001	Oct 27, 2020
AB1			5MG	A208824 002	Oct 27, 2020
AB1			10MG	A208824 003	Oct 27, 2020
AB1			20MG	A208824 004	Oct 27, 2020
AB1		CIPLA	2.5MG	A209539 001	Mar 26, 2019
AB1			5MG	A209539 002	Mar 26, 2019
AB1			10MG	A209539 003	Mar 26, 2019
AB1			20MG	A209539 004	Mar 26, 2019
AB1		DR REDDYS	2.5MG	A210069 001	Mar 26, 2019
AB1			5MG	A210069 002	Mar 26, 2019
AB1			10MG	A210069 003	Mar 26, 2019
AB1			20MG	A210069 004	Mar 26, 2019
AB1		HETERO LABS LTD III	2.5MG	A209908 001	Mar 26, 2019
AB1			5MG	A209908 002	Mar 26, 2019
AB1			10MG	A209908 003	Mar 26, 2019
AB1			20MG	A209908 004	Mar 26, 2019
AB1		LUPIN LTD	2.5MG	A210567 001	Mar 26, 2019
AB1			5MG	A210567 002	Mar 26, 2019
AB1			10MG	A210567 003	Mar 26, 2019
AB1			20MG	A210567 004	Mar 26, 2019
AB1		MACLEODS PHARMS LTD	2.5MG	A207244 001	Oct 07, 2019
AB1			5MG	A207244 002	Oct 07, 2019
AB1			10MG	A207244 003	Oct 07, 2019
AB1			20MG	A207244 004	Oct 07, 2019
AB1		NOVITIUM PHARMA	5MG	A215949 001	Mar 03, 2023
AB1			10MG	A215949 002	Mar 03, 2023
AB1			20MG	A215949 003	Mar 03, 2023
AB1		PRINSTON INC	2.5MG	A210609 001	Aug 11, 2022
AB1			5MG	A210609 002	Aug 11, 2022
AB1			10MG	A210609 003	Aug 11, 2022
AB1			20MG	A210609 004	Aug 11, 2022

PRESCRIPTION DRUG PRODUCT LIST

TADALAFIL

TABLET; ORAL

TADALAFIL

<u>AB1</u>	QILU PHARM HAINAN	<u>2.5MG</u>	<u>A210420</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A210420</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A210420</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A210420</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>	SUN PHARM	<u>2.5MG</u>	<u>A208934</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A208934</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A208934</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A208934</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>	SUNSHINE	<u>2.5MG</u>	<u>A211335</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A211335</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A211335</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A211335</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>	TEVA PHARMS USA	<u>2.5MG</u>	<u>A090141</u>	<u>001</u>	May 22, 2018
<u>AB1</u>		<u>5MG</u>	<u>A090141</u>	<u>002</u>	May 22, 2018
<u>AB1</u>		<u>10MG</u>	<u>A090141</u>	<u>003</u>	May 22, 2018
<u>AB1</u>		<u>20MG</u>	<u>A090141</u>	<u>004</u>	May 22, 2018
<u>AB1</u>	TORRENT	<u>2.5MG</u>	<u>A211839</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A211839</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A211839</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A211839</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>	UMEDICA	<u>2.5MG</u>	<u>A211298</u>	<u>001</u>	Oct 23, 2020
<u>AB1</u>		<u>5MG</u>	<u>A211298</u>	<u>002</u>	Oct 23, 2020
<u>AB1</u>		<u>10MG</u>	<u>A211298</u>	<u>003</u>	Oct 23, 2020
<u>AB1</u>		<u>20MG</u>	<u>A211298</u>	<u>004</u>	Oct 23, 2020
<u>AB1</u>	UNICHEM	<u>2.5MG</u>	<u>A209250</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A209250</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A209250</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A209250</u>	<u>004</u>	Mar 26, 2019
<u>AB1</u>	VKT PHARMA	<u>2.5MG</u>	<u>A215556</u>	<u>001</u>	Nov 04, 2021
<u>AB1</u>		<u>5MG</u>	<u>A215556</u>	<u>002</u>	Nov 04, 2021
<u>AB1</u>		<u>10MG</u>	<u>A215556</u>	<u>003</u>	Nov 04, 2021
<u>AB1</u>		<u>20MG</u>	<u>A215556</u>	<u>004</u>	Nov 04, 2021
<u>AB1</u>	WATSON LABS INC	<u>2.5MG</u>	<u>A205885</u>	<u>001</u>	Mar 29, 2019
<u>AB1</u>		<u>5MG</u>	<u>A205885</u>	<u>002</u>	Mar 29, 2019
<u>AB1</u>		<u>10MG</u>	<u>A205885</u>	<u>003</u>	Mar 29, 2019
<u>AB1</u>		<u>20MG</u>	<u>A205885</u>	<u>004</u>	Mar 29, 2019
<u>AB1</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A206693</u>	<u>001</u>	Mar 26, 2019
<u>AB1</u>		<u>5MG</u>	<u>A206693</u>	<u>002</u>	Mar 26, 2019
<u>AB1</u>		<u>10MG</u>	<u>A206693</u>	<u>003</u>	Mar 26, 2019
<u>AB1</u>		<u>20MG</u>	<u>A206693</u>	<u>004</u>	Mar 26, 2019

ADCIRCA

<u>AB2</u>	<u>+</u> ELI LILLY CO	<u>20MG</u>	<u>N022332</u>	<u>001</u>	May 22, 2009
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ALYO

<u>AB2</u>	TEVA PHARMS INC	<u>20MG</u>	<u>A216932</u>	<u>001</u>	Oct 12, 2022
<u>AB2</u>	TEVA PHARMS USA	<u>20MG</u>	<u>A209942</u>	<u>001</u>	Feb 05, 2019

TADALAFIL

<u>AB2</u>	AJANTA PHARMA LTD	<u>20MG</u>	<u>A210392</u>	<u>001</u>	Feb 05, 2019
<u>AB2</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A206286</u>	<u>001</u>	Feb 05, 2019
<u>AB2</u>	CHARTWELL RX	<u>20MG</u>	<u>A210572</u>	<u>001</u>	Feb 05, 2019
<u>AB2</u>	CIPLA	<u>20MG</u>	<u>A210255</u>	<u>001</u>	Feb 05, 2019
<u>AB2</u>	DR REDDYS	<u>20MG</u>	<u>A210145</u>	<u>001</u>	Feb 05, 2019
<u>AB2</u>	HETERO LABS LTD III	<u>20MG</u>	<u>A209907</u>	<u>001</u>	Feb 05, 2019
<u>AB2</u>	MACLEODS PHARMS LTD	<u>20MG</u>	<u>A207290</u>	<u>001</u>	Oct 16, 2019
<u>AB2</u>	PRINSTON INC	<u>20MG</u>	<u>A210608</u>	<u>001</u>	Aug 11, 2022
<u>AB2</u>	SUNSHINE	<u>20MG</u>	<u>A213496</u>	<u>001</u>	Nov 23, 2020
<u>AB2</u>	TORRENT	<u>20MG</u>	<u>A212062</u>	<u>001</u>	Mar 26, 2019

TAFAMIDIS

CAPSULE; ORAL

VYNDAMAX

<u>+</u>	FOLDRX PHARMS	61MG	N212161	001	May 03, 2019
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TAFAMIDIS MEGLUMINE

CAPSULE; ORAL

VYNDAQEL

<u>+</u>	FOLDRX PHARMS	20MG	N211996	001	May 03, 2019
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PRESCRIPTION DRUG PRODUCT LIST

TAFENOQUINE SUCCINATE

TABLET; ORAL

ARAKODA

+! 60 DEGREES PHARMS EQ 100MG BASE N210607 001 Aug 08, 2018

KRINTAFEL

+! GLAXOSMITHKLINE EQ 150MG BASE N210795 001 Jul 20, 2018

TAFLUPROST

SOLUTION/DROPS; OPHTHALMIC

TAFLUPROST**AT** MICRO LABS **0.0015%** **A209051 001** Aug 19, 2019**AT** SANDOZ **0.0015%** **A209040 001** Jan 28, 2022ZIOPATAN**AT** +! THEA PHARMA **0.0015%** **N202514 001** Feb 10, 2012TALAZOPARIB TOSYLATE

CAPSULE; ORAL

TALZENNA

+ PFIZER EQ 0.1MG BASE N211651 005 Jun 20, 2023

+ EQ 0.25MG BASE N211651 001 Oct 16, 2018

+ EQ 0.35MG BASE N211651 006 Jun 20, 2023

+ EQ 0.5MG BASE N211651 003 Sep 20, 2021

+ EQ 0.75MG BASE N211651 004 Sep 20, 2021

+! EQ 1MG BASE N211651 002 Oct 16, 2018

TALC

AEROSOL; INTRAPLEURAL

SCLEROSOL

+! SCIARRA LABS 4GM/SPRAY N020587 001 Dec 24, 1997

POWDER; INTRAPLEURAL

STERITALC

+ NOVATECH SA 2GM/VIAL N205555 001 May 01, 2017

+ 3GM/VIAL N205555 002 May 01, 2017

+! 4GM/VIAL N205555 003 May 01, 2017

TALC

+! SCIARRA LABS 5GM/BOT N021388 001 Dec 15, 2003

TAMOXIFEN CITRATE

SOLUTION; ORAL

SOLTAMOX

+! MAYNE PHARMA EQ 20MG BASE/10ML N021807 001 Oct 29, 2005

COMMRCL

TABLET; ORAL

TAMOXIFEN CITRATE**AB** ACTAVIS LABS FL INC **EQ 10MG BASE** **A070929 001** Feb 20, 2003**AB** **EQ 20MG BASE** **A070929 002** Feb 20, 2003**AB** APOTEX **EQ 10MG BASE** **A090878 001** Sep 23, 2011**AB** **EQ 20MG BASE** **A090878 002** Sep 23, 2011**AB** DR REDDYS LABS SA **EQ 10MG BASE** **A075797 001** Feb 20, 2003**AB** ! **EQ 20MG BASE** **A075797 002** Feb 20, 2003**AB** EUGIA PHARMA **EQ 10MG BASE** **A213358 001** Aug 14, 2020**AB** **EQ 20MG BASE** **A213358 002** Aug 14, 2020**AB** MYLAN **EQ 10MG BASE** **A074732 002** Feb 20, 2003**AB** **EQ 20MG BASE** **A074732 001** Feb 20, 2003**AB** ZYDUS PHARMS **EQ 10MG BASE** **A206694 001** Oct 27, 2017**AB** **EQ 20MG BASE** **A206694 002** Oct 27, 2017TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

FLOMAX**AB** +! SANOFI **0.4MG** **N020579 001** Apr 15, 1997TAMSULOSIN HYDROCHLORIDE**AB** ALKEM LABS LTD **0.4MG** **A207405 001** Aug 11, 2017**AB** AUROBINDO PHARMA **0.4MG** **A202433 001** Apr 30, 2013

LTD

AB CHARTWELL RX **0.4MG** **A211885 001** Oct 17, 2019**AB** IMPAX LABS **0.4MG** **A090377 001** Mar 02, 2010**AB** MACLEODS PHARMS LTD **0.4MG** **A204645 001** Jan 20, 2017**AB** SANDOZ **0.4MG** **A078015 001** Apr 27, 2010**AB** SUN PHARM INDS LTD **0.4MG** **A090931 001** Jul 15, 2010**AB** SYNTHON PHARMS **0.4MG** **A078801 001** Apr 27, 2010**AB** TEVA PHARMS **0.4MG** **A077630 001** Apr 27, 2010**AB** WOCHKHARDT **0.4MG** **A078938 001** Apr 27, 2010**AB** ZYDUS PHARMS USA **0.4MG** **A078225 001** Apr 27, 2010

INC

PRESCRIPTION DRUG PRODUCT LIST

TAPENTADOL HYDROCHLORIDE

TABLET; ORAL

NUCYNTA

+	COLLEGIUM PHARM INC	EQ 50MG BASE	N022304 001	Nov 20, 2008
+		EQ 75MG BASE	N022304 002	Nov 20, 2008
+		EQ 100MG BASE	N022304 003	Nov 20, 2008

TABLET, EXTENDED RELEASE; ORAL

NUCYNTA ER

+	COLLEGIUM PHARM INC	EQ 50MG BASE	N200533 001	Aug 25, 2011
+		EQ 100MG BASE	N200533 002	Aug 25, 2011
+		EQ 150MG BASE	N200533 003	Aug 25, 2011
+		EQ 200MG BASE	N200533 004	Aug 25, 2011
+		EQ 250MG BASE	N200533 005	Aug 25, 2011

TAPINAROF

CREAM; TOPICAL

VTAMA

+	DERMAVANT SCI	1%	N215272 001	May 23, 2022
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TASIMELTEON

CAPSULE; ORAL

HETLIOZ

AB	+	VANDA PHARMS INC	20MG	N205677 001	Jan 31, 2014
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TASIMELTEON

AB		APOTEX	20MG	A211607 001	Dec 20, 2022
AB		MSN	20MG	A211654 001	Jan 12, 2023
AB		TEVA PHARMS USA INC	20MG	A211601 001	Dec 12, 2022

SUSPENSION; ORAL

HETLIOZ LQ

+	VANDA PHARMS INC	4MG/ML	N214517 001	Dec 01, 2020
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TAVABOROLE

SOLUTION; TOPICAL

KERYDIN

AB	+	ANACOR PHARMS INC	5%	N204427 001	Jul 07, 2014
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TAVABOROLE

AB		ALEMBIC	5%	A212188 001	Oct 21, 2020
AB		AMNEAL	5%	A212256 001	Nov 25, 2020
AB		CHARTWELL RX	5%	A211963 001	Feb 03, 2021
AB		CIPLA	5%	A212224 001	Feb 09, 2021
AB		ENCUBE	5%	A211297 001	Oct 13, 2020
AB		LUPIN LTD	5%	A212168 001	Feb 08, 2021
AB		PADAGIS US	5%	A211848 001	Oct 13, 2020
AB		TARO	5%	A212215 001	May 07, 2021
AB		ZYDUS LIFESCIENCES	5%	A212294 001	Apr 10, 2023

TAZAROTENE

AEROSOL, FOAM; TOPICAL

FABIOR

+	MAYNE PHARMA	0.1%	N202428 001	May 11, 2012
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CREAM; TOPICAL

AVAGE

AB	+	ALLERGAN	0.1%	N021184 003	Sep 30, 2002
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TAZAROTENE

AB		COSETTE	0.1%	A208662 001	Dec 22, 2017
AB		TARO	0.1%	A208258 001	Apr 03, 2017

TAZORAC

AB	+	ALLERGAN	0.1%	N021184 002	Sep 29, 2000
	+		0.05%	N021184 001	Sep 29, 2000

GEL; TOPICAL

TAZAROTENE

AB		COSETTE	0.05%	A215433 001	Sep 13, 2022
AB			0.1%	A214136 001	Sep 13, 2022
AB		PADAGIS ISRAEL	0.05%	A213079 001	Apr 25, 2023
AB			0.1%	A213079 002	Apr 25, 2023
AB		SOLARIS PHARMA CORP	0.05%	A213644 001	Mar 20, 2023
AB			0.1%	A213644 002	Mar 20, 2023

TAZORAC

AB	+	ALLERGAN	0.05%	N020600 001	Jun 13, 1997
AB	+		0.1%	N020600 002	Jun 13, 1997

LOTION; TOPICAL

ARAZLO

+	BAUSCH	0.045%	N211882 001	Dec 18, 2019
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PRESCRIPTION DRUG PRODUCT LIST

TAZEMETOSTAT HYDROBROMIDE

TABLET; ORAL

TAZVERIK

+! EPIZYME INC EQ 200MG BASE

N211723 001 Jan 23, 2020

TECHNETIUM TC-99M BICISATE KIT

INJECTABLE; INJECTION

NEUROLITE

+! LANTHEUS MEDCL N/A

N020256 001 Nov 23, 1994

TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE; INJECTION

CERETEC

+! GE HEALTHCARE N/A

N019829 001 Dec 30, 1988

POWDER; INTRAVENOUS

DRAX EXAMETAZIME

JUBILANT N/A

N208870 001 Aug 17, 2017

TECHNETIUM TC-99M LABELED CARBON

AEROSOL; INHALATION

TECHNEGAS KIT

+! CYCLOMEDICA N/A

N022335 001 Sep 29, 2023

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETEC**AP +! BRACCO N/A****N018963 001** Jan 21, 1987**TECHNETIUM TC-99M MEBROFENIN****AP SUN PHARM INDS INC N/A****A078242 001** Jan 29, 2008TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-25

+! JUBILANT N/A

N018035 002 Feb 27, 2004

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

CIS-MDP**AP SUN PHARM INDS INC N/A****N018124 001****TECHNETIUM TC-99M MEDRONATE KIT****AP CARDINAL HEALTH 414 N/A****N018107 001**TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE; INJECTION

TECHNESCAN MAG3**AP +! CURIUM N/A****N019882 001** Jun 15, 1990**TECHNETIUM TC99M MERTIATIDE KIT****AP MEDI-RADIOPHARMA N/A****A206489 001** Feb 06, 2020**AP SUN PHARM INDS INC N/A****A208994 001** Jul 12, 2019

POWDER; INTRAVENOUS

TECHNETIUM TC 99M MERTIATIDE KIT

+! JUBILANT DRAXIMAGE N/A

N216820 001 Jan 30, 2023

TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE; INJECTION

TECHNESCAN

+! CURIUM N/A

N018321 001

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

DRAXIMAGE DTPA

+! JUBILANT N/A

N018511 001 Dec 29, 1989

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYRO**AP SUN PHARM INDS INC N/A****N019039 001** Jun 30, 1987**TECHNESCAN PYP KIT****AP CURIUM N/A****N017538 001**TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

ULTRATAG

+! CURIUM N/A

N019981 001 Jun 10, 1991

PRESCRIPTION DRUG PRODUCT LISTTECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

CARDIOLITE

AP	+	LANTHEUS MEDCL	N/A	N019785 001	Dec 21, 1990
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TECHNETIUM TC 99M SESTAMIBI

AP		CARDINAL HEALTH 414	N/A	A078809 001	Apr 28, 2009
AP		CURIUM	N/A	A078098 001	Sep 22, 2008
AP		JUBILANT DRAXIMAGE	N/A	A078806 001	Apr 29, 2009
AP		SUN PHARM INDS INC	10-30mCi	A079157 001	Jul 10, 2009

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INTRAVENOUS

TECHNELITE

	+	LANTHEUS MEDCL	1-20 CI/GENERATOR	N017771 002	Feb 12, 2014
	+	ULTRA-TECHNEKOW FM			
	+	CURIUM	1-19 CI/GENERATOR	N017243 003	Feb 18, 2014

TECHNETIUM TC-99M SUCCIMER

POWDER; INTRAVENOUS

NEPHROSCAN

	+	THERAGNOSTICS	N/A	N214993 001	Feb 18, 2022
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TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

AN-SULFUR COLLOID

AP	+	SUN PHARM INDS INC	N/A	N017858 001	
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TECHNETIUM TC-99M SULFUR COLLOID KIT

AP		JUBILANT	N/A	A213516 001	Nov 09, 2023
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TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVUE

	+	GE HEALTHCARE	N/A	N020372 001	Feb 09, 1996
		MYOVUE 30ML			
	+	GE HEALTHCARE	N/A	N020372 002	Jul 07, 2005

TECHNETIUM TC-99M TILMANOCEPT

INJECTABLE; INJECTION

LYMPHOSEEK KIT

	+	CARDINAL HEALTH 414	N/A	N202207 001	Mar 13, 2013
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TECOVIRIMAT

CAPSULE; ORAL

TPOXX

	+	SIGA TECHNOLOGIES	200MG	N208627 001	Jul 13, 2018
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SOLUTION; INTRAVENOUS

TPOXX

	+	SIGA TECHNOLOGIES	200MG/20ML (10MG/ML)	N214518 001	May 18, 2022
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TEDIZOLID PHOSPHATE

POWDER; INTRAVENOUS

SIVEXTRO

	+	CUBIST PHARMS LLC	200MG/VIAL	N205436 001	Jun 20, 2014
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TABLET; ORAL

SIVEXTRO

	+	CUBIST PHARMS LLC	200MG	N205435 001	Jun 20, 2014
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TEDUGLUTIDE RECOMBINANT

POWDER; SUBCUTANEOUS

GATTEX KIT

	+	TAKEDA PHARMS USA	5MG/VIAL	N203441 001	Dec 21, 2012
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TELAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

VIBATIV

	+	CUMBERLAND	EQ 750MG BASE/VIAL	N022110 002	Sep 11, 2009
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TELMISARTAN

TABLET; ORAL

MICARDIS

AB	+	BOEHRINGER INGELHEIM	20MG	N020850 003	Apr 04, 2000
AB	+		40MG	N020850 001	Nov 10, 1998
AB	+		80MG	N020850 002	Nov 10, 1998

TELMISARTAN

AB		ALEMBIC	20MG	A202130 001	Jul 07, 2014
AB			40MG	A202130 002	Jul 07, 2014
AB			80MG	A202130 003	Jul 07, 2014

PRESCRIPTION DRUG PRODUCT LIST

TELMISARTAN

TABLET; ORAL

TELMISARTAN

<u>AB</u>	AMNEAL PHARMS	<u>20MG</u>	<u>A204415 001</u>	Sep 08, 2015
<u>AB</u>		<u>40MG</u>	<u>A204415 002</u>	Sep 08, 2015
<u>AB</u>		<u>80MG</u>	<u>A204415 003</u>	Sep 08, 2015
<u>AB</u>	AUROBINDO PHARMA	<u>20MG</u>	<u>A206511 001</u>	Sep 03, 2015
<u>AB</u>		<u>40MG</u>	<u>A206511 002</u>	Sep 03, 2015
<u>AB</u>		<u>80MG</u>	<u>A206511 003</u>	Sep 03, 2015
<u>AB</u>	CADILA PHARMS LTD	<u>20MG</u>	<u>A208605 001</u>	Jul 25, 2017
<u>AB</u>		<u>40MG</u>	<u>A208605 002</u>	Jul 25, 2017
<u>AB</u>		<u>80MG</u>	<u>A208605 003</u>	Jul 25, 2017
<u>AB</u>	CHARTWELL RX	<u>20MG</u>	<u>A078710 001</u>	Jan 08, 2014
<u>AB</u>		<u>40MG</u>	<u>A078710 002</u>	Jan 08, 2014
<u>AB</u>		<u>80MG</u>	<u>A078710 003</u>	Jan 08, 2014
<u>AB</u>	GLENMARK PHARMS LTD	<u>20MG</u>	<u>A090032 001</u>	Jul 07, 2014
<u>AB</u>		<u>40MG</u>	<u>A090032 002</u>	Jul 07, 2014
<u>AB</u>		<u>80MG</u>	<u>A090032 003</u>	Jul 07, 2014
<u>AB</u>	HETERO LABS LTD V	<u>20MG</u>	<u>A205901 001</u>	Apr 22, 2016
<u>AB</u>		<u>40MG</u>	<u>A205901 002</u>	Apr 22, 2016
<u>AB</u>		<u>80MG</u>	<u>A205901 003</u>	Apr 22, 2016
<u>AB</u>	INVENTIA	<u>20MG</u>	<u>A205150 001</u>	Oct 30, 2015
<u>AB</u>		<u>40MG</u>	<u>A205150 002</u>	Oct 30, 2015
<u>AB</u>		<u>80MG</u>	<u>A205150 003</u>	Oct 30, 2015
<u>AB</u>	MICRO LABS	<u>20MG</u>	<u>A207016 001</u>	Oct 03, 2017
<u>AB</u>		<u>40MG</u>	<u>A207016 002</u>	Oct 03, 2017
<u>AB</u>		<u>80MG</u>	<u>A207016 003</u>	Oct 03, 2017
<u>AB</u>	MYLAN	<u>20MG</u>	<u>A202397 001</u>	Jul 07, 2014
<u>AB</u>		<u>40MG</u>	<u>A202397 002</u>	Jul 07, 2014
<u>AB</u>		<u>80MG</u>	<u>A202397 003</u>	Jul 07, 2014
<u>AB</u>	PRINSTON INC	<u>20MG</u>	<u>A207882 001</u>	May 03, 2017
<u>AB</u>		<u>40MG</u>	<u>A207882 002</u>	May 03, 2017
<u>AB</u>		<u>80MG</u>	<u>A207882 003</u>	May 03, 2017
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>A203867 001</u>	Nov 03, 2014
<u>AB</u>		<u>40MG</u>	<u>A203867 002</u>	Nov 03, 2014
<u>AB</u>		<u>80MG</u>	<u>A203867 003</u>	Nov 03, 2014
<u>AB</u>	ZYDUS PHARMS	<u>20MG</u>	<u>A203325 001</u>	Aug 26, 2014
<u>AB</u>		<u>40MG</u>	<u>A203325 002</u>	Aug 26, 2014
<u>AB</u>		<u>80MG</u>	<u>A203325 003</u>	Aug 26, 2014

TELOTRISTAT ETIPRATE

TABLET; ORAL

XERMELO

+! TERSERA

EQ 250MG BASE

N208794 001 Feb 28, 2017

TEMAZEPAM

CAPSULE; ORAL

RESTORIL

<u>AB</u>	+	SPECGX LLC	<u>7.5MG</u>	<u>N018163 003</u>	Oct 25, 1991
<u>AB</u>	+		<u>15MG</u>	<u>N018163 001</u>	
<u>AB</u>	+		<u>22.5MG</u>	<u>N018163 004</u>	Nov 02, 2004
<u>AB</u>	+!		<u>30MG</u>	<u>N018163 002</u>	

TEMAZEPAM

<u>AB</u>		ACTAVIS ELIZABETH	<u>15MG</u>	<u>A071620 002</u>	Aug 07, 1987
<u>AB</u>			<u>30MG</u>	<u>A071620 001</u>	Aug 07, 1987
<u>AB</u>		ALEMBIC	<u>7.5MG</u>	<u>A211542 001</u>	Nov 23, 2018
<u>AB</u>			<u>15MG</u>	<u>A211542 002</u>	Nov 23, 2018
<u>AB</u>			<u>22.5MG</u>	<u>A211542 003</u>	Nov 23, 2018
<u>AB</u>			<u>30MG</u>	<u>A211542 004</u>	Nov 23, 2018
<u>AB</u>		ALKEM LABS LTD	<u>7.5MG</u>	<u>A217875 001</u>	Aug 24, 2023
<u>AB</u>			<u>15MG</u>	<u>A217875 002</u>	Aug 24, 2023
<u>AB</u>			<u>22.5MG</u>	<u>A217875 003</u>	Aug 24, 2023
<u>AB</u>			<u>30MG</u>	<u>A217875 004</u>	Aug 24, 2023
<u>AB</u>		AMNEAL PHARMS	<u>7.5MG</u>	<u>A203482 001</u>	May 23, 2016
<u>AB</u>			<u>15MG</u>	<u>A203482 002</u>	May 23, 2016
<u>AB</u>			<u>22.5MG</u>	<u>A203482 003</u>	May 23, 2016
<u>AB</u>			<u>30MG</u>	<u>A203482 004</u>	May 23, 2016
<u>AB</u>		NOVEL LABS INC	<u>7.5MG</u>	<u>A071457 002</u>	Jun 22, 2012
<u>AB</u>			<u>15MG</u>	<u>A071456 001</u>	Apr 21, 1987
<u>AB</u>			<u>22.5MG</u>	<u>A071457 003</u>	Jun 22, 2012
<u>AB</u>			<u>30MG</u>	<u>A071457 001</u>	Apr 21, 1987
<u>AB</u>		PRINSTON INC	<u>7.5MG</u>	<u>A201781 001</u>	Jun 04, 2015
<u>AB</u>			<u>15MG</u>	<u>A201781 002</u>	Jun 04, 2015

PRESCRIPTION DRUG PRODUCT LIST

TEMAZEPAM

CAPSULE; ORAL

TEMAZEPAM

<u>AB</u>		<u>22.5MG</u>	<u>A201781 003</u>	Jun 04, 2015
<u>AB</u>		<u>30MG</u>	<u>A201781 004</u>	Jun 04, 2015
<u>AB</u>	SANDOZ	<u>15MG</u>	<u>A071427 001</u>	Jan 12, 1988
<u>AB</u>		<u>30MG</u>	<u>A071428 001</u>	Jan 12, 1988
<u>AB</u>	SUN PHARM INDUSTRIES	<u>7.5MG</u>	<u>A078581 001</u>	Sep 08, 2009

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A201528 001</u>	Feb 27, 2017
<u>AB</u>		<u>20MG</u>	<u>A201528 002</u>	Feb 27, 2017
<u>AB</u>		<u>100MG</u>	<u>A201528 003</u>	Feb 27, 2017
<u>AB</u>		<u>140MG</u>	<u>A201528 004</u>	Feb 27, 2017
<u>AB</u>		<u>180MG</u>	<u>A201528 005</u>	Feb 27, 2017
<u>AB</u>	!	<u>250MG</u>	<u>A201528 006</u>	Feb 27, 2017
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A203691 001</u>	May 08, 2015
<u>AB</u>		<u>20MG</u>	<u>A203691 002</u>	May 08, 2015
<u>AB</u>		<u>100MG</u>	<u>A203691 003</u>	May 08, 2015
<u>AB</u>		<u>140MG</u>	<u>A203691 004</u>	May 08, 2015
<u>AB</u>		<u>180MG</u>	<u>A203691 005</u>	May 08, 2015
<u>AB</u>		<u>250MG</u>	<u>A203691 006</u>	May 08, 2015
<u>AB</u>	CHARTWELL	<u>5MG</u>	<u>A206413 001</u>	Apr 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A206413 002</u>	Apr 12, 2016
<u>AB</u>		<u>100MG</u>	<u>A206413 003</u>	Apr 12, 2016
<u>AB</u>		<u>140MG</u>	<u>A206413 004</u>	Apr 12, 2016
<u>AB</u>		<u>180MG</u>	<u>A206413 005</u>	Apr 12, 2016
<u>AB</u>		<u>250MG</u>	<u>A206413 006</u>	Apr 12, 2016
<u>AB</u>	CHEMI SPA	<u>5MG</u>	<u>A204639 001</u>	Nov 23, 2016
<u>AB</u>		<u>20MG</u>	<u>A204639 002</u>	Nov 23, 2016
<u>AB</u>		<u>100MG</u>	<u>A204639 003</u>	Nov 23, 2016
<u>AB</u>		<u>140MG</u>	<u>A204639 004</u>	Nov 23, 2016
<u>AB</u>		<u>180MG</u>	<u>A204639 005</u>	Nov 23, 2016
<u>AB</u>		<u>250MG</u>	<u>A204639 006</u>	Nov 23, 2016
<u>AB</u>	DEVA HOLDING AS	<u>5MG</u>	<u>A207658 001</u>	Apr 26, 2017
<u>AB</u>		<u>20MG</u>	<u>A207658 002</u>	Apr 26, 2017
<u>AB</u>		<u>100MG</u>	<u>A207658 003</u>	Apr 26, 2017
<u>AB</u>		<u>140MG</u>	<u>A207658 004</u>	Apr 26, 2017
<u>AB</u>		<u>180MG</u>	<u>A207658 005</u>	Apr 26, 2017
<u>AB</u>		<u>250MG</u>	<u>A207658 006</u>	Apr 26, 2017
<u>AB</u>	NIVAGEN PHARMS INC	<u>5MG</u>	<u>A213328 001</u>	Nov 23, 2021
<u>AB</u>		<u>20MG</u>	<u>A213328 002</u>	Nov 23, 2021
<u>AB</u>		<u>100MG</u>	<u>A213328 003</u>	Nov 23, 2021
<u>AB</u>		<u>140MG</u>	<u>A213328 004</u>	Nov 23, 2021
<u>AB</u>		<u>180MG</u>	<u>A213328 005</u>	Nov 23, 2021
<u>AB</u>		<u>250MG</u>	<u>A213328 006</u>	Nov 23, 2021
<u>AB</u>	RISING	<u>5MG</u>	<u>A206309 001</u>	Apr 27, 2016
<u>AB</u>		<u>20MG</u>	<u>A206309 002</u>	Apr 27, 2016
<u>AB</u>		<u>100MG</u>	<u>A206309 003</u>	Apr 27, 2016
<u>AB</u>		<u>140MG</u>	<u>A206309 004</u>	Apr 27, 2016
<u>AB</u>		<u>180MG</u>	<u>A206309 005</u>	Apr 27, 2016
<u>AB</u>		<u>250MG</u>	<u>A206309 006</u>	Apr 27, 2016
<u>AB</u>	SUN PHARM	<u>5MG</u>	<u>A201742 001</u>	Feb 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A201742 002</u>	Feb 12, 2014
<u>AB</u>		<u>100MG</u>	<u>A201742 003</u>	Feb 12, 2014
<u>AB</u>		<u>140MG</u>	<u>A201742 004</u>	Feb 12, 2014
<u>AB</u>		<u>180MG</u>	<u>A201742 005</u>	Feb 12, 2014
<u>AB</u>		<u>250MG</u>	<u>A201742 006</u>	Feb 12, 2014
<u>AB</u>	ZYDUS PHARMS	<u>5MG</u>	<u>A206750 001</u>	Jul 31, 2017
<u>AB</u>		<u>20MG</u>	<u>A206750 002</u>	Jul 31, 2017
<u>AB</u>		<u>100MG</u>	<u>A206750 003</u>	Jul 31, 2017
<u>AB</u>		<u>140MG</u>	<u>A206750 004</u>	Jul 31, 2017
<u>AB</u>		<u>180MG</u>	<u>A206750 005</u>	Jul 31, 2017
<u>AB</u>		<u>250MG</u>	<u>A206750 006</u>	Jul 31, 2017

POWDER; INTRAVENOUS

TEMODAR

+	!	MERCK SHARP DOHME	100MG/VIAL	N022277 001	Feb 27, 2009
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PRESCRIPTION DRUG PRODUCT LIST

TEMSIROLIMUS

SOLUTION;INTRAVENOUS

TEMSIROLIMUS

AP	ACCORD HLTHCARE	25MG/ML (25MG/ML)	A203153 001	Jul 30, 2018
AP	GLAND PHARMA LTD	25MG/ML (25MG/ML)	A207383 001	Aug 16, 2019

TORISEL

AP	+! PF PRISM CV	25MG/ML (25MG/ML)	N022088 001	May 30, 2007
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TENAPANOR HYDROCHLORIDE

TABLET;ORAL

IBSRELA

+!	ARDELYX INC	EQ 50MG BASE	N211801 001	Sep 12, 2019
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XPHOZAH

+	ARDELYX INC	EQ 10MG BASE	N213931 001	Oct 17, 2023
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+		EQ 20MG BASE	N213931 002	Oct 17, 2023
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+!		EQ 30MG BASE	N213931 003	Oct 17, 2023
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TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

TENOFOVIR ALAFENAMIDE

AB	LUPIN LTD	EQ 25MG BASE	A214226 001	Mar 30, 2023
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VEMLIDY

AB	+! GILEAD SCIENCES INC	EQ 25MG BASE	N208464 001	Nov 10, 2016
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TENOFOVIR DISOPROXIL FUMARATE

POWDER;ORAL

VIREAD

+!	GILEAD SCIENCES INC	40MG/SCOOPFUL	N022577 001	Jan 18, 2012
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TABLET;ORAL

TENOFOVIR DISOPROXIL FUMARATE

AB	AUROBINDO PHARMA	150MG	A090647 001	Jan 26, 2018
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AB		200MG	A090647 002	Jan 26, 2018
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AB		250MG	A090647 003	Jan 26, 2018
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AB		300MG	A090647 004	Jan 26, 2018
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AB	CHARTWELL	300MG	A206481 001	Jul 26, 2018
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AB	CIPLA	300MG	A078800 001	Jan 26, 2018
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AB	HETERO LABS LTD III	300MG	A090636 001	Jan 26, 2018
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AB	MACLEODS PHARMS LTD	300MG	A203232 001	Jan 26, 2018
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AB	QILU	200MG	A209498 001	Mar 02, 2018
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AB		250MG	A209498 002	Mar 02, 2018
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AB		300MG	A209498 003	Mar 02, 2018
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AB	REYOUNG	300MG	A211337 001	Mar 02, 2023
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AB	STRIDES PHARMA	300MG	A090742 001	Jan 26, 2018
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AB	TEVA PHARMS USA	150MG	A091612 002	Jan 26, 2018
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AB		200MG	A091612 003	Jan 26, 2018
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AB		250MG	A091612 004	Jan 26, 2018
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AB		300MG	A091612 001	Mar 18, 2015
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VIREAD

AB	+ GILEAD SCIENCES INC	150MG	N021356 002	Jan 18, 2012
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AB	+	200MG	N021356 003	Jan 18, 2012
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AB	+	250MG	N021356 004	Jan 18, 2012
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AB	+!	300MG	N021356 001	Oct 26, 2001
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TEPOTINIB HYDROCHLORIDE

TABLET;ORAL

TEPMETKO

+!	EMD SERONO INC	EQ 225MG BASE	N214096 001	Feb 03, 2021
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TERAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

TERAZOSIN HYDROCHLORIDE

AB	APNAR PHARMA LP	EQ 1MG BASE	A074823 001	Mar 30, 1998
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AB		EQ 2MG BASE	A074823 002	Mar 30, 1998
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AB		EQ 5MG BASE	A074823 003	Mar 30, 1998
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AB		EQ 10MG BASE	A074823 004	Mar 30, 1998
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AB	HERITAGE PHARMA	EQ 1MG BASE	A075614 002	Jan 30, 2001
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AVET

AB		EQ 2MG BASE	A075614 001	Jan 30, 2001
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AB		EQ 5MG BASE	A075614 003	Jan 30, 2001
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AB		EQ 10MG BASE	A075614 004	Jan 30, 2001
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AB	! JUBILANT CADISTA	EQ 1MG BASE	A075317 001	Dec 20, 2004
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AB		EQ 2MG BASE	A075317 002	Dec 20, 2004
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AB		EQ 5MG BASE	A075317 003	Dec 20, 2004
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AB		EQ 10MG BASE	A075317 004	Dec 20, 2004
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PRESCRIPTION DRUG PRODUCT LIST

TERBINAFINE HYDROCHLORIDE

TABLET; ORAL

TERBINAFINE HYDROCHLORIDE

<u>AB</u>	!	AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A078297 001</u>	Jul 02, 2007
<u>AB</u>		BRECKENRIDGE PHARM	<u>EQ 250MG BASE</u>	<u>A077714 001</u>	Jun 04, 2010
<u>AB</u>		CHARTWELL	<u>EQ 250MG BASE</u>	<u>A078199 001</u>	Jul 02, 2007
<u>AB</u>		CIPLA	<u>EQ 250MG BASE</u>	<u>A077137 001</u>	Jul 02, 2007
<u>AB</u>		DR REDDYS LABS INC	<u>EQ 250MG BASE</u>	<u>A076390 001</u>	Jul 02, 2007
<u>AB</u>		EMED MEDCL	<u>EQ 250MG BASE</u>	<u>A077919 001</u>	Jul 02, 2007
<u>AB</u>		GLENMARK GENERICS	<u>EQ 250MG BASE</u>	<u>A078157 001</u>	Jul 02, 2007
<u>AB</u>		INVAGEN PHARMS	<u>EQ 250MG BASE</u>	<u>A077533 001</u>	Jul 02, 2007
<u>AB</u>		ORBION PHARMS	<u>EQ 250MG BASE</u>	<u>A078163 001</u>	Jul 02, 2007

TERBUTALINE SULFATE

INJECTABLE; INJECTION

TERBUTALINE SULFATE

<u>AP</u>	!	CHARTWELL INJECTABLE	<u>1MG/ML</u>	<u>A076770 001</u>	Apr 23, 2004
<u>AP</u>		FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A076887 001</u>	May 26, 2004
<u>AP</u>		HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A078630 001</u>	May 20, 2009
<u>AP</u>		UNITED BIOMEDCL	<u>1MG/ML</u>	<u>A200122 001</u>	Nov 08, 2013

TABLET; ORAL

BRETHINE

<u>AB</u>	+	ANI PHARMS	<u>2.5MG</u>	<u>N017849 001</u>	
<u>AB</u>	+		<u>5MG</u>	<u>N017849 002</u>	

TERBUTALINE SULFATE

<u>AB</u>		IMPAX LABS	<u>2.5MG</u>	<u>A075877 001</u>	Jun 26, 2001
<u>AB</u>			<u>5MG</u>	<u>A075877 002</u>	Jun 26, 2001
<u>AB</u>		LANNETT CO INC	<u>2.5MG</u>	<u>A077152 001</u>	Mar 25, 2005
<u>AB</u>	!		<u>5MG</u>	<u>A077152 002</u>	Mar 25, 2005
<u>AB</u>		TWI PHARMS	<u>2.5MG</u>	<u>A211832 001</u>	Jun 19, 2020
<u>AB</u>			<u>5MG</u>	<u>A211832 002</u>	Jun 19, 2020

TERCONAZOLE

CREAM; VAGINAL

TERCONAZOLE

<u>AB</u>		FOUGERA PHARMS	<u>0.4%</u>	<u>A076712 001</u>	Feb 18, 2005
<u>AB</u>	!	TARO	<u>0.4%</u>	<u>A076043 001</u>	Jan 19, 2005
BX	+	FOUGERA PHARMS INC	0.8%	N021735 001	Oct 01, 2004
BX	!	TARO	0.8%	A075953 001	Apr 06, 2004

SUPPOSITORY; VAGINAL

TERCONAZOLE

<u>AB</u>	!	PADAGIS ISRAEL	<u>80MG</u>	<u>A077149 001</u>	Mar 17, 2006
<u>AB</u>		TARO	<u>80MG</u>	<u>A077553 001</u>	Mar 09, 2007

TERIFLUNOMIDE

TABLET; ORAL

AUBAGIO

<u>AB</u>	+	SANOFI AVENTIS US	<u>7MG</u>	<u>N202992 001</u>	Sep 12, 2012
<u>AB</u>	+		<u>14MG</u>	<u>N202992 002</u>	Sep 12, 2012

TERIFLUNOMIDE

<u>AB</u>		ACCORD HLTHCARE	<u>7MG</u>	<u>A209690 001</u>	Jan 07, 2019
<u>AB</u>			<u>14MG</u>	<u>A209690 002</u>	Jan 07, 2019
<u>AB</u>		ALEMBIC	<u>7MG</u>	<u>A209572 001</u>	Apr 19, 2019
<u>AB</u>			<u>14MG</u>	<u>A209572 002</u>	Apr 19, 2019
<u>AB</u>		AMNEAL PHARMS CO	<u>7MG</u>	<u>A209613 001</u>	Sep 28, 2018
<u>AB</u>			<u>14MG</u>	<u>A209613 002</u>	Sep 28, 2018
<u>AB</u>		APOTEX	<u>7MG</u>	<u>A209601 001</u>	Nov 02, 2018
<u>AB</u>			<u>14MG</u>	<u>A209601 002</u>	Nov 02, 2018
<u>AB</u>		AUROBINDO PHARMA	<u>7MG</u>	<u>A209638 001</u>	Oct 26, 2018
<u>AB</u>			<u>14MG</u>	<u>A209638 002</u>	Oct 26, 2018
<u>AB</u>		BIOCON PHARMA	<u>7MG</u>	<u>A209639 001</u>	Mar 13, 2023
<u>AB</u>			<u>14MG</u>	<u>A209639 002</u>	Mar 13, 2023
<u>AB</u>		GLENMARK PHARMS	<u>7MG</u>	<u>A209663 001</u>	Nov 15, 2018
<u>AB</u>			<u>14MG</u>	<u>A209663 002</u>	Nov 15, 2018
<u>AB</u>		HETERO LABS LTD V	<u>7MG</u>	<u>A209598 001</u>	Mar 13, 2023
<u>AB</u>			<u>14MG</u>	<u>A209598 002</u>	Mar 13, 2023
<u>AB</u>		MSN	<u>7MG</u>	<u>A209623 001</u>	Apr 24, 2019
<u>AB</u>			<u>14MG</u>	<u>A209623 002</u>	Apr 24, 2019
<u>AB</u>		MYLAN	<u>7MG</u>	<u>A209702 001</u>	Feb 28, 2020
<u>AB</u>			<u>14MG</u>	<u>A209702 002</u>	Feb 28, 2020
<u>AB</u>		NATCO	<u>7MG</u>	<u>A209555 001</u>	May 15, 2023
<u>AB</u>			<u>14MG</u>	<u>A209555 002</u>	May 15, 2023
<u>AB</u>		SANDOZ	<u>7MG</u>	<u>A209710 001</u>	Jan 03, 2019

PRESCRIPTION DRUG PRODUCT LIST

TERIFLUNOMIDE

TABLET; ORAL

TERIFLUNOMIDE

<u>AB</u>		<u>14MG</u>	<u>A209710 002</u>	Jan 03, 2019
<u>AB</u>	SOLA PHARMS	<u>7MG</u>	<u>A209677 002</u>	Sep 28, 2023
<u>AB</u>		<u>14MG</u>	<u>A209677 001</u>	Jun 17, 2020
<u>AB</u>	TEVA PHARMS USA	<u>7MG</u>	<u>A209700 001</u>	Sep 04, 2018
<u>AB</u>		<u>14MG</u>	<u>A209700 002</u>	Sep 04, 2018
<u>AB</u>	ZYDUS PHARMS	<u>7MG</u>	<u>A209668 001</u>	Nov 30, 2018
<u>AB</u>		<u>14MG</u>	<u>A209668 002</u>	Nov 30, 2018

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

FORTEO

<u>AP</u>	+! LILLY	<u>0.6MG/2.4ML (0.25MG/ML)</u>	<u>N021318 002</u>	Jun 25, 2008
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TERIPARATIDE

<u>AP</u>	APOTEX	<u>0.6MG/2.4ML (0.25MG/ML)</u>	<u>A211097 001</u>	Nov 16, 2023
<u>AP</u>	TEVA PHARMS USA	<u>0.6MG/2.4ML (0.25MG/ML)</u>	<u>A208569 001</u>	Nov 16, 2023
	BONSITY			
	ALVOGEN	0.62MG/2.48ML (0.25MG/ML)	N211939 001	Oct 04, 2019

TERLIPRESSIN ACETATE

POWDER; INTRAVENOUS

TERLIVAZ

	+! MALLINCKRODT IRELAND	EQ 0.85MG BASE/VIAL	N022231 001	Sep 14, 2022
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TESTOSTERONE

GEL; TRANSDERMAL

TESTOSTERONE

<u>AB1</u>	ACTAVIS LABS UT INC	<u>25MG/2.5GM PACKET</u>	<u>A076737 001</u>	Jan 27, 2006
<u>AB1</u>	!	<u>50MG/5GM PACKET</u>	<u>A076737 002</u>	Jan 27, 2006
<u>AB1</u>	ENCUBE	<u>50MG/5GM PACKET</u>	<u>A212984 001</u>	Nov 09, 2021
<u>AB1</u>	STRIDES PHARMA	<u>25MG/2.5GM PACKET</u>	<u>A076744 001</u>	May 23, 2007
<u>AB1</u>		<u>50MG/5GM PACKET</u>	<u>A076744 002</u>	May 23, 2007

TESTIM

<u>AB2</u>	+! AUXILIUM PHARMS LLC	<u>50MG/5GM PACKET</u>	<u>N021454 001</u>	Oct 31, 2002
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TESTOSTERONE

<u>AB2</u>	ACTAVIS LABS UT INC	<u>1.62% (20.25MG/1.25GM PACKET)</u>	<u>A204570 002</u>	Jul 17, 2020
<u>AB2</u>		<u>1.62% (40.5MG/2.5GM PACKET)</u>	<u>A204570 003</u>	Jul 17, 2020
<u>AB2</u>		<u>50MG/5GM PACKET</u>	<u>A091073 001</u>	Sep 18, 2017
<u>AB2</u>	PADAGIS ISRAEL	<u>1.62% (20.25MG/1.25GM PACKET)</u>	<u>A205781 001</u>	Jul 12, 2017
<u>AB2</u>	!	<u>1.62% (40.5MG/2.5GM PACKET)</u>	<u>A205781 002</u>	Jul 12, 2017

VOGELXO

<u>AB2</u>	UPSHER SMITH LABS	<u>50MG/5GM PACKET</u>	<u>N204399 002</u>	Jun 04, 2014
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GEL, METERED; NASAL

NATESTO

	+! ACERUS	5.5MG/0.122GM ACTUATION	N205488 001	May 28, 2014
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GEL, METERED; TRANSDERMAL

ANDROGEL

<u>AB</u>	+! BESINS HLTHCARE	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>N022309 001</u>	Apr 29, 2011
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FORTESTA

<u>AB</u>	+! ENDO PHARMS	<u>10MG/0.5GM ACTUATION</u>	<u>N021463 001</u>	Dec 29, 2010
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TESTOSTERONE

<u>AB</u>	ACTAVIS LABS UT INC	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A204570 001</u>	Apr 10, 2019
<u>AB</u>		<u>10MG/0.5GM ACTUATION</u>	<u>A204571 001</u>	Aug 05, 2015
<u>AB</u>	AMNEAL	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A207373 001</u>	Apr 10, 2019
<u>AB</u>	ENCUBE	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A208620 001</u>	Apr 10, 2019
<u>AB</u>	LUPIN	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A208560 001</u>	Apr 10, 2019
<u>AB</u>	PADAGIS ISRAEL	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A204268 001</u>	Aug 04, 2015
<u>AB</u>	TWI PHARMS	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A209390 001</u>	Sep 23, 2019
<u>AB</u>	XIROMED	<u>1.62% (20.25MG/1.25GM ACTUATION)</u>	<u>A210835 001</u>	Apr 16, 2020
BX	! ACTAVIS LABS UT INC	12.5MG/1.25GM ACTUATION	A076737 003	Mar 09, 2015
	VOGELXO			
BX	UPSHER SMITH LABS	12.5MG/1.25GM ACTUATION	N204399 003	Jun 04, 2014

PELLET; IMPLANTATION

TESTOPEL

	! AUXILIUM PHARMS INC	75MG	A080911 001	
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SOLUTION, METERED; TRANSDERMAL

TESTOSTERONE

<u>AT</u>	ACTAVIS LABS UT INC	<u>30MG/1.5ML ACTUATION</u>	<u>A205328 001</u>	Aug 07, 2017
<u>AT</u>	CIPLA	<u>30MG/1.5ML ACTUATION</u>	<u>A209533 001</u>	Jan 29, 2018
<u>AT</u>	DASH PHARMS	<u>30MG/1.5ML ACTUATION</u>	<u>A212301 001</u>	Jan 11, 2021
<u>AT</u>	LUPIN LTD	<u>30MG/1.5ML ACTUATION</u>	<u>A208061 001</u>	Oct 23, 2017

PRESCRIPTION DRUG PRODUCT LIST

TESTOSTERONE

SOLUTION, METERED; TRANSDERMAL

TESTOSTERONE

AT	!	PADAGIS ISRAEL	30MG/1.5ML ACTUATION	A204255	001	Feb 28, 2017
AT		TWI PHARMS	30MG/1.5ML ACTUATION	A209836	001	Sep 03, 2021

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

AO	+	!	PFIZER	100MG/ML	A085635	002
AO	+			200MG/ML	A085635	003

TESTOSTERONE CYPIONATE

AO			AM REGENT	200MG/ML	A207742	001	Jun 16, 2017
AO			CIPLA	100MG/ML	A210362	001	Jun 19, 2018
AO				200MG/ML	A210362	002	Jun 19, 2018
AO			EUGIA PHARMA	100MG/ML	A211817	001	Oct 20, 2023
AO				200MG/ML	A211817	002	Oct 20, 2023
AO			HIKMA	100MG/ML	A090387	001	Jul 15, 2010
AO				200MG/ML	A090387	002	Jul 15, 2010
AO			HIKMA FARMACEUTICA	200MG/ML	A091244	001	May 01, 2012
AO			PADAGIS US	200MG/ML	A040530	001	Jan 31, 2005
AO			SANDOZ	100MG/ML	A040615	001	Aug 10, 2006
AO				200MG/ML	A040615	002	Aug 10, 2006
AO			SUN PHARM INDS LTD	100MG/ML	A201720	001	Jun 03, 2013
AO				200MG/ML	A201720	002	Jun 03, 2013
AO			WATSON PHARMS INC	200MG/ML	A086030	001	
AO			WILSHIRE PHARMS INC	200MG/ML	A206368	001	Apr 24, 2019
AO			XIROMED	100MG/ML	A215351	001	Apr 13, 2023
AO				200MG/ML	A215351	002	Apr 13, 2023

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

TESTOSTERONE ENANTHATE

AO			HIKMA FARMACEUTICA	200MG/ML	A091120	001	Sep 18, 2012
AO			NEXUS	200MG/ML	A040575	001	Jun 14, 2006
AO	!		WATSON PHARMS INC	200MG/ML	A085598	001	

SOLUTION; SUBCUTANEOUS

XYOSTED (AUTOINJECTOR)

	+	!	ANTARES PHARMA INC	50MG/0.5ML (50MG/0.5ML)	N209863	001	Sep 28, 2018
	+			75MG/0.5ML (75MG/0.5ML)	N209863	002	Sep 28, 2018
	+			100MG/0.5ML (100MG/0.5ML)	N209863	003	Sep 28, 2018

TESTOSTERONE UNDECANOATE

CAPSULE; ORAL

JATENZO

	+		TOLMAR	158MG	N206089	001	Mar 27, 2019
	+			198MG	N206089	002	Mar 27, 2019
	+	!		237MG	N206089	003	Mar 27, 2019

KYZATREX

	+		MARIUS PHARMS LLC	100MG	N213953	001	Jul 27, 2022
	+			150MG	N213953	002	Jul 27, 2022
	+	!		200MG	N213953	003	Jul 27, 2022

TLANDO

	+	!	ANTARES PHARMA INC	112.5MG	N208088	001	Mar 28, 2022
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INJECTABLE; INTRAMUSCULAR

AVEED

	+	!	ENDO PHARMS INC	750MG/3ML (250MG/ML)	N022219	001	Mar 05, 2014
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TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

AB			ACTAVIS LABS FL INC	25MG	A206686	001	Jul 07, 2017
AB			APOTEX	12.5MG	A206093	001	Mar 17, 2020
AB				25MG	A206093	002	Mar 17, 2020
AB			BIONPHARMA	12.5MG	A208826	001	Dec 18, 2017
AB				25MG	A208826	002	Dec 18, 2017
AB			CHARTWELL RX	12.5MG	A210544	001	Apr 20, 2018
AB				25MG	A210544	002	Apr 20, 2018
AB			DR REDDYS	12.5MG	A209284	001	Jan 08, 2018
AB				25MG	A209284	002	Jan 08, 2018
AB			HETERO LABS LTD V	12.5MG	A204574	001	Feb 03, 2016
AB				25MG	A204574	002	Feb 03, 2016
AB			MYLAN	12.5MG	A207682	001	Jan 31, 2017
AB				25MG	A207682	002	Jan 31, 2017

PRESCRIPTION DRUG PRODUCT LIST

TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

AB	PIRAMAL HLTHCARE UK	12.5MG	A213316 001	Jan 22, 2020
AB		25MG	A213316 002	Jan 22, 2020

XENAZINE

AB	+ BAUSCH	12.5MG	N021894 001	Aug 15, 2008
AB	+!	25MG	N021894 002	Aug 15, 2008

TETRACAINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

TETRACAINE HYDROCHLORIDE

+	ALCON LABS	0.5%	N208135 001	Feb 29, 2016
+	BAUSCH LOMB IRELAND	0.5%	N210821 001	Mar 12, 2019

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

AB	AMNEAL PHARMS NY	250MG	A210674 001	Sep 18, 2018
AB	!	500MG	A210674 002	Sep 18, 2018
AB	BRECKENRIDGE	250MG	A210662 001	Nov 07, 2018
AB		500MG	A210662 002	Nov 07, 2018
AB	CHARTWELL TETRA	250MG	A062752 001	Aug 12, 1988
AB		500MG	A062752 002	Aug 12, 1988
AB	STRIDES PHARMA	250MG	A212635 001	Mar 03, 2020
AB		500MG	A212635 002	Mar 03, 2020
AB	WATSON LABS	250MG	A061837 001	
AB		500MG	A061837 002	

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION; NASAL

TYZINE

!	FOUGERA PHARMS	0.05%	A086576 002	
		0.1%	A086576 001	

SPRAY; NASAL

TYZINE

!	FOUGERA PHARMS	0.1%	A086576 003	
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THALIDOMIDE

CAPSULE; ORAL

THALOMID

+	BRISTOL-MYERS	50MG	N020785 001	Jul 16, 1998
+		100MG	N020785 002	Jan 17, 2003
+		150MG	N020785 004	Jan 10, 2007
+	!	200MG	N020785 003	Jan 17, 2003

THALLOUS CHLORIDE TL-201

INJECTABLE; INJECTION

THALLOUS CHLORIDE TL 201

+	CURIUM	1mCi/ML	N018150 001	
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THEOPHYLLINE

CAPSULE, EXTENDED RELEASE; ORAL

THEO-24

	AUXILIUM PHARMS LLC	100MG	A087943 002	Aug 22, 1983
+		200MG	A087943 001	Aug 22, 1983
		300MG	A087943 003	Aug 22, 1983
!		400MG	A087943 004	Feb 28, 1992

SOLUTION; ORAL

THEOPHYLLINE

AA	!	CHARTWELL MOLECULAR	80MG/15ML	A091156 001	Apr 13, 2011
AA		TRIS PHARMA INC	80MG/15ML	A091586 001	Jun 15, 2012

SOLUTION, ELIXIR; ORAL

ELIXOPHYLLIN

AA	+	NOSTRUM LABS INC	80MG/15ML	A085186 001	
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THEOPHYLLINE

AA		PHARM ASSOC	80MG/15ML	A206344 001	Dec 16, 2016
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TABLET, EXTENDED RELEASE; ORAL

THEOCHRON

AB		NOSTRUM PHARMS LLC	300MG	A087400 002	Jan 11, 1983
AB			450MG	A087400 005	Aug 09, 2023

THEOPHYLLINE

AB		ALEMBIC	300MG	A090430 001	Oct 27, 2010
AB	!		450MG	A090430 002	Oct 27, 2010
AB		AMNEAL	300MG	A216276 001	Mar 20, 2023

PRESCRIPTION DRUG PRODUCT LIST

THEOPHYLLINE

TABLET, EXTENDED RELEASE;ORAL

THEOPHYLLINE

<u>AB</u>		<u>450MG</u>	<u>A216276 002</u>	Mar 20, 2023
<u>AB</u>	ANNORA PHARMA	<u>300MG</u>	<u>A217422 001</u>	Sep 08, 2023
<u>AB</u>		<u>450MG</u>	<u>A217422 002</u>	Sep 08, 2023
<u>AB</u>	BIONPHARMA	<u>300MG</u>	<u>A216300 001</u>	Apr 26, 2023
<u>AB</u>		<u>450MG</u>	<u>A216300 002</u>	Apr 26, 2023
<u>AB</u>	GLENMARK GENERICS	<u>400MG</u>	<u>A090355 001</u>	Jul 13, 2010
<u>AB</u>		<u>600MG</u>	<u>A090355 002</u>	Jul 13, 2010
<u>AB</u>	GLENMARK PHARMS LTD	<u>300MG</u>	<u>A212184 001</u>	Jun 03, 2021
<u>AB</u>		<u>450MG</u>	<u>A212184 002</u>	Jun 03, 2021
<u>AB</u>	HARMAN FINOCHEM	<u>300MG</u>	<u>A214806 001</u>	Oct 24, 2023
<u>AB</u>		<u>450MG</u>	<u>A214806 002</u>	Oct 24, 2023
<u>AB</u>	HERITAGE PHARMA AVET	<u>300MG</u>	<u>A089763 001</u>	Apr 30, 1990
<u>AB</u>		<u>450MG</u>	<u>A081236 001</u>	Nov 09, 1992
<u>AB</u>	LEADING	<u>300MG</u>	<u>A214950 001</u>	Jan 30, 2023
<u>AB</u>		<u>450MG</u>	<u>A214950 002</u>	Jan 30, 2023
<u>AB</u>	NOSTRUM LABS INC	<u>400MG</u>	<u>A040560 003</u>	Apr 21, 2006
<u>AB</u>	!	<u>600MG</u>	<u>A040560 002</u>	Apr 21, 2006
<u>AB</u>	RHODES PHARMS	<u>300MG</u>	<u>A214113 001</u>	May 11, 2023
<u>AB</u>	+	<u>400MG</u>	<u>A040086 002</u>	Sep 01, 1982
<u>AB</u>		<u>600MG</u>	<u>A040086 001</u>	Apr 15, 1996
<u>AB</u>	TEVA PHARMS INC	<u>300MG</u>	<u>A216961 001</u>	Oct 12, 2022
<u>AB</u>		<u>450MG</u>	<u>A216961 002</u>	Oct 12, 2022
	THEOCHRON			
	NOSTRUM PHARMS LLC	100MG	A087400 003	Feb 21, 1985
		200MG	A087400 004	Feb 21, 1985

THIAMINE HYDROCHLORIDE

INJECTABLE;INJECTION

THIAMINE HYDROCHLORIDE

<u>AP</u>	CAPLIN	<u>100MG/ML</u>	<u>A215692 001</u>	Mar 06, 2023
<u>AP</u>	DR REDDYS	<u>100MG/ML</u>	<u>A080571 001</u>	
<u>AP</u>	EUGIA PHARMA	<u>100MG/ML</u>	<u>A208703 001</u>	Apr 20, 2022
<u>AP</u>	+!	<u>100MG/ML</u>	<u>A080556 001</u>	
<u>AP</u>	FRESENIUS KABI USA	<u>100MG/ML</u>	<u>A091623 001</u>	Jun 25, 2012
<u>AP</u>	MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<u>A206106 001</u>	Dec 01, 2017
<u>AP</u>	SAGENT PHARMS INC	<u>100MG/ML</u>	<u>A080575 001</u>	
<u>AP</u>	WEST-WARD PHARMS INT	<u>100MG/ML</u>		
	!	DR REDDYS	200MG/ML	A080571 002

THIOGUANINE

TABLET;ORAL

THIOGUANINE

+! WAYLIS THERAP 40MG N012429 001

THIORIDAZINE HYDROCHLORIDE

TABLET;ORAL

THIORIDAZINE HYDROCHLORIDE

MYLAN 10MG A088004 002 Mar 15, 1983
 25MG A088004 003 Mar 15, 1983
 50MG A088004 004 Mar 15, 1983
 ! 100MG A088004 001 Nov 18, 1983

THIOTEPA

INJECTABLE;INJECTION

THIOTEPA

<u>AP</u>	BELOTECA	<u>15MG/VIAL</u>	<u>A211831 001</u>	Dec 08, 2021
<u>AP</u>	DR REDDYS	<u>15MG/VIAL</u>	<u>A210337 001</u>	May 04, 2018
<u>AP</u>	HENGRUI PHARMA	<u>15MG/VIAL</u>	<u>A209150 001</u>	May 04, 2018
<u>AP</u>	PENN LIFE	<u>15MG/VIAL</u>	<u>A208242 001</u>	Jan 10, 2020
<u>AP</u>	!	WEST-WARD PHARMS	<u>15MG/VIAL</u>	<u>A075547 001</u> Apr 02, 2001
	INT			

POWDER;INTRACAVITARY, INTRAVENOUS, INTRAVESICAL

TEPADINA

<u>AP</u>	+!	ADIENNE SA	<u>15MG/VIAL</u>	<u>N208264 001</u> Jan 26, 2017
<u>AP</u>	+!		<u>100MG/VIAL</u>	<u>N208264 002</u> Jan 26, 2017

THIOTEPA

<u>AP</u>	GLAND PHARMA LTD	<u>15MG/VIAL</u>	<u>A214222 001</u>	Mar 08, 2021
<u>AP</u>		<u>100MG/VIAL</u>	<u>A214222 002</u>	Jan 03, 2022
<u>AP</u>	HIKMA	<u>100MG/VIAL</u>	<u>A211755 001</u>	Sep 05, 2023
<u>AP</u>	KINDOS	<u>15MG/VIAL</u>	<u>A216037 001</u>	Dec 26, 2023
<u>AP</u>		<u>100MG/VIAL</u>	<u>A216037 002</u>	Dec 26, 2023

PRESCRIPTION DRUG PRODUCT LIST

THIOTEPA

POWDER; INTRACAVITARY, INTRAVENOUS, INTRAVESICAL

THIOTEPA

<u>AP</u>	MSN	<u>15MG/VIAL</u>	<u>A213049 001</u>	Mar 04, 2020
<u>AP</u>		<u>100MG/VIAL</u>	<u>A213049 002</u>	Mar 04, 2020

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

<u>AB</u>	AMNEAL	<u>1MG</u>	<u>A215456 001</u>	Feb 28, 2022
<u>AB</u>		<u>2MG</u>	<u>A215456 002</u>	Feb 28, 2022
<u>AB</u>		<u>5MG</u>	<u>A215456 003</u>	Feb 28, 2022
<u>AB</u>		<u>10MG</u>	<u>A215456 004</u>	Feb 28, 2022
<u>AB</u>	NOVITIUM PHARMA	<u>1MG</u>	<u>A211642 001</u>	Apr 05, 2019
<u>AB</u>		<u>2MG</u>	<u>A211642 002</u>	Apr 05, 2019
<u>AB</u>	!	<u>5MG</u>	<u>A211642 003</u>	Apr 05, 2019
<u>AB</u>		<u>10MG</u>	<u>A211642 004</u>	Apr 05, 2019
<u>AB</u>	RISING	<u>1MG</u>	<u>A071093 002</u>	Jun 23, 1987
<u>AB</u>		<u>2MG</u>	<u>A071093 003</u>	Jun 23, 1987
<u>AB</u>		<u>5MG</u>	<u>A071093 004</u>	Jun 23, 1987
<u>AB</u>		<u>10MG</u>	<u>A071093 001</u>	Jun 23, 1987

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

<u>AB</u>	+ CEPHALON	<u>2MG</u>	<u>N020646 005</u>	Apr 16, 1999
<u>AB</u>	+	<u>4MG</u>	<u>N020646 001</u>	Sep 30, 1997
<u>AB</u>	+	<u>12MG</u>	<u>N020646 002</u>	Sep 30, 1997
<u>AB</u>	+	<u>16MG</u>	<u>N020646 003</u>	Sep 30, 1997

TIAGABINE HYDROCHLORIDE

<u>AB</u>	MSN	<u>2MG</u>	<u>A214816 001</u>	Nov 16, 2021
<u>AB</u>		<u>4MG</u>	<u>A214816 002</u>	Nov 16, 2021
<u>AB</u>		<u>12MG</u>	<u>A214816 003</u>	Nov 16, 2021
<u>AB</u>		<u>16MG</u>	<u>A214816 004</u>	Nov 16, 2021
<u>AB</u>	SUN PHARM INDS	<u>2MG</u>	<u>A077555 001</u>	Nov 04, 2011
<u>AB</u>		<u>4MG</u>	<u>A077555 002</u>	Nov 04, 2011

TICAGRELOR

TABLET; ORAL

BRILINTA

<u>AB</u>	+	ASTRAZENECA	<u>90MG</u>	<u>N022433 001</u>	Jul 20, 2011
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TICAGRELOR

<u>AB</u>	ALKEM LABS LTD	<u>90MG</u>	<u>A208567 001</u>	Apr 21, 2023
<u>AB</u>	HISUN PHARM HANGZHOU	<u>90MG</u>	<u>A208575 001</u>	Jan 23, 2019
	BRILINTA + ASTRAZENECA	60MG	N022433 002	Sep 03, 2015

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

!	APOTEX	250MG	A075089 001	Jul 01, 1999
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TIGECYCLINE

POWDER; INTRAVENOUS

TIGECYCLINE

<u>AP</u>	AMNEAL	<u>50MG/VIAL</u>	<u>N211158 001</u>	Aug 02, 2018	
<u>AP</u>	APOTEX	<u>50MG/VIAL</u>	<u>A204439 001</u>	Dec 21, 2018	
<u>AP</u>	EUGIA PHARMA	<u>50MG/VIAL</u>	<u>A206335 001</u>	Jun 11, 2019	
<u>AP</u>	+	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>N205645 001</u>	Dec 01, 2016
<u>AP</u>	MEITHEAL	<u>50MG/VIAL</u>	<u>A214020 001</u>	May 13, 2021	
<u>AP</u>	SANDOZ	<u>50MG/VIAL</u>	<u>A091620 001</u>	May 27, 2015	

TYGACIL

<u>AP</u>	+	PF PRISM CV	<u>50MG/VIAL</u>	<u>N021821 001</u>	Jun 15, 2005
	TIGECYCLINE ACCORD HLTHCARE INC	50MG/VIAL	N208744 001	Jan 18, 2018	

TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

BETIMOL

+	THEA PHARMA	EQ 0.25% BASE	N020439 001	Mar 31, 1995
+		EQ 0.5% BASE	N020439 002	Mar 31, 1995

PRESCRIPTION DRUG PRODUCT LIST

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS;OPHTHALMIC

TIMOLOL MALEATE

<u>AB</u>	!	ALEMBIC	<u>EQ 0.25% BASE</u>	<u>A212942 001</u>	Oct 22, 2020
<u>AB</u>	!		<u>EQ 0.5% BASE</u>	<u>A212942 002</u>	Oct 22, 2020
<u>AB</u>		DR REDDYS	<u>EQ 0.25% BASE</u>	<u>A215733 001</u>	Sep 22, 2022
<u>AB</u>			<u>EQ 0.5% BASE</u>	<u>A215733 002</u>	Sep 22, 2022
<u>AB</u>		EUGIA PHARMA	<u>EQ 0.5% BASE</u>	<u>A213540 001</u>	Mar 23, 2023
<u>AB</u>		GLAND PHARMA LTD	<u>EQ 0.25% BASE</u>	<u>A214645 001</u>	Jun 24, 2022
<u>AB</u>			<u>EQ 0.5% BASE</u>	<u>A214645 002</u>	Feb 14, 2023
<u>AB</u>	+	SANDOZ	<u>EQ 0.25% BASE</u>	<u>N020963 001</u>	Oct 21, 1998
<u>AB</u>	+		<u>EQ 0.5% BASE</u>	<u>N020963 002</u>	Oct 21, 1998

TIMOPTIC-XE

<u>AB</u>	+	BAUSCH AND LOMB INC	<u>EQ 0.25% BASE</u>	<u>N020330 001</u>	Nov 04, 1993
<u>AB</u>	+		<u>EQ 0.5% BASE</u>	<u>N020330 002</u>	Nov 04, 1993

SOLUTION/DROPS;OPHTHALMIC

TIMOLOL MALEATE

<u>AT1</u>		BAUSCH AND LOMB	<u>EQ 0.25% BASE</u>	<u>A074778 001</u>	Mar 25, 1997
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A074776 001</u>	Mar 25, 1997
<u>AT1</u>		FDC LTD	<u>EQ 0.25% BASE</u>	<u>A077259 001</u>	Apr 30, 2008
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A077259 002</u>	Apr 30, 2008
<u>AT1</u>		MANKIND PHARMA	<u>EQ 0.25% BASE</u>	<u>A078771 001</u>	Sep 28, 2009
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A078771 002</u>	Sep 28, 2009
<u>AT1</u>		MICRO LABS	<u>EQ 0.25% BASE</u>	<u>A217343 001</u>	May 26, 2023
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A217343 002</u>	May 26, 2023
<u>AT1</u>		PACIFIC PHARMA	<u>EQ 0.25% BASE</u>	<u>A074746 001</u>	Mar 25, 1997
<u>AT1</u>			<u>EQ 0.5% BASE</u>	<u>A074747 001</u>	Mar 25, 1997
<u>AT1</u>	!	SANDOZ	<u>EQ 0.25% BASE</u>	<u>A074261 001</u>	Apr 28, 1995
<u>AT1</u>	!		<u>EQ 0.5% BASE</u>	<u>A074262 001</u>	Apr 28, 1995

TIMOPTIC

<u>AT1</u>	+	BAUSCH AND LOMB INC	<u>EQ 0.25% BASE</u>	<u>N018086 001</u>	
<u>AT1</u>	+		<u>EQ 0.5% BASE</u>	<u>N018086 002</u>	

ISTALOL

<u>AT2</u>	+	BAUSCH AND LOMB	<u>EQ 0.5% BASE</u>	<u>N021516 001</u>	Jun 04, 2004
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TIMOLOL MALEATE

<u>AT2</u>		APOTEX	<u>EQ 0.5% BASE</u>	<u>A204936 001</u>	Apr 17, 2015
<u>AT3</u>		AMRING PHARMS	<u>EQ 0.25% BASE</u>	<u>A212592 001</u>	Dec 13, 2021
<u>AT3</u>			<u>EQ 0.5% BASE</u>	<u>A212592 002</u>	Dec 13, 2021
<u>AT3</u>		EPIC PHARMA LLC	<u>EQ 0.5% BASE</u>	<u>A212291 001</u>	Sep 11, 2020
<u>AT3</u>		INGENUS PHARMS LLC	<u>EQ 0.25% BASE</u>	<u>A216533 001</u>	Sep 13, 2022
<u>AT3</u>			<u>EQ 0.5% BASE</u>	<u>A216533 002</u>	Sep 13, 2022
<u>AT3</u>		MICRO LABS	<u>EQ 0.5% BASE</u>	<u>A216596 001</u>	Sep 07, 2022
<u>AT3</u>		SENTISS PHARMA	<u>EQ 0.5% BASE</u>	<u>A217195 001</u>	May 08, 2023

TIMOPTIC IN OCULOSE

<u>AT3</u>	+	BAUSCH AND LOMB INC	<u>EQ 0.25% BASE</u>	<u>N019463 001</u>	Nov 05, 1986
<u>AT3</u>	+		<u>EQ 0.5% BASE</u>	<u>N019463 002</u>	Nov 05, 1986

TABLET;ORAL

TIMOLOL MALEATE

<u>AB</u>		MYLAN	<u>5MG</u>	<u>A072668 002</u>	Jun 08, 1990
<u>AB</u>			<u>10MG</u>	<u>A072668 003</u>	Jun 08, 1990
<u>AB</u>	!		<u>20MG</u>	<u>A072668 001</u>	Jun 08, 1990
<u>AB</u>		RISING	<u>5MG</u>	<u>A207556 001</u>	May 02, 2019
<u>AB</u>			<u>10MG</u>	<u>A207556 002</u>	May 02, 2019
<u>AB</u>			<u>20MG</u>	<u>A207556 003</u>	May 02, 2019

TINIDAZOLE

TABLET;ORAL

TINDAMAX

<u>AB</u>	+	MISSION PHARMA	<u>250MG</u>	<u>N021618 001</u>	May 17, 2004
<u>AB</u>	+		<u>500MG</u>	<u>N021618 002</u>	May 17, 2004

TINIDAZOLE

<u>AB</u>		CHARTWELL RX	<u>250MG</u>	<u>A202044 001</u>	Apr 30, 2012
<u>AB</u>			<u>500MG</u>	<u>A202044 002</u>	Apr 30, 2012
<u>AB</u>		EDENBRIDGE PHARMS	<u>250MG</u>	<u>A203808 001</u>	Aug 04, 2015
<u>AB</u>			<u>500MG</u>	<u>A203808 002</u>	Aug 04, 2015
<u>AB</u>		THINQ PHARM-CRO PVT	<u>250MG</u>	<u>A202489 001</u>	Oct 09, 2013
<u>AB</u>			<u>500MG</u>	<u>A202489 002</u>	Oct 09, 2013

PRESCRIPTION DRUG PRODUCT LIST

TIOPRONIN

TABLET; ORAL

THIOLA

AB	+ !	MISSION PHARMA	100MG	N019569	001	Aug 11, 1988
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TIOPRONIN

AB		PAR PHARM INC	100MG	A216198	001	Jun 02, 2022
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AB		TEVA PHARMS USA INC	100MG	A214326	001	Apr 26, 2021
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TABLET, DELAYED RELEASE; ORAL

THIOLA EC

AB	+ !	MISSION PHARMACAL	300MG	N211843	002	Jun 28, 2019
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TIOPRONIN

AB		AMNEAL	300MG	A216278	001	Aug 15, 2023
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THIOLA EC

	+	MISSION PHARMACAL	100MG	N211843	001	Jun 28, 2019
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TIOTROPIUM BROMIDE

POWDER; INHALATION

SPIRIVA

AB	+ !	BOEHRINGER INGELHEIM	EQ 0.018MG BASE/INH	N021395	001	Jan 30, 2004
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TIOTROPIUM BROMIDE

AB		LUPIN	EQ 0.018MG BASE/INH	A211287	001	Jun 20, 2023
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SPRAY, METERED; INHALATION

SPIRIVA RESPIMAT

	+ !	BOEHRINGER INGELHEIM	EQ 0.00125MG BASE/INH	N021936	002	Sep 15, 2015
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	+ !		EQ 0.0025MG BASE/INH	N021936	001	Sep 24, 2014
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TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE

TABLET; ORAL

LONSURF

AB	+	TAIHO ONCOLOGY	EQ 6.14MG BASE;15MG	N207981	001	Sep 22, 2015
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AB	+ !		EQ 8.19MG BASE;20MG	N207981	002	Sep 22, 2015
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TIPIRACIL HYDROCHLORIDE AND TRIFLURIDINE

AB		NATCO	EQ 6.14MG BASE;15MG	A214008	001	Jun 13, 2023
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AB			EQ 8.19MG BASE;20MG	A214008	002	Jun 13, 2023
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TIPRANAVIR

CAPSULE; ORAL

APTIVUS

	+ !	BOEHRINGER INGELHEIM	250MG	N021814	001	Jun 22, 2005
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TIRBANIBULIN

OINTMENT; TOPICAL

KLISYRI

	+ !	ALMIRALL	1%	N213189	001	Dec 14, 2020
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TIROFIBAN HYDROCHLORIDE

SOLUTION; INJECTION

AGGRASTAT

	+ !	MEDICURE	EQ 3.75MG BASE/15ML (EQ 0.25MG BASE/ML)	N020912	002	Aug 31, 2016
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SOLUTION; INTRAVENOUS

AGGRASTAT

AP	+	MEDICURE	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	N020913	002	May 17, 2002
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AP	+ !		EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	N020913	003	Apr 20, 2000
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TIROFIBAN HYDROCHLORIDE

AP		EUGIA PHARMA	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	A216379	001	May 01, 2023
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AP			EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	A216379	002	May 01, 2023
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AP		GLAND PHARMA LTD	EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	A206888	001	Apr 08, 2021
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AP		NEXUS	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	A213947	001	Feb 07, 2023
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AP			EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML)	A213947	002	Jul 24, 2023
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TIRZEPATIDE

SOLUTION; SUBCUTANEOUS

MOUNJARO

	+ !	ELI LILLY AND CO	2.5MG/0.5ML (2.5MG/0.5ML)	N215866	001	May 13, 2022
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	+ !		5MG/0.5ML (5MG/0.5ML)	N215866	002	May 13, 2022
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	+ !		7.5MG/0.5ML (7.5MG/0.5ML)	N215866	003	May 13, 2022
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	+ !		10MG/0.5ML (10MG/0.5ML)	N215866	004	May 13, 2022
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	+ !		12.5MG/0.5ML (12.5MG/0.5ML)	N215866	005	May 13, 2022
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	+ !		15MG/0.5ML (15MG/0.5ML)	N215866	006	May 13, 2022
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PRESCRIPTION DRUG PRODUCT LIST

TIRZEPATIDE

SOLUTION;SUBCUTANEOUS

ZEPBOUND

+	!	ELI LILLY AND CO	2.5MG/0.5ML (2.5MG/0.5ML)	N217806	001	Nov 08, 2023
+	!		5MG/0.5ML (5MG/0.5ML)	N217806	002	Nov 08, 2023
+	!		7.5MG/0.5ML (7.5MG/0.5ML)	N217806	003	Nov 08, 2023
+	!		10MG/0.5ML (10MG/0.5ML)	N217806	004	Nov 08, 2023
+	!		12.5MG/0.5ML (12.5MG/0.5ML)	N217806	005	Nov 08, 2023
+	!		15MG/0.5ML (15MG/0.5ML)	N217806	006	Nov 08, 2023

TIVOZANIB HYDROCHLORIDE

CAPSULE;ORAL

FOTIVDA

+		AVEO PHARMS	EQ 0.89MG BASE	N212904	001	Mar 10, 2021
+	!		EQ 1.34MG BASE	N212904	002	Mar 10, 2021

TIZANIDINE HYDROCHLORIDE

CAPSULE;ORAL

TIZANIDINE HYDROCHLORIDE

<u>AB</u>		ALEMBIC	<u>EQ 2MG BASE</u>	<u>A213223</u>	<u>001</u>	Jan 13, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A213223</u>	<u>002</u>	Jan 13, 2020
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A213223</u>	<u>003</u>	Jan 13, 2020
<u>AB</u>		ALKEM LABS LTD	<u>EQ 2MG BASE</u>	<u>A212196</u>	<u>001</u>	Mar 27, 2019
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A212196</u>	<u>002</u>	Mar 27, 2019
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A212196</u>	<u>003</u>	Mar 27, 2019
<u>AB</u>		APOTEX INC	<u>EQ 2MG BASE</u>	<u>A078868</u>	<u>001</u>	Feb 03, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A078868</u>	<u>002</u>	Feb 03, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A078868</u>	<u>003</u>	Feb 03, 2012
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 2MG BASE</u>	<u>A213544</u>	<u>001</u>	Mar 24, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A213544</u>	<u>002</u>	Mar 24, 2020
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A213544</u>	<u>003</u>	Mar 24, 2020
<u>AB</u>		CADILA PHARMS LTD	<u>EQ 2MG BASE</u>	<u>A210021</u>	<u>001</u>	Mar 26, 2019
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A210021</u>	<u>002</u>	Mar 26, 2019
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A210021</u>	<u>003</u>	Mar 26, 2019
<u>AB</u>		JUBILANT GENERICS	<u>EQ 2MG BASE</u>	<u>A209605</u>	<u>001</u>	Aug 04, 2017
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A209605</u>	<u>002</u>	Aug 04, 2017
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A209605</u>	<u>003</u>	Aug 04, 2017
<u>AB</u>		NOVAST LABS	<u>EQ 2MG BASE</u>	<u>A210267</u>	<u>001</u>	Mar 12, 2019
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A210267</u>	<u>002</u>	Mar 12, 2019
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A210267</u>	<u>003</u>	Mar 12, 2019
<u>AB</u>		RUBICON	<u>EQ 2MG BASE</u>	<u>A213798</u>	<u>001</u>	May 27, 2020
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A213798</u>	<u>002</u>	May 27, 2020
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A213798</u>	<u>003</u>	May 27, 2020
<u>AB</u>		ZYDUS PHARMS	<u>EQ 2MG BASE</u>	<u>A208622</u>	<u>001</u>	Mar 03, 2017
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A208622</u>	<u>002</u>	Mar 03, 2017
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A208622</u>	<u>003</u>	Mar 03, 2017
		<u>ZANAFLEX</u>				
<u>AB</u>	+	LEGACY PHARMA USA	<u>EQ 2MG BASE</u>	<u>N021447</u>	<u>001</u>	Aug 29, 2002
<u>AB</u>	+		<u>EQ 4MG BASE</u>	<u>N021447</u>	<u>002</u>	Aug 29, 2002
<u>AB</u>	+	!	<u>EQ 6MG BASE</u>	<u>N021447</u>	<u>003</u>	Aug 29, 2002

TABLET;ORAL

TIZANIDINE HYDROCHLORIDE

<u>AB</u>		ALKEM LABS LTD	<u>EQ 2MG BASE</u>	<u>A211798</u>	<u>001</u>	Jan 25, 2019	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A211798</u>	<u>002</u>	Jan 25, 2019	
<u>AB</u>		APOTEX	<u>EQ 2MG BASE</u>	<u>A076533</u>	<u>001</u>	Jan 16, 2004	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A076533</u>	<u>002</u>	Jan 16, 2004	
<u>AB</u>		CADILA	<u>EQ 2MG BASE</u>	<u>A208187</u>	<u>001</u>	Mar 09, 2018	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A208187</u>	<u>002</u>	Mar 09, 2018	
<u>AB</u>		CASI PHARMS INC	<u>EQ 2MG BASE</u>	<u>A076280</u>	<u>001</u>	Nov 26, 2002	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A076280</u>	<u>002</u>	Jun 27, 2002	
<u>AB</u>		DR REDDYS LABS INC	<u>EQ 2MG BASE</u>	<u>A076286</u>	<u>001</u>	Jul 03, 2002	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A076286</u>	<u>002</u>	Jul 03, 2002	
<u>AB</u>		EPIC PHARMA LLC	<u>EQ 2MG BASE</u>	<u>A076347</u>	<u>001</u>	Oct 11, 2002	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A076347</u>	<u>002</u>	Oct 11, 2002	
<u>AB</u>		OXFORD PHARMS	<u>EQ 2MG BASE</u>	<u>A076281</u>	<u>001</u>	Oct 20, 2003	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A076281</u>	<u>002</u>	Oct 20, 2003	
<u>AB</u>		SUN PHARM INDS INC	<u>EQ 2MG BASE</u>	<u>A076416</u>	<u>001</u>	Sep 29, 2003	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A076416</u>	<u>002</u>	Sep 29, 2003	
<u>AB</u>		UNICHEM LABS LTD	<u>EQ 2MG BASE</u>	<u>A091283</u>	<u>001</u>	Nov 28, 2012	
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A091283</u>	<u>002</u>	Nov 28, 2012	
		<u>ZANAFLEX</u>					
<u>AB</u>	+	!	LEGACY PHARMA USA	<u>EQ 4MG BASE</u>	<u>N020397</u>	<u>001</u>	Nov 27, 1996

PRESCRIPTION DRUG PRODUCT LIST

TOBRAMYCIN

OINTMENT;OPHTHALMIC

TOBREX

+! NOVARTIS 0.3% N050555 001

POWDER; INHALATION

TOBI PODHALER

+! MYLAN SPECIALITY LP 28MG N201688 001 Mar 22, 2013

SOLUTION; INHALATION

BETHKIS**AN** +! CHIESI **300MG/4ML** **N201820 001** Oct 12, 2012KITABIS PAK**AN** PULMOFLOW INC **300MG/5ML** **N205433 001** Dec 02, 2014TOBI**AN** +! MYLAN SPECIALITY LP **300MG/5ML** **N050753 001** Dec 22, 1997TOBRAMYCIN**AN** ALKEM LABS LTD **300MG/5ML** **A212848 001** Apr 01, 2021**AN** AMNEAL PHARMS **300MG/5ML** **A205501 001** Jul 13, 2015**AN** DR REDDYS LABS SA **300MG/5ML** **A207080 001** Jul 09, 2018**AN** HIKMA **300MG/5ML** **A201422 001** May 28, 2014**AN** LUPIN **300MG/5ML** **A208964 001** Mar 22, 2017**AN** MANKIND PHARMA **300MG/5ML** **A214478 001** Jun 30, 2023**AN** **300MG/4ML** **A216725 001** Sep 22, 2022**AN** MICRO LABS **300MG/5ML** **A217344 001** Jul 27, 2023**AN** SUN PHARM **300MG/5ML** **A207136 001** Dec 26, 2019**AN** TEVA PHARMS USA **300MG/5ML** **A091589 001** Oct 10, 2013**AN** **300MG/4ML** **A210915 001** Jun 26, 2019

SOLUTION/DROPS;OPHTHALMIC

TOBRAMYCIN**AT** ALEMBIC **0.3%** **A211847 001** Apr 19, 2019**AT** BAUSCH AND LOMB **0.3%** **A064052 001** Nov 29, 1993**AT** CHARTWELL RX **0.3%** **A065026 001** Sep 11, 2001**AT** GLAND PHARMA LTD **0.3%** **A212628 001** Mar 09, 2021**AT** SOMERSET THERAPS LLC **0.3%** **A207444 001** Jun 28, 2017TOBREX**AT** +! NOVARTIS **0.3%** **N050541 001****AT** SANDOZ **0.3%** **A062535 001** Dec 13, 1984TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE**AP** BAXTER HLTHCARE **EQ 40MG BASE/ML** **A206965 001** Jul 01, 2016**AP** EUGIA PHARMA **EQ 1.2GM BASE/VIAL** **A217519 001** Jun 21, 2023**AP** ! FRESENIUS KABI USA **EQ 10MG BASE/ML** **A065122 001** Nov 29, 2002**AP** ! **EQ 40MG BASE/ML** **A065122 002** Nov 29, 2002**AP** **EQ 1.2GM BASE/VIAL** **N050789 001** Jul 13, 2004**AP** GLAND PHARMA LTD **EQ 10MG BASE/ML** **A209621 001** Feb 11, 2021**AP** **EQ 40MG BASE/ML** **A209621 002** Feb 11, 2021**AP** **EQ 1.2GM BASE/VIAL** **A211189 001** May 18, 2021**AP** HAINAN POLY **EQ 1.2GM BASE/VIAL** **A217029 001** Feb 01, 2023**AP** HIKMA **EQ 40MG BASE/ML** **A063117 001** Apr 26, 1991**AP** HOSPIRA **EQ 40MG BASE/ML** **A063111 001** Apr 30, 1991**AP** MYLAN LABS LTD **EQ 40MG BASE/ML** **A065407 001** Mar 11, 2008**AP** XELLIA PHARMS APS **EQ 1.2GM BASE/VIAL** **A205685 001** Sep 16, 2014**AP** ! XGEN PHARMS **EQ 1.2GM BASE/VIAL** **A065013 001** Aug 17, 2001TOBRAMYCIN SULFATE (PHARMACY BULK)**AP** ! FRESENIUS KABI USA **EQ 40MG BASE/ML** **A065120 001** Nov 29, 2002**AP** GLAND PHARMA LTD **EQ 40MG BASE/ML** **A209346 001** Feb 09, 2021TOFACITINIB CITRATE

SOLUTION; ORAL

TOFACITINIB CITRATE**AA** SLAYBACK PHARMA LLC **EQ 1MG BASE/ML** **A216878 001** Sep 25, 2023XELJANZ**AA** +! PFIZER **EQ 1MG BASE/ML** **N213082 001** Sep 25, 2020

TABLET; ORAL

XELJANZ

+ PF PRISM CV EQ 5MG BASE N203214 001 Nov 06, 2012

+! EQ 10MG BASE N203214 002 May 30, 2018

TABLET, EXTENDED RELEASE; ORAL

XELJANZ XR

+ PFIZER EQ 11MG BASE N208246 001 Feb 23, 2016

+! EQ 22MG BASE N208246 002 Dec 12, 2019

PRESCRIPTION DRUG PRODUCT LIST

TOFERSEN

SOLUTION; INTRATHECAL

QALSODY

+! BIOGEN MA

100MG/15ML (6.7MG/ML)

N215887 001 Apr 25, 2023

TOLCAPONE

TABLET; ORAL

TASMARAB +! BAUSCH100MGN020697 001 Jan 29, 1998TOLCAPONEAB NOVAST LABS100MGA208937 001 Aug 07, 2018AB PAR PHARM INC100MGA204584 001 Mar 26, 2015TOLMETIN SODIUM

CAPSULE; ORAL

TOLMETIN SODIUM

RISING

EQ 400MG BASE

A073393 001 May 27, 1993

TABLET; ORAL

TOLMETIN SODIUM

! RISING

EQ 600MG BASE

A074473 001 Aug 30, 1994

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

DETROL LAAB + UPJOHN2MGN021228 001 Dec 22, 2000AB +!4MGN021228 002 Dec 22, 2000TOLTERODINE TARTRATEAB AJANTA PHARMA LTD2MGA213397 001 May 19, 2020AB4MGA213397 002 May 19, 2020AB AMTA2MGA213858 001 Feb 02, 2021AB4MGA213858 002 Feb 02, 2021AB HETERO LABS LTD III2MGA206419 001 Dec 12, 2017AB4MGA206419 002 Dec 12, 2017AB INVENTIA HLTHCARE2MGA204562 001 Aug 19, 2019AB4MGA204562 002 Aug 19, 2019AB TEVA PHARMS USA2MGA079141 001 Nov 22, 2016AB4MGA079141 002 Nov 22, 2016AB TORRENT2MGA203016 001 Aug 11, 2015AB4MGA203016 002 Aug 11, 2015

TABLET; ORAL

DETROLAB + UPJOHN1MGN020771 001 Mar 25, 1998AB +!2MGN020771 002 Mar 25, 1998TOLTERODINE TARTRATEAB ELYSIUM1MGA210775 001 Dec 30, 2019AB2MGA210775 002 Dec 30, 2019AB HETERO LABS LTD V1MGA204397 001 Aug 02, 2021AB2MGA204397 002 Aug 02, 2021AB IVAX SUB TEVA1MGA077006 001 Feb 23, 2015AB

PHARMS

2MGA077006 002 Feb 23, 2015AB MACLEODS PHARMS LTD1MGA203409 001 Aug 31, 2015AB2MGA203409 002 Aug 31, 2015AB UNICHEM1MGA205399 001 Aug 05, 2020AB2MGA205399 002 Aug 05, 2020AB UNIQUE1MGA204721 001 Jan 24, 2020AB2MGA204721 002 Jan 24, 2020TOLVAPTAN

TABLET; ORAL

SAMSCAAB + OTSUKA15MGN022275 001 May 19, 2009AB +!30MGN022275 002 May 19, 2009TOLVAPTANAB ALKEM LABS LTD15MGA211891 003 Sep 06, 2022AB30MGA211891 001 May 19, 2020AB APOTEX15MGA207605 002 Sep 06, 2022AB30MGA207605 001 May 19, 2020AB HETERO LABS LTD V15MGA205646 002 Sep 06, 2022AB30MGA205646 001 Jul 16, 2021AB MSN15MGA216949 001 Jun 21, 2023AB30MGA216949 002 Jun 21, 2023AB PAR PHARM INC15MGA206119 001 Feb 15, 2022AB30MGA206119 002 Feb 15, 2022

PRESCRIPTION DRUG PRODUCT LIST

TOLVAPTAN

TABLET; ORAL

JYNARQUE

+	OTSUKA	15MG	N204441	001	Apr 23, 2018
+		30MG	N204441	002	Apr 23, 2018
+	!	45MG	N204441	003	Apr 23, 2018
+		60MG	N204441	004	Apr 23, 2018
+		90MG	N204441	005	Apr 23, 2018

TOLVAPTAN

	ALKEM LABS LTD	60MG	A211891	002	May 19, 2020
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TOPIRAMATE

CAPSULE; ORAL

TOPAMAX

<u>AB</u>	+	JANSSEN PHARMS	<u>15MG</u>	<u>N020844</u>	<u>001</u>	Oct 26, 1998
<u>AB</u>	+	!	<u>25MG</u>	<u>N020844</u>	<u>002</u>	Oct 26, 1998

TOPIRAMATE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>15MG</u>	<u>A215449</u>	<u>001</u>	Sep 27, 2023
<u>AB</u>			<u>25MG</u>	<u>A215449</u>	<u>002</u>	Sep 27, 2023
<u>AB</u>		TEVA	<u>15MG</u>	<u>A076575</u>	<u>001</u>	Apr 17, 2009
<u>AB</u>			<u>25MG</u>	<u>A076575</u>	<u>002</u>	Apr 17, 2009
<u>AB</u>		TWI PHARMS	<u>15MG</u>	<u>A217694</u>	<u>001</u>	Dec 05, 2023
<u>AB</u>			<u>25MG</u>	<u>A217694</u>	<u>002</u>	Dec 05, 2023
<u>AB</u>		ZYDUS PHARMS USA INC	<u>15MG</u>	<u>A078877</u>	<u>001</u>	Oct 14, 2009
<u>AB</u>			<u>25MG</u>	<u>A078877</u>	<u>002</u>	Oct 14, 2009

CAPSULE, EXTENDED RELEASE; ORAL

TOPIRAMATE

<u>AB1</u>		ACTAVIS LABS FL	<u>25MG</u>	<u>A206210</u>	<u>001</u>	Mar 10, 2023
<u>AB1</u>			<u>50MG</u>	<u>A206210</u>	<u>002</u>	Mar 10, 2023
<u>AB1</u>			<u>100MG</u>	<u>A206210</u>	<u>003</u>	Mar 10, 2023
<u>AB1</u>			<u>200MG</u>	<u>A206210</u>	<u>004</u>	Mar 10, 2023
<u>AB1</u>		AJANTA PHARMA LTD	<u>25MG</u>	<u>A215663</u>	<u>001</u>	Aug 15, 2023
<u>AB1</u>			<u>50MG</u>	<u>A215663</u>	<u>002</u>	Aug 15, 2023
<u>AB1</u>			<u>100MG</u>	<u>A215663</u>	<u>003</u>	Aug 15, 2023
<u>AB1</u>			<u>200MG</u>	<u>A215663</u>	<u>004</u>	Aug 15, 2023
<u>AB1</u>		ALKEM LABS LTD	<u>25MG</u>	<u>A217248</u>	<u>001</u>	Jun 20, 2023
<u>AB1</u>			<u>50MG</u>	<u>A217248</u>	<u>002</u>	Jun 20, 2023
<u>AB1</u>			<u>100MG</u>	<u>A217248</u>	<u>003</u>	Jun 20, 2023
<u>AB1</u>			<u>200MG</u>	<u>A217248</u>	<u>004</u>	Jun 20, 2023
<u>AB1</u>		DR REDDYS	<u>25MG</u>	<u>A217231</u>	<u>001</u>	Nov 17, 2023
<u>AB1</u>			<u>50MG</u>	<u>A217231</u>	<u>002</u>	Nov 17, 2023
<u>AB1</u>			<u>100MG</u>	<u>A217231</u>	<u>003</u>	Nov 17, 2023
<u>AB1</u>			<u>200MG</u>	<u>A217231</u>	<u>004</u>	Nov 17, 2023
<u>AB1</u>		PAR PHARM	<u>25MG</u>	<u>A205976</u>	<u>002</u>	May 04, 2023
<u>AB1</u>			<u>50MG</u>	<u>A205976</u>	<u>003</u>	May 04, 2023
<u>AB1</u>			<u>100MG</u>	<u>A205976</u>	<u>004</u>	May 04, 2023
<u>AB1</u>			<u>200MG</u>	<u>A205976</u>	<u>001</u>	Mar 01, 2023
<u>AB1</u>		ZYDUS	<u>25MG</u>	<u>A207382</u>	<u>001</u>	Nov 24, 2017
<u>AB1</u>			<u>50MG</u>	<u>A207382</u>	<u>002</u>	Nov 24, 2017
<u>AB1</u>			<u>100MG</u>	<u>A207382</u>	<u>003</u>	Nov 24, 2017
<u>AB1</u>			<u>200MG</u>	<u>A207382</u>	<u>004</u>	Oct 30, 2023

TROKENDI XR

<u>AB1</u>	+	SUPERMUS PHARMS	<u>25MG</u>	<u>N201635</u>	<u>001</u>	Aug 16, 2013
<u>AB1</u>	+		<u>50MG</u>	<u>N201635</u>	<u>002</u>	Aug 16, 2013
<u>AB1</u>	+		<u>100MG</u>	<u>N201635</u>	<u>003</u>	Aug 16, 2013
<u>AB1</u>	+	!	<u>200MG</u>	<u>N201635</u>	<u>004</u>	Aug 16, 2013

OUDEXY XR

<u>AB2</u>	+	UPSHER SMITH LABS	<u>25MG</u>	<u>N205122</u>	<u>001</u>	Mar 11, 2014
<u>AB2</u>	+		<u>50MG</u>	<u>N205122</u>	<u>002</u>	Mar 11, 2014
<u>AB2</u>	+		<u>100MG</u>	<u>N205122</u>	<u>003</u>	Mar 11, 2014
<u>AB2</u>	+		<u>150MG</u>	<u>N205122</u>	<u>004</u>	Mar 11, 2014
<u>AB2</u>	+	!	<u>200MG</u>	<u>N205122</u>	<u>005</u>	Mar 11, 2014

TOPIRAMATE

<u>AB2</u>		GLENMARK PHARMS LTD	<u>25MG</u>	<u>A210278</u>	<u>001</u>	Feb 01, 2021
<u>AB2</u>			<u>50MG</u>	<u>A210278</u>	<u>002</u>	Feb 01, 2021
<u>AB2</u>			<u>100MG</u>	<u>A210278</u>	<u>003</u>	Feb 01, 2021
<u>AB2</u>			<u>150MG</u>	<u>A210278</u>	<u>004</u>	Feb 01, 2021
<u>AB2</u>			<u>200MG</u>	<u>A210278</u>	<u>005</u>	Feb 01, 2021
<u>AB2</u>		ZYDUS	<u>25MG</u>	<u>A208949</u>	<u>001</u>	Nov 29, 2022
<u>AB2</u>			<u>50MG</u>	<u>A208949</u>	<u>002</u>	Nov 29, 2022
<u>AB2</u>			<u>100MG</u>	<u>A208949</u>	<u>003</u>	Nov 29, 2022

PRESCRIPTION DRUG PRODUCT LIST

TOPIRAMATE

CAPSULE, EXTENDED RELEASE;ORAL

TOPIRAMATE

<u>AB2</u>		<u>150MG</u>	<u>A208949 004</u>	Nov 29, 2022
<u>AB2</u>		<u>200MG</u>	<u>A208949 005</u>	Nov 29, 2022

SOLUTION;ORAL

EPRONTIA

+! AZURITY

25MG/ML

N214679 001 Nov 05, 2021

TABLET;ORAL

TOPAMAX

<u>AB</u>	+	JANSSEN PHARMS	<u>25MG</u>	<u>N020505 004</u>	Dec 24, 1996
<u>AB</u>	+		<u>50MG</u>	<u>N020505 005</u>	Dec 24, 1996
<u>AB</u>	+!		<u>100MG</u>	<u>N020505 001</u>	Dec 24, 1996
<u>AB</u>	+		<u>200MG</u>	<u>N020505 002</u>	Dec 24, 1996

TOPIRAMATE

<u>AB</u>		ACCORD HLTHCARE	<u>25MG</u>	<u>A076311 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A076311 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076311 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A076311 004</u>	Mar 27, 2009
<u>AB</u>		ASCENT PHARMS INC	<u>25MG</u>	<u>A215414 001</u>	Aug 26, 2021
<u>AB</u>			<u>50MG</u>	<u>A215414 002</u>	Aug 26, 2021
<u>AB</u>			<u>100MG</u>	<u>A215414 003</u>	Aug 26, 2021
<u>AB</u>			<u>200MG</u>	<u>A215414 004</u>	Aug 26, 2021
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A078462 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A078462 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A078462 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A078462 004</u>	Mar 27, 2009
<u>AB</u>		CIPLA LTD	<u>25MG</u>	<u>A076343 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A076343 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076343 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A076343 004</u>	Mar 27, 2009
<u>AB</u>		GLENMARK GENERICS	<u>25MG</u>	<u>A077627 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A077627 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A077627 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A077627 004</u>	Mar 27, 2009
<u>AB</u>		INVAGEN PHARMS	<u>25MG</u>	<u>A079162 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A079162 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A079162 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A079162 004</u>	Mar 27, 2009
<u>AB</u>		SUN PHARM	<u>25MG</u>	<u>A090278 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A090278 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A090278 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A090278 004</u>	Mar 27, 2009
<u>AB</u>		SUN PHARM INDS LTD	<u>25MG</u>	<u>A076327 001</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076327 002</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A076327 003</u>	Mar 27, 2009
<u>AB</u>		UNICHEM LABS LTD	<u>25MG</u>	<u>A090162 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A090162 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A090162 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A090162 004</u>	Feb 19, 2013
<u>AB</u>		VIWIT PHARM	<u>25MG</u>	<u>A077733 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A077733 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A077733 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A077733 004</u>	Mar 27, 2009
<u>AB</u>		ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078235 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A078235 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A078235 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A078235 004</u>	Mar 27, 2009

TOPOTECAN HYDROCHLORIDE

CAPSULE;ORAL

HYCAMTIN

+ SANDOZ

EQ 0.25MG BASE

N020981 001 Oct 11, 2007

+!

EQ 1MG BASE

N020981 002 Oct 11, 2007

INJECTABLE;INJECTION

HYCAMTIN

<u>AP</u>	+	SANDOZ	<u>EQ 4MG BASE/VIAL</u>	<u>N020671 001</u>	May 28, 1996
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TOPOTECAN HYDROCHLORIDE

<u>AP</u>		ACCORD HLTHCARE	<u>EQ 4MG BASE/VIAL</u>	<u>A202351 001</u>	Jun 26, 2013
<u>AP</u>		ACTAVIS TOTOWA	<u>EQ 4MG BASE/VIAL</u>	<u>A090620 001</u>	Dec 02, 2010
<u>AP</u>		DR REDDYS LABS LTD	<u>EQ 4MG BASE/VIAL</u>	<u>A201191 001</u>	Mar 09, 2011
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 4MG BASE/VIAL</u>	<u>A091089 001</u>	Nov 29, 2010

PRESCRIPTION DRUG PRODUCT LIST

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

TOPOTECAN HYDROCHLORIDE

<u>AP</u>	NOVAST LABS	<u>EQ 4MG BASE/VIAL</u>	<u>A206962 001</u>	Nov 30, 2016
<u>AP</u>	SAGENT PHARMS	<u>EQ 4MG BASE/VIAL</u>	<u>A091284 001</u>	Jan 26, 2011
<u>AP</u>	TEYRO LABS	<u>EQ 4MG BASE/VIAL</u>	<u>A091199 001</u>	Dec 01, 2010

SOLUTION; INTRAVENOUS

TOPOTECAN HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A204406 002</u>	Jul 06, 2017
<u>AP</u>	+! HOSPIRA INC	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N200582 001</u>	Feb 02, 2011
<u>AP</u>	MEITHEAL	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N022453 001</u>	Dec 20, 2012
	ACCORD HLTHCARE	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A204406 001	Jul 06, 2017

TOREMIFENE CITRATE

TABLET; ORAL

FARESTON

<u>AB</u>	+! KYOWA KIRIN	<u>EQ 60MG BASE</u>	<u>N020497 001</u>	May 29, 1997
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TOREMIFENE CITRATE

<u>AB</u>	MSN	<u>EQ 60MG BASE</u>	<u>A212818 001</u>	Aug 18, 2020
<u>AB</u>	RISING	<u>EQ 60MG BASE</u>	<u>A208813 001</u>	Dec 04, 2018

TORSEMIDE

TABLET; ORAL

TORSEMIDE

<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078249 001</u>	Oct 17, 2007
<u>AB</u>		<u>10MG</u>	<u>A078249 002</u>	Oct 17, 2007
<u>AB</u>		<u>20MG</u>	<u>A078249 003</u>	Oct 17, 2007
<u>AB</u>		<u>100MG</u>	<u>A078249 004</u>	Oct 17, 2007
<u>AB</u>	COREPHARMA	<u>5MG</u>	<u>A076894 001</u>	May 31, 2005
<u>AB</u>		<u>10MG</u>	<u>A076894 002</u>	May 31, 2005
<u>AB</u>		<u>20MG</u>	<u>A076894 003</u>	May 31, 2005
<u>AB</u>		<u>100MG</u>	<u>A076894 004</u>	May 31, 2005
<u>AB</u>	HETERO LABS LTD III	<u>5MG</u>	<u>A079234 001</u>	Jan 27, 2009
<u>AB</u>		<u>10MG</u>	<u>A079234 002</u>	Jan 27, 2009
<u>AB</u>		<u>20MG</u>	<u>A079234 003</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079234 004</u>	Jan 27, 2009
<u>AB</u>	HIKMA	<u>5MG</u>	<u>A076943 001</u>	Mar 01, 2005
<u>AB</u>		<u>10MG</u>	<u>A076943 002</u>	Mar 01, 2005
<u>AB</u>		<u>20MG</u>	<u>A076943 003</u>	Mar 01, 2005
<u>AB</u>	PLIVA PHARM IND	<u>5MG</u>	<u>A076346 001</u>	May 30, 2003
<u>AB</u>		<u>10MG</u>	<u>A076346 002</u>	May 30, 2003
<u>AB</u>	!	<u>20MG</u>	<u>A076346 003</u>	May 30, 2003
<u>AB</u>		<u>100MG</u>	<u>A076346 004</u>	Oct 19, 2004
<u>AB</u>	STRIDES PHARMA	<u>5MG</u>	<u>A076226 001</u>	May 27, 2003
<u>AB</u>		<u>5MG</u>	<u>A090613 001</u>	Mar 22, 2011
<u>AB</u>		<u>10MG</u>	<u>A076226 002</u>	May 27, 2003
<u>AB</u>		<u>10MG</u>	<u>A090613 002</u>	Mar 22, 2011
<u>AB</u>		<u>20MG</u>	<u>A076226 003</u>	May 27, 2003
<u>AB</u>		<u>20MG</u>	<u>A090613 003</u>	Mar 22, 2011
<u>AB</u>		<u>100MG</u>	<u>A076226 004</u>	May 27, 2003
<u>AB</u>		<u>100MG</u>	<u>A090613 004</u>	Mar 22, 2011
	SOAANZ			
	+ SARFE PHARMS	40MG	N213218 003	Nov 17, 2021

TRABECTEDIN

POWDER; INTRAVENOUS

YONDELIS

+!	JANSSEN PRODS	1MG/VIAL	N207953 001	Oct 23, 2015
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TRAMADOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CONZIP

+!	CIPHER PHARMS INC	100MG	N022370 001	May 07, 2010
+		200MG	N022370 002	May 07, 2010
+		300MG	N022370 003	May 07, 2010

SOLUTION; ORAL

QDOLO

+!	ATHENA	5MG/ML	N214044 001	Sep 01, 2020
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TABLET; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB</u>	ACI	<u>50MG</u>	<u>A202075 001</u>	Nov 28, 2011
<u>AB</u>	AMNEAL PHARMS	<u>50MG</u>	<u>A076003 001</u>	Jun 20, 2002
<u>AB</u>	APOTEX	<u>50MG</u>	<u>A075981 001</u>	Jul 10, 2002
<u>AB</u>	AUROBINDO PHARMA	<u>50MG</u>	<u>A203494 001</u>	Mar 31, 2014

PRESCRIPTION DRUG PRODUCT LIST

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

LTD

AB	CSPC OUYI PHARM CO	50MG	A091498 001	Mar 29, 2013
AB	IPCA LABS LTD	50MG	A201973 001	Nov 16, 2012
AB	MERRO PHARM USA	50MG	A206706 001	Jul 02, 2019
AB	RUBICON	50MG	A208708 001	Jun 28, 2019
AB	! SUN PHARM INDS INC	50MG	A075964 001	Jun 19, 2002
AB	TEVA	50MG	A075977 001	Jun 19, 2002
AB	UNICHEM	50MG	A211825 001	Aug 09, 2019
AB	ZYDUS PHARMS USA	50MG	A090404 001	Jan 31, 2011
	INC			
	RUBICON	25MG	A208708 003	Nov 27, 2023
		100MG	A208708 002	Jun 28, 2019

TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HYDROCHLORIDE

AB1	! LUPIN LTD	100MG	A200503 001	Aug 29, 2011
AB1		200MG	A200503 002	Aug 29, 2011
AB1		300MG	A200503 003	Aug 29, 2011
AB1	SUN PHARM	100MG	A201384 001	Dec 07, 2011
AB1		200MG	A201384 002	Dec 07, 2011
AB1		300MG	A201384 003	Dec 07, 2011
	!	100MG	A091607 001	Dec 30, 2011
		300MG	A091607 003	Dec 30, 2011

TRAMETINIB DIMETHYL SULFOXIDE

SOLUTION; ORAL

MEKINIST

+! NOVARTIS

EQ 0.05MG BASE/ML

N217513 001 Mar 16, 2023

TABLET; ORAL

MEKINIST

+ NOVARTIS

EQ 0.5MG

N204114 001 May 29, 2013

+!

EQ 2MG

N204114 003 May 29, 2013

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL

AB	AUROBINDO PHARMA	1MG	A078438 001	Jun 12, 2007
AB		2MG	A078438 002	Jun 12, 2007
AB		4MG	A078438 003	Jun 12, 2007
AB	EPIC PHARMA	1MG	A078508 003	Jun 18, 2008
AB		2MG	A078508 001	Jun 18, 2008
AB		4MG	A078508 002	Jun 18, 2008
AB	LUPIN	1MG	A077522 001	Jun 12, 2007
AB		2MG	A077522 002	Jun 12, 2007
AB	!	4MG	A077522 003	Jun 12, 2007

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE

GLENMARK GENERICS

1MG; 240MG

A079135 004 Aug 30, 2010

2MG; 180MG

A079135 001 May 26, 2010

2MG; 240MG

A079135 002 May 26, 2010

!

4MG; 240MG

A079135 003 May 05, 2010

TRANEXAMIC ACID

INJECTABLE; INJECTION

CYKLOKAPRON

AP	+! PFIZER	100MG/ML	N019281 001	Dec 30, 1986
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TRANEXAMIC ACID

AP	ACIC PHARMS	100MG/ML	A202436 001	Feb 11, 2014
AP	AM REGENT	100MG/ML	A201885 001	Aug 10, 2011
AP	AMNEAL PHARMS CO	100MG/ML	A208840 001	Feb 28, 2017
AP	APOTEX	100MG/ML	A209860 001	Jan 14, 2020
AP	AVET LIFESCIENCES	100MG/ML	A203521 001	Aug 12, 2014
AP	CAPLIN	100MG/ML	A212360 001	Jul 17, 2019
AP	EPIC PHARMA LLC	100MG/ML	A202373 001	Nov 17, 2011
AP	EUGIA PHARMA	100MG/ML	A205035 001	Jan 14, 2016
AP	FRESENIUS KABI USA	100MG/ML	A091596 001	Mar 02, 2012
AP	GLAND	100MG/ML	A207239 001	Feb 13, 2017
AP	MICRO LABS	100MG/ML	A206713 001	Jun 27, 2017
AP	MYLAN INSTITUTIONAL	100MG/ML	A091657 001	Nov 03, 2011
AP	PROVEPHARM SAS	100MG/ML	A212676 001	Jul 17, 2019
AP	XGEN PHARMS	100MG/ML	A201580 001	Jun 14, 2013

PRESCRIPTION DRUG PRODUCT LIST

TRANEXAMIC ACID

SOLUTION;INTRAVENOUS

TRANEXAMIC ACID

<u>AB</u>		AMNEAL	<u>1GM/100ML (10MG/ML)</u>	<u>A217155</u>	<u>001</u>	Oct 16, 2023
<u>AB</u>	+	EXELA PHARMA	<u>1GM/100ML (10MG/ML)</u>	<u>N212020</u>	<u>001</u>	Apr 15, 2019

TABLET;ORAL

LYSTEDA

<u>AB</u>	+	AMRING PHARMS	<u>650MG</u>	<u>N022430</u>	<u>001</u>	Nov 13, 2009
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TRANEXAMIC ACID

<u>AB</u>		ACTAVIS LABS FL INC	<u>650MG</u>	<u>A202093</u>	<u>001</u>	Dec 27, 2012
<u>AB</u>		ANI PHARMS	<u>650MG</u>	<u>A203256</u>	<u>001</u>	Jul 25, 2016
<u>AB</u>		AUROBINDO PHARMA USA	<u>650MG</u>	<u>A205133</u>	<u>001</u>	Sep 21, 2015

TRANLYCYPROMINE SULFATE

TABLET;ORAL

PARNATE

<u>AB</u>	+	CONCORDIA	<u>EQ 10MG BASE</u>	<u>N012342</u>	<u>003</u>	Aug 16, 1985
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TRANLYCYPROMINE SULFATE

<u>AB</u>		CROSSMEDIKA SA	<u>EQ 10MG BASE</u>	<u>A213503</u>	<u>001</u>	Jun 27, 2022
<u>AB</u>		NOVITIUM PHARMA	<u>EQ 10MG BASE</u>	<u>A206856</u>	<u>001</u>	Apr 17, 2018
<u>AB</u>		STRIDES PHARMA	<u>EQ 10MG BASE</u>	<u>A040640</u>	<u>001</u>	Jun 29, 2006

TRAVOPROST

IMPLANT;INTRACAMERAL

IDOSE TR

+! GLAUKOS

75MCG

N218010 001 Dec 13, 2023

SOLUTION/DROPS;OPHTHALMIC

TRAVATAN Z

<u>AT</u>	+	SANDOZ	<u>0.004%</u>	<u>N021994</u>	<u>001</u>	Sep 21, 2006
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TRAVOPROST

<u>AT</u>		APOTEX	<u>0.004%</u>	<u>A203431</u>	<u>001</u>	Jul 10, 2015
<u>AT</u>		MICRO LABS	<u>0.004%</u>	<u>A203767</u>	<u>001</u>	Mar 19, 2021
<u>AT</u>		MYLAN	<u>0.004%</u>	<u>A205050</u>	<u>001</u>	Jul 07, 2017
<u>AT1</u>		ALEMBIC	<u>0.004%</u>	<u>A210458</u>	<u>001</u>	Dec 20, 2019
<u>AT1</u>	!	CHARTWELL RX	<u>0.004%</u>	<u>A091340</u>	<u>001</u>	Mar 01, 2013

TRAZODONE HYDROCHLORIDE

TABLET;ORAL

TRAZODONE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>50MG</u>	<u>A206923</u>	<u>001</u>	Sep 08, 2017
<u>AB</u>			<u>100MG</u>	<u>A206923</u>	<u>002</u>	Sep 08, 2017
<u>AB</u>			<u>150MG</u>	<u>A206923</u>	<u>003</u>	Sep 08, 2017
<u>AB</u>			<u>300MG</u>	<u>A206923</u>	<u>004</u>	Sep 08, 2017
<u>AB</u>		APOTEX	<u>50MG</u>	<u>A071258</u>	<u>001</u>	Mar 25, 1987
<u>AB</u>	!	APOTEX INC	<u>100MG</u>	<u>A071196</u>	<u>001</u>	Mar 25, 1987
<u>AB</u>			<u>150MG</u>	<u>A071196</u>	<u>002</u>	Apr 26, 1999
<u>AB</u>			<u>300MG</u>	<u>A071196</u>	<u>003</u>	Apr 26, 1999
<u>AB</u>		AUROBINDO PHARMA	<u>50MG</u>	<u>A204852</u>	<u>001</u>	Feb 05, 2020
<u>AB</u>			<u>100MG</u>	<u>A204852</u>	<u>002</u>	Feb 05, 2020
<u>AB</u>			<u>150MG</u>	<u>A204852</u>	<u>003</u>	Feb 05, 2020
<u>AB</u>			<u>300MG</u>	<u>A204852</u>	<u>004</u>	Feb 05, 2020
<u>AB</u>		OXFORD PHARMS	<u>50MG</u>	<u>A072192</u>	<u>001</u>	Feb 02, 1989
<u>AB</u>			<u>100MG</u>	<u>A072193</u>	<u>001</u>	Feb 02, 1989
<u>AB</u>		PLIVA	<u>150MG</u>	<u>A071525</u>	<u>001</u>	Mar 09, 1988
<u>AB</u>		SUN PHARM INDUSTRIES	<u>50MG</u>	<u>A073137</u>	<u>002</u>	Mar 24, 1993
<u>AB</u>			<u>100MG</u>	<u>A073137</u>	<u>001</u>	Mar 24, 1993
<u>AB</u>			<u>150MG</u>	<u>A073137</u>	<u>003</u>	Dec 22, 1995
<u>AB</u>		TEVA PHARMS USA	<u>50MG</u>	<u>A071523</u>	<u>001</u>	Dec 11, 1987
<u>AB</u>			<u>100MG</u>	<u>A071524</u>	<u>001</u>	Dec 11, 1987
<u>AB</u>		TORRENT	<u>50MG</u>	<u>A202180</u>	<u>001</u>	Nov 27, 2013
<u>AB</u>			<u>100MG</u>	<u>A202180</u>	<u>002</u>	Nov 27, 2013
<u>AB</u>			<u>150MG</u>	<u>A202180</u>	<u>003</u>	Nov 27, 2013
<u>AB</u>			<u>300MG</u>	<u>A202180</u>	<u>004</u>	Nov 27, 2013
<u>AB</u>		ZYDUS PHARMS	<u>50MG</u>	<u>A205253</u>	<u>001</u>	Oct 10, 2017
<u>AB</u>			<u>100MG</u>	<u>A205253</u>	<u>002</u>	Oct 10, 2017
<u>AB</u>			<u>150MG</u>	<u>A205253</u>	<u>003</u>	Oct 10, 2017
<u>AB</u>			<u>300MG</u>	<u>A205253</u>	<u>004</u>	Oct 10, 2017

PRESCRIPTION DRUG PRODUCT LIST

TREPROSTINIL

INJECTABLE; IV (INFUSION), SUBCUTANEOUS

REMODULIN

<u>AP</u>	+	UNITED THERAP	<u>1MG/ML</u>	<u>N021272</u>	<u>001</u>	May 21, 2002
<u>AP</u>	+		<u>2.5MG/ML</u>	<u>N021272</u>	<u>002</u>	May 21, 2002
<u>AP</u>	+		<u>5MG/ML</u>	<u>N021272</u>	<u>003</u>	May 21, 2002
<u>AP</u>	+		<u>10MG/ML</u>	<u>N021272</u>	<u>004</u>	May 21, 2002

TREPROSTINIL

<u>AP</u>		ALEMBIC GLOBAL	<u>1MG/ML</u>	<u>A211574</u>	<u>001</u>	Feb 11, 2021
<u>AP</u>			<u>2.5MG/ML</u>	<u>A211574</u>	<u>002</u>	Feb 11, 2021
<u>AP</u>			<u>5MG/ML</u>	<u>A211574</u>	<u>003</u>	Feb 11, 2021
<u>AP</u>			<u>10MG/ML</u>	<u>A211574</u>	<u>004</u>	Feb 11, 2021
<u>AP</u>		DR REDDYS	<u>1MG/ML</u>	<u>A210214</u>	<u>001</u>	May 22, 2020
<u>AP</u>			<u>2.5MG/ML</u>	<u>A210214</u>	<u>002</u>	May 22, 2020
<u>AP</u>			<u>5MG/ML</u>	<u>A210214</u>	<u>003</u>	May 22, 2020
<u>AP</u>			<u>10MG/ML</u>	<u>A210214</u>	<u>004</u>	May 22, 2020
<u>AP</u>		PAR STERILE PRODUCTS	<u>1MG/ML</u>	<u>A209382</u>	<u>001</u>	Sep 24, 2019
<u>AP</u>			<u>2.5MG/ML</u>	<u>A209382</u>	<u>002</u>	Sep 24, 2019
<u>AP</u>			<u>5MG/ML</u>	<u>A209382</u>	<u>003</u>	Sep 24, 2019
<u>AP</u>			<u>10MG/ML</u>	<u>A209382</u>	<u>004</u>	Sep 24, 2019
<u>AP</u>		SANDOZ	<u>1MG/ML</u>	<u>A203649</u>	<u>001</u>	Nov 30, 2017
<u>AP</u>			<u>2.5MG/ML</u>	<u>A203649</u>	<u>002</u>	Nov 30, 2017
<u>AP</u>			<u>5MG/ML</u>	<u>A203649</u>	<u>003</u>	Nov 30, 2017
<u>AP</u>			<u>10MG/ML</u>	<u>A203649</u>	<u>004</u>	Nov 30, 2017
<u>AP</u>		TEVA PHARMS USA	<u>1MG/ML</u>	<u>A206648</u>	<u>001</u>	Sep 26, 2019
<u>AP</u>			<u>2.5MG/ML</u>	<u>A206648</u>	<u>002</u>	Sep 26, 2019
<u>AP</u>			<u>5MG/ML</u>	<u>A206648</u>	<u>003</u>	Sep 26, 2019
<u>AP</u>			<u>10MG/ML</u>	<u>A206648</u>	<u>004</u>	Sep 26, 2019

REMODULIN

	+	UNITED THERAP	0.1MG/ML	N021272	006	Sep 28, 2023
	+		0.2MG/ML	N021272	007	Sep 28, 2023
	+		0.4MG/ML	N021272	008	Sep 28, 2023
	+		20MG/ML	N021272	005	Jul 30, 2021

POWDER; INHALATION

TYVASO DPI

	+	UNITED THERAP	0.016MG/INH	N214324	001	May 23, 2022
	+		0.032MG/INH	N214324	002	May 23, 2022
	+		0.048MG/INH	N214324	003	May 23, 2022
	+		0.064MG/INH	N214324	004	May 23, 2022

SOLUTION; INHALATION

TYVASO

	+	UNITED THERAP	0.6MG/ML	N022387	001	Jul 30, 2009
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TREPROSTINIL DIOLAMINE

TABLET, EXTENDED RELEASE; ORAL

ORENITRAM

	+	UNITED THERAP	EQ 0.125MG BASE	N203496	001	Dec 20, 2013
	+		EQ 0.25MG BASE	N203496	002	Dec 20, 2013
	+		EQ 1MG BASE	N203496	003	Dec 20, 2013
	+		EQ 2.5MG BASE	N203496	004	Dec 20, 2013
	+		EQ 5MG BASE	N203496	005	Oct 07, 2016

TRETINOIN

CAPSULE; ORAL

TRETINOIN

<u>AB</u>		ANCHEN PHARMS	<u>10MG</u>	<u>A201687</u>	<u>001</u>	Oct 24, 2012
<u>AB</u>	!	BARR LABS INC	<u>10MG</u>	<u>A077684</u>	<u>001</u>	Jun 22, 2007
<u>AB</u>		GLENMARK PHARMS LTD	<u>10MG</u>	<u>A208279</u>	<u>001</u>	Dec 23, 2016

CREAM; TOPICAL

AVITA

<u>AB</u>		MYLAN PHARMS INC	<u>0.025%</u>	<u>N020404</u>	<u>003</u>	Jan 14, 1997
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RETIN-A

<u>AB</u>	+	BAUSCH	<u>0.025%</u>	<u>N019049</u>	<u>001</u>	Sep 16, 1988
<u>AB</u>	+	VALEANT PHARMS NORTH	<u>0.1%</u>	<u>N017340</u>	<u>001</u>	

TRETINOIN

<u>AB</u>		PADAGIS US	<u>0.025%</u>	<u>A075264</u>	<u>001</u>	Dec 24, 1998
<u>AB</u>			<u>0.1%</u>	<u>A075213</u>	<u>001</u>	Dec 24, 1998
<u>AB</u>		TARO	<u>0.025%</u>	<u>A215713</u>	<u>001</u>	Jan 05, 2023
<u>AB</u>			<u>0.1%</u>	<u>A211645</u>	<u>001</u>	Jan 22, 2019

RETIN-A

<u>AB1</u>	+	VALEANT BERMUDA	<u>0.05%</u>	<u>N017522</u>	<u>001</u>	
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PRESCRIPTION DRUG PRODUCT LIST

TRETINOIN

CREAM; TOPICAL

TRETINOIN

AB1	PADAGIS US	0.05%	A075265 001	Dec 24, 1998
AB1	TARO	0.05%	A211644 001	Jan 25, 2019
	RENOVA			
	+! VALEANT PHARMS NORTH	0.02%	N021108 001	Aug 31, 2000

GEL; TOPICAL

ATRALIN

AB	+! DOW PHARM	0.05%	N022070 001	Jul 26, 2007
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RETIN-A

AB	+! VALEANT INTL	0.01%	N017955 001	
AB	+!	0.025%	N017579 002	

RETIN-A-MICRO

AB	+! BAUSCH	0.08%	N020475 003	Jan 28, 2014
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TRETINOIN

AB	MYLAN	0.05%	A207955 001	Aug 13, 2015
AB	PADAGIS US	0.01%	A075589 001	Jun 11, 2002
AB		0.025%	A075529 001	Feb 22, 2000

TRETINOIN MICROSPHERE

AB	ENCUBE	0.08%	A215609 001	Aug 22, 2023
	RETIN-A MICRO			
	+! BAUSCH	0.04%	N020475 002	May 10, 2002
	+!	0.1%	N020475 001	Feb 07, 1997
	RETIN-A-MICRO			
	+! BAUSCH	0.06%	N020475 004	Oct 23, 2017

LOTION; TOPICAL

ALTRENO

	+! DOW PHARM	0.05%	N209353 001	Aug 23, 2018
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TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

TRIAMCINOLONE ACETONIDE

AT	ALKEM LABS LTD	0.025%	A207651 001	Dec 26, 2017
AT		0.1%	A207651 002	Dec 26, 2017
AT		0.5%	A207651 003	Dec 26, 2017
AT	CHARTWELL RX	0.025%	A208763 001	Feb 01, 2017
AT		0.1%	A208763 002	Feb 01, 2017
AT		0.5%	A208763 003	Feb 01, 2017
AT	COSETTE	0.025%	A089797 001	May 31, 1991
AT		0.1%	A089798 001	May 31, 1991
AT	ENCUBE	0.1%	A208848 001	Sep 18, 2017
AT	+! FOUGERA PHARMS	0.025%	A085692 001	
AT	+!	0.1%	A085692 003	
AT	+!	0.5%	A085692 002	
AT	GLENMARK PHARMS LTD	0.1%	A207117 001	Aug 05, 2016
AT	MACLEODS PHARMS LTD	0.025%	A209535 001	May 18, 2018
AT		0.1%	A209535 002	May 18, 2018
AT		0.5%	A209535 003	May 18, 2018
AT	MICRO LABS	0.025%	A040671 001	Jun 09, 2006
AT		0.1%	A040671 002	Jun 09, 2006
AT	PADAGIS US	0.025%	A086413 002	
AT		0.1%	A086413 003	
AT		0.5%	A086413 001	
AT	STRIDES PHARMA	0.025%	A210346 001	Feb 11, 2019
AT		0.1%	A210346 002	Feb 11, 2019
AT		0.5%	A210346 003	Feb 11, 2019
AT	TARO	0.1%	A040039 001	Nov 26, 1997

TRIDERM

AT	CROWN LABS	0.025%	A088042 002	Mar 25, 2015
AT		0.1%	A088042 001	Mar 19, 1984
AT		0.5%	A088042 003	Mar 25, 2015

FOR SUSPENSION, EXTENDED RELEASE; INTRA-ARTICULAR

ZILRETTA

	+! PACIRA PHARMS INC	32MG/VIAL	N208845 001	Oct 06, 2017
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INJECTABLE; INJECTION

KENALOG-40

AB	+! APOTHECON	40MG/ML	N014901 001	
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TRIAMCINOLONE ACETONIDE

AB	AMNEAL	40MG/ML	A207550 001	Dec 11, 2017
AB	EUGIA PHARMA	40MG/ML	A212400 001	Jul 05, 2022
AB	LONG GROVE PHARMS	40MG/ML	A213543 001	Jan 19, 2022

PRESCRIPTION DRUG PRODUCT LIST

TRIAMCINOLONE ACETONIDE

INJECTABLE; INJECTION

TRIAMCINOLONE ACETONIDE

AB	MYLAN LABS LTD	40MG/ML	A212567 001	Nov 02, 2022
AB	TEVA PHARMS USA	40MG/ML	A209852 001	Oct 05, 2018
	KENALOG-10			
	+ APOTHECON	10MG/ML	N012041 001	
	KENALOG-80			
	+! APOTHECON	80MG/ML	N014901 002	Apr 12, 2019
	INJECTABLE; INTRAVITREAL			
	TRIESENCE			
	+! HARROW EYE	40MG/ML (40MG/ML)	N022048 001	Nov 29, 2007

LOTION; TOPICAL

TRIAMCINOLONE ACETONIDE

AT	COSETTE	0.1%	A089129 001	Aug 14, 1986
AT	EPIC PHARMA LLC	0.025%	A202374 001	May 08, 2013
AT		0.1%	A202374 002	May 08, 2013
AT	FOUGERA PHARMS	0.025%	A040467 001	Apr 21, 2003
AT		0.1%	A040467 002	Apr 21, 2003
AT	MICRO LABS	0.1%	A040672 002	Dec 13, 2006
AT	QUAGEN	0.025%	A213559 001	Jul 01, 2020
AT		0.1%	A213559 002	Jul 01, 2020
AT	! WOCKHARDT BIO AG	0.025%	A088450 001	Apr 01, 1985
AT	!	0.1%	A088451 001	Apr 03, 1985

OINTMENT; TOPICAL

TRIAMCINOLONE ACETONIDE

AT	CHARTWELL RX	0.05%	A214532 001	Jan 27, 2023
AT		0.1%	A207365 001	Oct 12, 2018
AT	CINTEX SVCS	0.05%	A213619 001	Nov 10, 2020
AT	COSETTE	0.025%	A089795 001	Dec 23, 1988
AT		0.1%	A089796 001	Dec 23, 1988
AT		0.5%	A208925 001	Oct 06, 2017
AT	! ENCUBE	0.05%	A212384 001	Nov 29, 2019
AT		0.1%	A205373 001	May 13, 2016
AT	FOUGERA PHARMS	0.025%	A085691 001	
AT		0.1%	A085691 003	
AT		0.5%	A085691 002	
AT	GLENMARK PHARMS	0.1%	A208320 001	Aug 22, 2017
AT	GLENMARK PHARMS LTD	0.5%	A206379 001	Jul 22, 2016
AT	MACLEODS PHARMS LTD	0.025%	A209828 001	Nov 23, 2018
AT		0.05%	A216625 001	Aug 18, 2023
AT		0.1%	A209828 002	Nov 23, 2018
AT		0.5%	A209828 003	Nov 23, 2018
AT	PADAGIS ISRAEL	0.05%	A212460 001	Feb 05, 2021
AT	+! PADAGIS US	0.025%	A087385 002	
AT	+!	0.1%	A087385 003	
AT	+!	0.5%	A087385 001	
AT	STRIDES PHARMA	0.05%	A212356 001	Jun 01, 2020
AT	TARO	0.1%	A040037 001	Sep 30, 1994

PASTE; DENTAL

TRIAMCINOLONE ACETONIDE

AT	AKORN	0.1%	A206312 001	Aug 11, 2016
AT	COSETTE	0.1%	A205592 001	Jan 12, 2017
AT	LYNE	0.1%	A040771 001	Jul 01, 2010
AT	QUAGEN	0.1%	A214582 001	Dec 09, 2022
AT	! TARO	0.1%	A070730 001	Oct 01, 1986

SPRAY; TOPICAL

TRIAMCINOLONE ACETONIDE

AT	AKORN	0.147MG/GM	A207094 001	Dec 07, 2016
AT	! PADAGIS ISRAEL	0.147MG/GM	A205782 001	Apr 13, 2015
AT	RISING	0.147MG/GM	A206786 001	Sep 08, 2017

SUSPENSION; INJECTION

XIPERE

	+! BAUSCH AND LOMB INC	40MG/ML	N211950 001	Oct 22, 2021
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TRIAMTERENE

CAPSULE; ORAL

DYRENIUM

AB	+ CONCORDIA	50MG	N013174 001	
AB	+!	100MG	N013174 002	

TRIAMTERENE

AB	AGNITIO	50MG	A211581 001	Aug 19, 2019
AB		100MG	A211581 002	Aug 19, 2019

PRESCRIPTION DRUG PRODUCT LIST

TRIAZOLAM

TABLET; ORAL

HALCION

<u>AB</u>	+	PFIZER	<u>0.125MG</u>	<u>N017892</u>	<u>003</u>	Apr 26, 1985
<u>AB</u>	+	!	<u>0.25MG</u>	<u>N017892</u>	<u>001</u>	Nov 15, 1982

TRIAZOLAM

<u>AB</u>		HIKMA	<u>0.125MG</u>	<u>A074224</u>	<u>001</u>	Jun 01, 1994
<u>AB</u>			<u>0.25MG</u>	<u>A074224</u>	<u>002</u>	Jun 01, 1994
<u>AB</u>		NOVAST LABS	<u>0.125MG</u>	<u>A214219</u>	<u>001</u>	Oct 20, 2020
<u>AB</u>			<u>0.25MG</u>	<u>A214219</u>	<u>002</u>	Oct 20, 2020
<u>AB</u>		SANDOZ	<u>0.125MG</u>	<u>A216890</u>	<u>001</u>	Nov 08, 2022
<u>AB</u>			<u>0.25MG</u>	<u>A216890</u>	<u>002</u>	Nov 08, 2022
<u>AB</u>		ZYDUS PHARMS	<u>0.125MG</u>	<u>A213003</u>	<u>001</u>	Dec 28, 2022
<u>AB</u>			<u>0.25MG</u>	<u>A213003</u>	<u>002</u>	Dec 28, 2022

TRICLABENDAZOLE

TABLET; ORAL

EGATEN

+	!	NOVARTIS	250MG	N208711	001	Feb 13, 2019
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TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

SYPRINE

<u>AB</u>	+	!	BAUSCH	<u>250MG</u>	<u>N019194</u>	<u>001</u>	Nov 08, 1985
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TRIENTINE HYDROCHLORIDE

<u>AB</u>		DR REDDYS	<u>250MG</u>	<u>A211076</u>	<u>001</u>	Jul 03, 2019
<u>AB</u>		ECI PHARMS LLC	<u>250MG</u>	<u>A209945</u>	<u>001</u>	Aug 13, 2021
<u>AB</u>		HETERO LABS LTD III	<u>250MG</u>	<u>A216356</u>	<u>001</u>	Jun 23, 2022
<u>AB</u>		MSN	<u>250MG</u>	<u>A211134</u>	<u>001</u>	May 22, 2019
<u>AB</u>		NAVINTA LLC	<u>250MG</u>	<u>A211251</u>	<u>001</u>	Jan 16, 2019
<u>AB</u>		PAR PHARM INC	<u>250MG</u>	<u>A210096</u>	<u>001</u>	Sep 25, 2019
<u>AB</u>		RISING	<u>250MG</u>	<u>A212238</u>	<u>001</u>	Feb 20, 2020
<u>AB</u>		WATSON LABS TEVA	<u>250MG</u>	<u>A207567</u>	<u>001</u>	Feb 07, 2018
<u>AB</u>		ZYDUS PHARMS	<u>250MG</u>	<u>A211554</u>	<u>001</u>	Apr 26, 2019
		RISING	500MG	A212238	002	Sep 22, 2023

TRIENTINE TETRAHYDROCHLORIDE

TABLET; ORAL

CUVRIOR

+	!	ORPHALAN	300MG	N215760	001	Apr 28, 2022
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TRIFAROTENE

CREAM; TOPICAL

AKLIEF

+	!	GALDERMA LABS LP	0.005%	N211527	001	Oct 04, 2019
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TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL

TRIFLUOPERAZINE HYDROCHLORIDE

<u>AB</u>		MYLAN	<u>EQ 1MG BASE</u>	<u>A040209</u>	<u>001</u>	Jul 07, 1997
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A040209</u>	<u>002</u>	Jul 07, 1997
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A040209</u>	<u>003</u>	Jul 07, 1997
<u>AB</u>	!		<u>EQ 10MG BASE</u>	<u>A040209</u>	<u>004</u>	Jul 07, 1997
<u>AB</u>		SANDOZ	<u>EQ 1MG BASE</u>	<u>A085785</u>	<u>001</u>	
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A085786</u>	<u>001</u>	
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A085789</u>	<u>001</u>	
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A085788</u>	<u>001</u>	

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

<u>AT</u>		SANDOZ	<u>1%</u>	<u>A074311</u>	<u>001</u>	Oct 06, 1995
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VIROPTIC

<u>AT</u>	+	!	MONARCH PHARMS	<u>1%</u>	<u>N018299</u>	<u>001</u>
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TRIHHEPTANOIN

LIQUID; ORAL

DOJOLVI

+	!	ULTRAGENYX PHARM INC	100% w/w	N213687	001	Jun 30, 2020
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PRESCRIPTION DRUG PRODUCT LIST

TRIHEXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

AA	MIKART	2MG/5ML	A040251 001	Sep 27, 1999
AA	! PHARM ASSOC	2MG/5ML	A040177 001	Apr 17, 1997

TABLET; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

AA	NATCO PHARMA LTD	2MG	A091630 001	Nov 17, 2010
AA		5MG	A091630 002	Nov 17, 2010
AA	NOVITIUM PHARMA	2MG	A040254 001	Dec 24, 1998
AA		5MG	A040254 002	Dec 24, 1998
AA	! WATSON LABS	2MG	A084363 001	
AA	!	5MG	A084364 001	

TRILACICLIB DIHYDROCHLORIDE

POWDER; INTRAVENOUS

COSELA

+	! G1 THERAP	EQ 300MG BASE/VIAL	N214200 001	Feb 12, 2021
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TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TRIMETHOBENZAMIDE HYDROCHLORIDE

AB	AVET LIFESCIENCES	300MG	A205950 001	Nov 21, 2023
AB	LUPIN	300MG	A076546 001	Aug 20, 2003
AB	! SUN PHARM INDUSTRIES	300MG	A076570 001	Aug 28, 2003

INJECTABLE; INJECTION

TIGAN

+	! PAR STERILE PRODUCTS	100MG/ML	N017530 001	
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TRIMETHOPRIM

TABLET; ORAL

TRIMETHOPRIM

AB	! DR REDDYS LABS SA	100MG	N018679 001	Jul 30, 1982
AB	NOVEL LABS INC	100MG	A091437 001	Jun 15, 2011
AB	NOVITIUM PHARMA	100MG	A216393 001	Oct 28, 2022
AB	WATSON LABS	100MG	A070049 001	Jun 06, 1985

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

TRIMIPRAMINE MALEATE

AB	CROSSMEDIKA SA	EQ 25MG BASE	A208127 001	Apr 15, 2016
AB	!	EQ 50MG BASE	A208127 002	Apr 15, 2016
AB		EQ 100MG BASE	A208127 003	Apr 15, 2016
AB	ELITE LABS INC	EQ 25MG BASE	A077361 001	Aug 02, 2006
AB		EQ 50MG BASE	A077361 002	Aug 02, 2006
AB		EQ 100MG BASE	A077361 003	Aug 02, 2006

TRIPTORELIN PAMOATE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

TRIPTODUR KIT

+	! AZURITY	EQ 22.5MG BASE/VIAL	N208956 001	Jun 29, 2017
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INJECTABLE; INTRAMUSCULAR

TRELSTAR

+	! VERITY	EQ 3.75MG BASE/VIAL	N020715 001	Jun 15, 2000
+	!	EQ 11.25MG BASE/VIAL	N021288 001	Jun 29, 2001
+	!	EQ 22.5MG BASE/VIAL	N022437 001	Mar 10, 2010

TROFINETIDE

SOLUTION; ORAL

DAYBUE

+	! ACADIA PHARMS INC	200MG/ML	N217026 001	Mar 10, 2023
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TROMETHAMINE

SOLUTION; INJECTION

THAM

+	! HOSPIRA	18GM/500ML (3.6GM/100ML)	N013025 002	
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TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

MYDRIACYL

AT	! ALCON LABS INC	1%	A084306 001	
AT	! SANDOZ	0.5%	A084305 001	

TROPICACYL

AT	EPIC PHARMA LLC	0.5%	A040314 001	Sep 29, 2000
AT	RISING	1%	A040315 001	Sep 29, 2000

PRESCRIPTION DRUG PRODUCT LISTTROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

TROPICAMIDE

AT	BAUSCH AND LOMB	0.5%	A040067 001	Jul 27, 1994
AT		1%	A040064 001	Jul 27, 1994
AT	SOMERSET THERAPS LLC	1%	A207524 001	Dec 12, 2019

TROSPIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

TROSPIUM CHLORIDE

AB	!	ACTAVIS LABS FL INC	60MG	A091289 001	Oct 12, 2012
AB		AMTA	60MG	A214760 001	Apr 30, 2021
AB		GRANULES	60MG	A213185 001	Apr 23, 2020
AB		PADAGIS US	60MG	A201291 001	May 24, 2013

TABLET;ORAL

TROSPIUM CHLORIDE

AB		APOTEX	20MG	A091513 001	Dec 06, 2011
AB		CHARTWELL RX	20MG	A215781 001	Feb 16, 2022
AB	!	GLENMARK GENERICS	20MG	A091575 001	Aug 13, 2010
AB		HERITAGE PHARMS INC	20MG	A204945 001	Aug 30, 2016
AB		INVAGEN PHARMS	20MG	A091688 001	Aug 23, 2016
AB		PADAGIS US	20MG	A091573 001	Nov 17, 2010

TRYPAN BLUE

SOLUTION;OPHTHALMIC

VISIONBLUE

+	!	DORC	0.06%	N021670 001	Dec 16, 2004
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TUCATINIB

TABLET;ORAL

TUKYSA

+		SEAGEN	50MG	N213411 001	Apr 17, 2020
+	!		150MG	N213411 002	Apr 17, 2020

UBROGEPANT

TABLET;ORAL

UBRELVY

+		ABEVIE	50MG	N211765 001	Dec 23, 2019
+	!		100MG	N211765 002	Dec 23, 2019

ULIPRISTAL ACETATE

TABLET;ORAL

ELLA

AB	+	!	LAB HRA PHARMA	30MG	N022474 001	Aug 13, 2010
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LOGILIA

AB			TEVA PHARMS USA	30MG	A207952 001	Feb 13, 2017
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UMECLIDINIUM BROMIDE

POWDER; INHALATION

INCRUSE ELLIPTA

+	!	GLAXO GRP ENGLAND	EQ 62.5MCG BASE/INH	N205382 001	Apr 30, 2014
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UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER; INHALATION

ANORO ELLIPTA

+	!	GLAXOSMITHKLINE	EQ 0.0625MG BASE/INH;EQ 0.025MG BASE/INH	N203975 001	Dec 18, 2013
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UPADACITINIB

TABLET, EXTENDED RELEASE;ORAL

RINVOQ

+		ABEVIE	15MG	N211675 001	Aug 16, 2019
+			30MG	N211675 002	Jan 14, 2022
+	!		45MG	N211675 003	Mar 16, 2022

UREA, C-14

CAPSULE;ORAL

PYTEST

+	!	AVENT	1uCi	N020617 001	May 09, 1997
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URIDINE TRIACETATE

GRANULE;ORAL

VISTOGARD

+	!	BTG INTL	10GM/PACKET	N208159 001	Dec 11, 2015
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XURIDEN

+	!	BTG INTL	2GM/PACKET	N208169 001	Sep 04, 2015
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PRESCRIPTION DRUG PRODUCT LIST

URSODIOL

CAPSULE; ORAL

ACTIGALL

<u>AB</u>	<u>+!</u>	TEVA BRANDED PHARM	<u>300MG</u>	<u>N019594</u>	<u>002</u>	Dec 31, 1987
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URSODIOL

<u>AB</u>		ABHAI LLC	<u>300MG</u>	<u>A210707</u>	<u>001</u>	May 17, 2018
<u>AB</u>		AMNEAL PHARMS CO	<u>300MG</u>	<u>A211301</u>	<u>001</u>	Oct 16, 2018
<u>AB</u>		AUROBINDO PHARMA LTD	<u>300MG</u>	<u>A214849</u>	<u>001</u>	Nov 28, 2022
<u>AB</u>		CHARTWELL RX	<u>300MG</u>	<u>A213555</u>	<u>001</u>	Aug 17, 2020
<u>AB</u>		EPIC PHARMA	<u>300MG</u>	<u>A075517</u>	<u>001</u>	Mar 14, 2000
<u>AB</u>		EYWA PHARMA	<u>300MG</u>	<u>A212452</u>	<u>001</u>	Oct 30, 2019
<u>AB</u>		LANNETT CO INC	<u>300MG</u>	<u>A079082</u>	<u>001</u>	Dec 15, 2008
<u>AB</u>		RISING	<u>300MG</u>	<u>A213200</u>	<u>001</u>	Feb 12, 2020
<u>AB</u>		RK PHARMA	<u>300MG</u>	<u>A214329</u>	<u>001</u>	Jul 28, 2021
<u>AB</u>		STRIDES PHARMA	<u>300MG</u>	<u>A210344</u>	<u>001</u>	Jan 22, 2021
<u>AB</u>		ZYDUS	<u>300MG</u>	<u>A214295</u>	<u>001</u>	Oct 16, 2020
		LGM PHARMA	200MG	A205789	001	May 08, 2020
		!	400MG	A205789	002	May 08, 2020

TABLET; ORAL

URSO 250

<u>AB</u>	<u>+</u>	ALLERGAN	<u>250MG</u>	<u>N020675</u>	<u>001</u>	Dec 10, 1997
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URSO FORTE

<u>AB</u>	<u>+!</u>	ALLERGAN	<u>500MG</u>	<u>N020675</u>	<u>002</u>	Jul 21, 2004
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URSODIOL

<u>AB</u>		GLENMARK GENERICS	<u>250MG</u>	<u>A090801</u>	<u>001</u>	Jul 12, 2011
<u>AB</u>			<u>500MG</u>	<u>A090801</u>	<u>002</u>	Jul 12, 2011
<u>AB</u>		PAR PHARM	<u>250MG</u>	<u>A202540</u>	<u>001</u>	Feb 14, 2013
<u>AB</u>			<u>500MG</u>	<u>A202540</u>	<u>002</u>	Feb 14, 2013
<u>AB</u>		STRIDES PHARMA	<u>250MG</u>	<u>A213504</u>	<u>001</u>	Aug 20, 2020
<u>AB</u>			<u>500MG</u>	<u>A213504</u>	<u>002</u>	Aug 20, 2020
<u>AB</u>		ZYDUS	<u>250MG</u>	<u>A211145</u>	<u>001</u>	Oct 30, 2018
<u>AB</u>			<u>500MG</u>	<u>A211145</u>	<u>002</u>	Oct 30, 2018

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

<u>AB</u>		ADAPTIS	<u>EQ 500MG BASE</u>	<u>A079012</u>	<u>001</u>	May 24, 2010
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A079012</u>	<u>002</u>	May 24, 2010
<u>AB</u>		APOTEX	<u>EQ 500MG BASE</u>	<u>A090500</u>	<u>001</u>	Apr 04, 2014
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A090500</u>	<u>002</u>	Apr 04, 2014
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A090682</u>	<u>001</u>	May 24, 2010
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A090682</u>	<u>002</u>	May 24, 2010
<u>AB</u>		CADILA	<u>EQ 500MG BASE</u>	<u>A079137</u>	<u>001</u>	Dec 29, 2017
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A079137</u>	<u>002</u>	Dec 29, 2017
<u>AB</u>		CHARTWELL RX	<u>EQ 500MG BASE</u>	<u>A090216</u>	<u>001</u>	May 24, 2010
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A090216</u>	<u>002</u>	May 24, 2010
<u>AB</u>		CIPLA	<u>EQ 500MG BASE</u>	<u>A077135</u>	<u>001</u>	May 24, 2010
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A077135</u>	<u>002</u>	May 24, 2010
<u>AB</u>		HETERO LABS LTD V	<u>EQ 500MG BASE</u>	<u>A203047</u>	<u>001</u>	Apr 08, 2015
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A203047</u>	<u>002</u>	Apr 08, 2015
<u>AB</u>		JUBILANT GENERICS	<u>EQ 500MG BASE</u>	<u>A201506</u>	<u>001</u>	Apr 03, 2012
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A201506</u>	<u>002</u>	Apr 03, 2012
<u>AB</u>		MYLAN PHARMS INC	<u>EQ 500MG BASE</u>	<u>A078518</u>	<u>001</u>	May 24, 2010
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A078518</u>	<u>002</u>	May 24, 2010
<u>AB</u>		SANDOZ	<u>EQ 500MG BASE</u>	<u>A077478</u>	<u>001</u>	May 24, 2010
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A077478</u>	<u>002</u>	May 24, 2010
<u>AB</u>		SUN PHARM INDS LTD	<u>EQ 500MG BASE</u>	<u>A076588</u>	<u>001</u>	Jan 31, 2007
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A076588</u>	<u>002</u>	Jan 31, 2007
<u>AB</u>		YILING	<u>EQ 500MG BASE</u>	<u>A209553</u>	<u>001</u>	Mar 18, 2020
<u>AB</u>			<u>EQ 1GM BASE</u>	<u>A209553</u>	<u>002</u>	Mar 18, 2020

VALTREX

<u>AB</u>	<u>+</u>	GLAXOSMITHKLINE	<u>EQ 500MG BASE</u>	<u>N020487</u>	<u>001</u>	Jun 23, 1995
<u>AB</u>	<u>+!</u>		<u>EQ 1GM BASE</u>	<u>N020487</u>	<u>002</u>	Jun 23, 1995

VALBENZAZINE TOSYLATE

CAPSULE; ORAL

INGREZZA

	<u>+</u>	NEUROCRINE	EQ 40MG BASE	N209241	001	Apr 11, 2017
	<u>+</u>		EQ 60MG BASE	N209241	003	Apr 23, 2021
	<u>+!</u>		EQ 80MG BASE	N209241	002	Oct 04, 2017

PRESCRIPTION DRUG PRODUCT LIST

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION; ORAL

VALCYTE

AB	+!	CHEPLAPHARM	50MG/ML	N022257	001	Aug 28, 2009
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VALGANCICLOVIR HYDROCHLORIDE

AB		ACTAVIS LABS FL INC	50MG/ML	A205220	001	Jul 18, 2016
AB		AJANTA PHARMA LTD	50MG/ML	A212890	001	Jan 13, 2020
AB		APPCO	50MG/ML	A216317	001	Apr 11, 2023
AB		AUROBINDO PHARMA	50MG/ML	A215124	001	Nov 17, 2022
AB		GRANULES	50MG/ML	A213306	001	Jan 31, 2020
AB		HETERO LABS LTD V	50MG/ML	A211475	001	Oct 04, 2022
AB		MSN	50MG/ML	A210169	001	Feb 17, 2022

TABLET; ORAL

VALCYTE

AB	+!	CHEPLAPHARM	EQ 450MG BASE	N021304	001	Mar 29, 2001
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VALGANCICLOVIR HYDROCHLORIDE

AB		AJANTA PHARMA LTD	EQ 450MG BASE	A212234	001	Dec 26, 2019
AB		AUROBINDO PHARMA LTD	EQ 450MG BASE	A204750	001	Mar 31, 2016
AB		DR REDDYS	EQ 450MG BASE	A203511	001	Nov 04, 2014
AB			EQ 450MG BASE	A206876	001	Dec 12, 2017
AB		HETERO LABS LTD V	EQ 450MG BASE	A205166	001	Mar 18, 2016
AB		STRIDES PHARMA	EQ 450MG BASE	A200790	001	Nov 04, 2014

VALPROATE SODIUM

INJECTABLE; INJECTION

VALPROATE SODIUM

AP		FRESENIUS KABI USA	EQ 100MG BASE/ML	A076539	001	Jun 26, 2003
AP	!	HIKMA FARMACEUTICA	EQ 100MG BASE/ML	A078523	001	Feb 17, 2010
AP		SAGENT	EQ 100MG BASE/ML	A076295	001	Nov 14, 2002

VALPROIC ACID

CAPSULE; ORAL

VALPROIC ACID

AB	!	BIONPHARMA	250MG	A073484	001	Jun 29, 1993
AB		CATALENT	250MG	A073229	001	Oct 29, 1991
AB		SUN PHARM INDS LTD	250MG	A091037	001	Feb 22, 2013
BX		EYWA	250MG	A207611	001	Aug 05, 2019

SYRUP; ORAL

VALPROIC ACID

AA		ANI PHARMS	250MG/5ML	A073178	001	Aug 25, 1992
AA		CHARTWELL RX	250MG/5ML	A075782	001	Dec 22, 2000
AA		HIKMA	250MG/5ML	A074060	001	Jan 13, 1995
AA	!	PHARM ASSOC	250MG/5ML	A075379	001	Dec 15, 2000
AA		SCIEGEN PHARMS INC	250MG/5ML	A090517	001	May 28, 2010

VALRUBICIN

SOLUTION; INTRAVESICAL

VALRUBICIN

AO		HIKMA	40MG/ML	A206430	001	Apr 19, 2019
AO	+!	ENDO PHARM	40MG/ML	N020892	001	Sep 25, 1998

VALSARTAN

SOLUTION; ORAL

VALSARTAN

!	NOVITIUM PHARMA	20MG/5ML	A214102	001	Nov 02, 2021
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TABLET; ORAL

DIOVAN

AB	+	NOVARTIS	40MG	N021283	004	Aug 14, 2002
AB	+		80MG	N021283	001	Jul 18, 2001
AB	+		160MG	N021283	002	Jul 18, 2001
AB	+!		320MG	N021283	003	Jul 18, 2001

VALSARTAN

AB		ALEMBIC	40MG	A091367	001	Jan 05, 2015
AB			80MG	A091367	002	Jan 05, 2015
AB			160MG	A091367	003	Jan 05, 2015
AB			320MG	A091367	004	Jan 05, 2015
AB		ALKEM LABS LTD	40MG	A205536	001	Mar 12, 2019
AB			80MG	A205536	002	Mar 12, 2019
AB			160MG	A205536	003	Mar 12, 2019
AB			320MG	A205536	004	Mar 12, 2019
AB		AMNEAL PHARMS	40MG	A204011	001	Jan 11, 2016
AB			80MG	A204011	002	Jan 11, 2016
AB			160MG	A204011	003	Jan 11, 2016

PRESCRIPTION DRUG PRODUCT LIST

VALSARTAN

TABLET; ORAL

VALSARTAN

<u>AB</u>		<u>320MG</u>	<u>A204011</u>	<u>004</u>	Jan 11, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>40MG</u>	<u>A202223</u>	<u>001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A202223</u>	<u>002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A202223</u>	<u>003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A202223</u>	<u>004</u>	Jan 05, 2015
<u>AB</u>	DR REDDYS	<u>40MG</u>	<u>A201618</u>	<u>001</u>	Oct 05, 2021
<u>AB</u>		<u>80MG</u>	<u>A201618</u>	<u>002</u>	Oct 05, 2021
<u>AB</u>		<u>160MG</u>	<u>A201618</u>	<u>003</u>	Oct 05, 2021
<u>AB</u>		<u>320MG</u>	<u>A201618</u>	<u>004</u>	Oct 05, 2021
<u>AB</u>	HETERO LABS LTD V	<u>40MG</u>	<u>A203311</u>	<u>001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A203311</u>	<u>002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A203311</u>	<u>003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A203311</u>	<u>004</u>	Jan 05, 2015
<u>AB</u>	JUBILANT GENERICS	<u>40MG</u>	<u>A203536</u>	<u>001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A203536</u>	<u>002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A203536</u>	<u>003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A203536</u>	<u>004</u>	Jan 05, 2015
<u>AB</u>	LUPIN LTD	<u>40MG</u>	<u>A201677</u>	<u>001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A201677</u>	<u>002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A201677</u>	<u>003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A201677</u>	<u>004</u>	Jan 05, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>40MG</u>	<u>A202696</u>	<u>001</u>	Sep 16, 2016
<u>AB</u>		<u>80MG</u>	<u>A202696</u>	<u>002</u>	Sep 16, 2016
<u>AB</u>		<u>160MG</u>	<u>A202696</u>	<u>003</u>	Sep 16, 2016
<u>AB</u>		<u>320MG</u>	<u>A202696</u>	<u>004</u>	Sep 16, 2016
<u>AB</u>	MYLAN	<u>40MG</u>	<u>A090866</u>	<u>001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A090866</u>	<u>002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A090866</u>	<u>003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A090866</u>	<u>004</u>	Jan 05, 2015
<u>AB</u>	OHM LABS INC	<u>40MG</u>	<u>A077492</u>	<u>001</u>	Jun 26, 2014
<u>AB</u>		<u>80MG</u>	<u>A077492</u>	<u>002</u>	Jun 26, 2014
<u>AB</u>		<u>160MG</u>	<u>A077492</u>	<u>003</u>	Jun 26, 2014
<u>AB</u>		<u>320MG</u>	<u>A077492</u>	<u>004</u>	Jun 26, 2014
<u>AB</u>	PRINSTON INC	<u>40MG</u>	<u>A204821</u>	<u>001</u>	Jun 09, 2015
<u>AB</u>		<u>80MG</u>	<u>A204821</u>	<u>002</u>	Jun 09, 2015
<u>AB</u>		<u>160MG</u>	<u>A204821</u>	<u>003</u>	Jun 09, 2015
<u>AB</u>		<u>320MG</u>	<u>A204821</u>	<u>004</u>	Jun 09, 2015
<u>AB</u>	SCIEGEN PHARMS INC	<u>40MG</u>	<u>A204038</u>	<u>001</u>	Oct 27, 2021
<u>AB</u>		<u>80MG</u>	<u>A204038</u>	<u>002</u>	Oct 27, 2021
<u>AB</u>		<u>160MG</u>	<u>A204038</u>	<u>003</u>	Oct 27, 2021
<u>AB</u>		<u>320MG</u>	<u>A204038</u>	<u>004</u>	Oct 27, 2021
<u>AB</u>	SQUARE PHARMS	<u>40MG</u>	<u>A205347</u>	<u>001</u>	Apr 09, 2018
<u>AB</u>		<u>80MG</u>	<u>A205347</u>	<u>002</u>	Apr 09, 2018
<u>AB</u>		<u>160MG</u>	<u>A205347</u>	<u>003</u>	Apr 09, 2018
<u>AB</u>		<u>320MG</u>	<u>A205347</u>	<u>004</u>	Apr 09, 2018

VAMOROLONE

SUSPENSION; ORAL

AGAMREE

+! CATALYST PHARMS 40MG/ML N215239 001 Oct 26, 2023

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOCIN HYDROCHLORIDE

<u>AB</u>	+ ANI PHARMS	<u>EQ 125MG BASE</u>	<u>N050606</u>	<u>001</u>	Apr 15, 1986
<u>AB</u>	+!	<u>EQ 250MG BASE</u>	<u>N050606</u>	<u>002</u>	Apr 15, 1986
	<u>VANCOMYCIN HYDROCHLORIDE</u>				
<u>AB</u>	LUPIN LTD	<u>EQ 125MG BASE</u>	<u>A090439</u>	<u>001</u>	Jan 28, 2015
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A090439</u>	<u>002</u>	Jan 28, 2015
<u>AB</u>	ORIENT PHARMA CO LTD	<u>EQ 125MG BASE</u>	<u>A210729</u>	<u>001</u>	Apr 29, 2019
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A210729</u>	<u>002</u>	Apr 29, 2019
<u>AB</u>	PAI HOLDINGS PHARM	<u>EQ 125MG BASE</u>	<u>A065478</u>	<u>001</u>	Apr 09, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065478</u>	<u>002</u>	Apr 09, 2012
<u>AB</u>	STRIDES PHARMA	<u>EQ 125MG BASE</u>	<u>A065490</u>	<u>001</u>	Apr 09, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065490</u>	<u>002</u>	Apr 09, 2012

FOR SOLUTION; ORAL

VANCOCIN HYDROCHLORIDEAA ! ANI PHARMS EQ 250MG BASE/5ML A061667 002 Jul 13, 1983

PRESCRIPTION DRUG PRODUCT LIST

VANCOMYCIN HYDROCHLORIDE

FOR SOLUTION; ORAL

VANCOMYCIN HYDROCHLORIDE

<u>AA</u>	AMNEAL	<u>EQ 250MG BASE/5ML</u>	<u>A215338 001</u>	Jun 23, 2023
	<u>FIRVANQ KIT</u>			
<u>AB</u>	+! AZURITY	<u>EQ 25MG BASE/ML</u>	<u>N208910 001</u>	Jan 26, 2018
<u>AB</u>	+!	<u>EQ 50MG BASE/ML</u>	<u>N208910 002</u>	Jan 26, 2018
	<u>VANCOMYCIN HYDROCHLORIDE</u>			
<u>AB</u>	ALKEM LABS LTD	<u>EQ 25MG BASE/ML</u>	<u>A214913 001</u>	Nov 14, 2022
<u>AB</u>		<u>EQ 50MG BASE/ML</u>	<u>A214913 002</u>	Nov 14, 2022

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	ASPIRO	<u>EQ 500MG BASE/VIAL</u>	<u>A216591 001</u>	Jul 06, 2022
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A216591 002</u>	Jul 06, 2022
<u>AP</u>	EUGIA PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A205780 001</u>	Mar 31, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A205780 002</u>	Mar 31, 2016
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A205779 001</u>	Mar 29, 2016
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A205779 002</u>	Mar 29, 2016
<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A062663 001</u>	Mar 17, 1987
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062663 005</u>	Aug 17, 2016
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062663 002</u>	Jul 31, 1987
<u>AP</u>	!	<u>EQ 5GM BASE/VIAL</u>	<u>A062663 003</u>	Jun 03, 1988
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A062663 004</u>	Nov 28, 1997
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A205694 001</u>	Jan 21, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A205694 002</u>	Jan 21, 2016
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A205694 003</u>	Jun 06, 2023
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A205694 004</u>	Jun 06, 2023
<u>AP</u>	HAINAN POLY	<u>EQ 5GM BASE/VIAL</u>	<u>A215821 001</u>	Nov 18, 2021
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A215821 002</u>	Nov 18, 2021
<u>AP</u>	HAINAN POLY PHARM	<u>EQ 500MG BASE/VIAL</u>	<u>A212332 001</u>	Jun 12, 2019
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A212332 002</u>	Jun 12, 2019
<u>AP</u>	HIKMA	<u>EQ 1GM BASE/VIAL</u>	<u>A203300 002</u>	Aug 11, 2020
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A204360 001</u>	Oct 15, 2018
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A204360 002</u>	Oct 15, 2018
<u>AP</u>	HIKMA PHARMS	<u>EQ 750MG BASE/VIAL</u>	<u>A206616 001</u>	Oct 03, 2018
<u>AP</u>	! HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062911 001</u>	Aug 04, 1988
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A062931 001</u>	Oct 29, 1992
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062912 002</u>	Jan 07, 2009
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062933 002</u>	May 27, 2009
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062912 001</u>	Aug 04, 1988
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062933 001</u>	Oct 29, 1992
<u>AP</u>	!	<u>EQ 5GM BASE/VIAL</u>	<u>A063076 001</u>	Dec 21, 1990
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A063076 002</u>	Sep 14, 2020
<u>AP</u>	HOSPIRA INC	<u>EQ 10GM BASE/VIAL</u>	<u>A065455 001</u>	Apr 29, 2009
<u>AP</u>	MEITHEAL	<u>EQ 500MG BASE/VIAL</u>	<u>A215197 001</u>	Jul 28, 2022
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A215195 001</u>	Sep 15, 2022
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A215197 002</u>	Jul 28, 2022
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A215196 001</u>	Jul 27, 2022
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A215196 002</u>	Jul 27, 2022
<u>AP</u>	MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065397 001</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065397 002</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A065432 001</u>	Dec 30, 2008
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A091554 001</u>	Sep 19, 2011
<u>AP</u>	PHARM ASSOC	<u>EQ 500MG BASE/VIAL</u>	<u>A065401 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065401 002</u>	Jun 30, 2008
<u>AP</u>	SAGENT PHARMS	<u>EQ 5GM BASE/VIAL</u>	<u>A200837 001</u>	Aug 10, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A200837 002</u>	Sep 02, 2014
<u>AP</u>	SANDOZ INC	<u>EQ 5GM BASE/VIAL</u>	<u>A201048 001</u>	Aug 10, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A201048 002</u>	Aug 10, 2012
<u>AP</u>	XELLIA PHARMS APS	<u>EQ 5GM BASE/VIAL</u>	<u>A204125 001</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A204125 002</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A204107 001</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A204107 002</u>	Dec 28, 2015
	VANCOXIN HYDROCHLORIDE IN PLASTIC CONTAINER			
	+! BAXTER HLTHCARE	EQ 500MG BASE/100ML	N050671 001	Apr 29, 1993
	+!	EQ 750MG BASE/150ML	N050671 002	Dec 20, 2010
	+!	EQ 1GM BASE/200ML	N050671 003	Mar 01, 1999
	VANCOMYCIN HYDROCHLORIDE			
	! HOSPIRA	EQ 1.5GM BASE/VIAL	A062912 003	Jul 10, 2020
	POWDER; INTRAVENOUS			
	<u>VANCOMYCIN HYDROCHLORIDE</u>			
<u>AP</u>	EUGIA PHARMA	<u>EQ 1.25GM BASE/VIAL</u>	<u>A217401 001</u>	Aug 04, 2023

PRESCRIPTION DRUG PRODUCT LIST

VANCOMYCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

AP		<u>EQ 1.5GM BASE/VIAL</u>	<u>A217401 002</u>	Aug 04, 2023
AP	HIKMA	<u>EQ 1.25GM BASE/VIAL</u>	<u>A217489 001</u>	Jun 29, 2023
AP		<u>EQ 1.5GM BASE/VIAL</u>	<u>A217489 002</u>	Jun 29, 2023
AP	+! MYLAN LABS LTD	<u>EQ 1.25GM BASE/VIAL</u>	<u>N209481 003</u>	Jul 10, 2018
AP	+!	<u>EQ 1.5GM BASE/VIAL</u>	<u>N209481 004</u>	Jul 10, 2018
	+!	EQ 750MG BASE/VIAL	N209481 002	Jul 10, 2018
	VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER			
	SAMSON MEDCL	EQ 100GM BASE	A091532 001	Jan 06, 2016
	POWDER; INTRAVENOUS, ORAL			
	VANCOMYCIN HYDROCHLORIDE			
	+! ZHEJIANG NOVUS PHARM	EQ 500MG BASE/VIAL	N210274 001	Jan 20, 2023
	+!	EQ 1GM BASE/VIAL	N210274 002	Jan 20, 2023
	+!	EQ 5GM BASE/VIAL	N210274 003	Jan 20, 2023
	+!	EQ 10GM BASE/VIAL	N210274 004	Jan 20, 2023
	SOLUTION; INTRAVENOUS			
	VANCOMYCIN HYDROCHLORIDE			
	+! XELLIA PHARMS APS	EQ 500MG BASE/100ML (EQ 5MG BASE/ML)	N211962 001	Feb 15, 2019
	+!	EQ 750MG BASE/150ML (EQ 5MG BASE/ML)	N211962 005	May 13, 2020
	+!	EQ 1GM BASE/200ML (EQ 5MG BASE/ML)	N211962 002	Feb 15, 2019
	+!	EQ 1.25GM BASE/250ML (EQ 5MG BASE/ML)	N211962 006	May 13, 2020
	+!	EQ 1.5GM BASE/300ML (EQ 5MG BASE/ML)	N211962 003	Feb 15, 2019
	+!	EQ 1.75GM BASE/350ML (EQ 5MG BASE/ML)	N211962 007	May 13, 2020
	+!	EQ 2GM BASE/400ML (EQ 5MG BASE/ML)	N211962 004	Feb 15, 2019

VANDETANIB

TABLET; ORAL

CAPRELSA

	+ GENZYME CORP	100MG	N022405 001	Apr 06, 2011
	+!	300MG	N022405 002	Apr 06, 2011

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

VARDENAFIL HYDROCHLORIDE

AB	ALEMBIC	<u>EQ 2.5MG BASE</u>	<u>A214031 001</u>	Aug 04, 2020
AB		<u>EQ 5MG BASE</u>	<u>A214031 002</u>	Aug 04, 2020
AB		<u>EQ 10MG BASE</u>	<u>A214031 003</u>	Aug 04, 2020
AB		<u>EQ 20MG BASE</u>	<u>A214031 004</u>	Aug 04, 2020
AB	CROSSMEDIKA SA	<u>EQ 2.5MG BASE</u>	<u>A209057 001</u>	Oct 31, 2018
AB		<u>EQ 5MG BASE</u>	<u>A209057 002</u>	Oct 31, 2018
AB		<u>EQ 10MG BASE</u>	<u>A209057 003</u>	Oct 31, 2018
AB		<u>EQ 20MG BASE</u>	<u>A209057 004</u>	Oct 31, 2018
AB	MACLEODS PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A204632 001</u>	Oct 22, 2019
AB		<u>EQ 5MG BASE</u>	<u>A204632 002</u>	Oct 22, 2019
AB		<u>EQ 10MG BASE</u>	<u>A204632 003</u>	Oct 22, 2019
AB		<u>EQ 20MG BASE</u>	<u>A204632 004</u>	Oct 22, 2019
AB	TEVA PHARMS	<u>EQ 2.5MG BASE</u>	<u>A091347 001</u>	May 03, 2012
AB		<u>EQ 5MG BASE</u>	<u>A091347 002</u>	May 03, 2012
AB		<u>EQ 10MG BASE</u>	<u>A091347 003</u>	May 03, 2012
AB		<u>EQ 20MG BASE</u>	<u>A091347 004</u>	May 03, 2012
AB	ZYDUS PHARMS	<u>EQ 2.5MG BASE</u>	<u>A208960 001</u>	Oct 31, 2018
AB		<u>EQ 5MG BASE</u>	<u>A208960 002</u>	Oct 31, 2018
AB		<u>EQ 10MG BASE</u>	<u>A208960 003</u>	Oct 31, 2018
AB	!	<u>EQ 20MG BASE</u>	<u>A208960 004</u>	Oct 31, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

VARDENAFIL HYDROCHLORIDE

AB	!	ALEMBIC	<u>10MG</u>	<u>A208324 001</u>	Nov 16, 2018
AB		MACLEODS PHARMS LTD	<u>10MG</u>	<u>A205988 001</u>	Mar 10, 2020

VARENICLINE TARTRATE

SPRAY; NASAL

TYRVAYA

	+! OYSTER POINT PHARMA	EQ 0.03MG BASE/SPRAY	N213978 001	Oct 15, 2021
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TABLET; ORAL

VARENICLINE TARTRATE

AB	ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A214557 001</u>	Aug 23, 2023
AB		<u>EQ 1MG BASE</u>	<u>A214557 002</u>	Aug 23, 2023
AB	APOTEX	<u>EQ 0.5MG BASE</u>	<u>A201962 001</u>	Jan 25, 2023
AB		<u>EQ 1MG BASE</u>	<u>A201962 002</u>	Jan 25, 2023
AB	HETERO LABS LTD III	<u>EQ 0.5MG BASE</u>	<u>A214571 001</u>	Oct 23, 2023
AB		<u>EQ 1MG BASE</u>	<u>A214571 002</u>	Oct 23, 2023

PRESCRIPTION DRUG PRODUCT LIST

VARENICLINE TARTRATE

TABLET; ORAL

VARENICLINE TARTRATE

<u>AB</u>	LEADING PHARMA	<u>EQ 0.5MG BASE</u>	<u>A217151 001</u>	Jul 25, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A217151 002</u>	Jul 25, 2023
<u>AB</u>	LUPIN LTD	<u>EQ 0.5MG BASE</u>	<u>A211862 001</u>	Dec 04, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A211862 002</u>	Dec 04, 2023
<u>AB</u>	MANKIND PHARMA	<u>EQ 0.5MG BASE</u>	<u>A214255 001</u>	Aug 01, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A214255 002</u>	Aug 01, 2023
<u>AB</u>	PAR PHARM INC	<u>EQ 0.5MG BASE</u>	<u>A201785 001</u>	Aug 11, 2021
<u>AB</u>	!	<u>EQ 1MG BASE</u>	<u>A201785 002</u>	Aug 11, 2021
<u>AB</u>	ZYDUS	<u>EQ 0.5MG BASE</u>	<u>A216723 001</u>	Jun 12, 2023
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A216723 002</u>	Jun 12, 2023

VASOPRESSIN

SOLUTION; INTRAVENOUS

VASOPRESSIN

<u>AP</u>	AMNEAL	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A212944 001</u>	Aug 05, 2022
<u>AP</u>	AMPHASTAR PHARMS INC	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A211857 001</u>	Jul 18, 2022
<u>AP</u>	CIPLA	<u>20UNITS/100ML (0.2UNITS/ML)</u>	<u>A217987 001</u>	Dec 06, 2023
<u>AP</u>	EAGLE PHARMS	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A211538 001</u>	Dec 15, 2021
<u>AP</u>	EUGIA PHARMA	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A214314 001</u>	Aug 15, 2022
<u>AP</u>	FRESENIUS KABI USA	<u>20UNITS/ML (20UNITS/ML)</u>	<u>A213206 001</u>	May 26, 2023

VASOSTRICT

<u>AP</u>	+!	PAR STERILE PRODUCTS	<u>20UNITS/ML (20UNITS/ML)</u>	<u>N204485 001</u>	Apr 17, 2014
<u>AP</u>	+!		<u>20UNITS/100ML (0.2UNITS/ML)</u>	<u>N204485 005</u>	Apr 21, 2021
		VASOPRESSIN			
	+!	AM REGENT	20UNITS/ML (20UNITS/ML)	N212593 001	Aug 03, 2020
	+!		200UNITS/10ML (20UNITS/ML)	N212593 002	Jun 09, 2023
		VASOPRESSIN IN SODIUM CHLORIDE 0.9%			
	+!	BAXTER HLTHCARE CORP	20UNITS/100ML (0.2UNITS/ML)	N217569 001	Sep 29, 2023
	+!		40UNITS/100ML (0.4UNITS/ML)	N217569 002	Sep 29, 2023
		VASOSTRICT			
	+!	PAR STERILE PRODUCTS	40UNITS/100ML (0.4UNITS/ML)	N204485 003	Apr 15, 2020
	+!		200UNITS/10ML (20UNITS/ML)	N204485 002	Dec 17, 2016

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>	EUGIA PHARMA	<u>10MG/VIAL</u>	<u>A206670 001</u>	Dec 20, 2018	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A206670 002</u>	Dec 20, 2018	
<u>AP</u>	GLAND	<u>10MG/VIAL</u>	<u>A205390 001</u>	May 26, 2016	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A205390 002</u>	May 26, 2016	
<u>AP</u>	HIKMA	<u>10MG/VIAL</u>	<u>A075549 001</u>	Jun 13, 2000	
<u>AP</u>		<u>10MG/VIAL</u>	<u>A203725 001</u>	Jul 30, 2019	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A075549 002</u>	Jun 13, 2000	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A203725 002</u>	Jul 30, 2019	
<u>AP</u>	HOSPIRA	<u>10MG/VIAL</u>	<u>A075164 001</u>	Oct 21, 1999	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A075164 002</u>	Oct 21, 1999	
<u>AP</u>	MEITHEAL	<u>10MG/VIAL</u>	<u>A074688 001</u>	Aug 25, 1999	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A074688 002</u>	Aug 25, 1999	
<u>AP</u>	MYLAN LABS LTD	<u>10MG/VIAL</u>	<u>A090243 001</u>	May 11, 2010	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A090243 002</u>	May 11, 2010	
<u>AP</u>	SAGENT PHARMS INC	<u>10MG/VIAL</u>	<u>A078274 001</u>	Dec 29, 2008	
<u>AP</u>		<u>20MG/VIAL</u>	<u>A078274 002</u>	Dec 29, 2008	
<u>AP</u>	!	SUN PHARM	<u>10MG/VIAL</u>	<u>A079001 001</u>	Jun 17, 2009
<u>AP</u>	!		<u>20MG/VIAL</u>	<u>A079001 002</u>	Jun 17, 2009

VEMURAFENIB

TABLET; ORAL

ZELBORAF

+!	HOFFMANN LA ROCHE	240MG	N202429 001	Aug 17, 2011
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VENETOCLAX

TABLET; ORAL

VENCLEXTA

+	ABBVIE	10MG	N208573 001	Apr 11, 2016
+		50MG	N208573 002	Apr 11, 2016
+!		100MG	N208573 003	Apr 11, 2016

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE BESYLATE

TABLET, EXTENDED RELEASE;ORAL

VENLAFAXINE BESYLATE

+! ALMATICA

EQ 112.5MG BASE

N215429 001 Jun 29, 2022

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

EFFEXOR XRAB + UPJOHNEQ 37.5MG BASEN020699 001 Oct 20, 1997AB +EQ 75MG BASEN020699 002 Oct 20, 1997AB +!EQ 150MG BASEN020699 004 Oct 20, 1997VENLAFAXINE HYDROCHLORIDEAB ANNORA PHARMAEQ 37.5MG BASEA212277 001 Jul 08, 2019ABEQ 75MG BASEA212277 002 Jul 08, 2019ABEQ 150MG BASEA212277 003 Jul 08, 2019AB AUROBINDO PHARMA
LTDEQ 37.5MG BASEA200834 001 Apr 14, 2011ABEQ 75MG BASEA200834 002 Apr 14, 2011ABEQ 150MG BASEA200834 003 Apr 14, 2011AB DR REDDYS LABS LTDEQ 37.5MG BASEA078421 001 May 06, 2011ABEQ 75MG BASEA078421 002 May 06, 2011ABEQ 150MG BASEA078421 003 May 06, 2011AB GRANULESEQ 37.5MG BASEA217390 001 May 18, 2023ABEQ 75MG BASEA217390 002 May 18, 2023ABEQ 150MG BASEA217390 003 May 18, 2023AB INTELLIPHARMACEUTIC
SEQ 37.5MG BASEA201272 001 Nov 23, 2018ABEQ 75MG BASEA201272 002 Nov 23, 2018ABEQ 150MG BASEA201272 003 Nov 23, 2018AB INVENTIA HLTHCAREEQ 37.5MG BASEA203332 001 Mar 12, 2020ABEQ 75MG BASEA203332 002 Mar 12, 2020ABEQ 150MG BASEA203332 003 Mar 12, 2020AB MACLEODS PHARMS LTDEQ 37.5MG BASEA204889 001 Oct 05, 2017ABEQ 75MG BASEA204889 002 Oct 05, 2017ABEQ 150MG BASEA204889 003 Oct 05, 2017AB NOSTRUM PHARMS LLCEQ 37.5MG BASEA200430 001 Apr 04, 2023ABEQ 75MG BASEA200430 002 Apr 04, 2023ABEQ 150MG BASEA200430 003 Apr 04, 2023AB ORBION PHARMSEQ 37.5MG BASEA091123 001 Jul 11, 2011ABEQ 75MG BASEA091123 002 Jul 11, 2011ABEQ 150MG BASEA091123 003 Jul 11, 2011AB TEVAEQ 37.5MG BASEA076565 001 Jun 28, 2010ABEQ 75MG BASEA076565 002 Jun 28, 2010ABEQ 150MG BASEA076565 003 Jun 28, 2010AB VALEANT PHARMS
NORTHEQ 37.5MG BASEA090071 001 Apr 15, 2011ABEQ 75MG BASEA090071 002 Apr 15, 2011ABEQ 150MG BASEA090071 003 Apr 15, 2011AB WOCKHARDT BIO AGEQ 37.5MG BASEA078865 001 Apr 14, 2011ABEQ 75MG BASEA078865 002 Apr 14, 2011ABEQ 150MG BASEA078865 003 Apr 14, 2011AB YICHANG HUMANWELLEQ 37.5MG BASEA214654 001 Aug 06, 2021ABEQ 75MG BASEA214654 002 Aug 06, 2021ABEQ 150MG BASEA214654 003 Aug 06, 2021AB ZYDUS PHARMS USA
INCEQ 37.5MG BASEA090174 001 Apr 14, 2011ABEQ 75MG BASEA090174 002 Apr 14, 2011ABEQ 150MG BASEA090174 003 Apr 14, 2011

TABLET;ORAL

VENLAFAXINE HYDROCHLORIDEAB ALEMbic PHARMS LTDEQ 25MG BASEA078932 001 Dec 14, 2010ABEQ 37.5MG BASEA078932 002 Dec 14, 2010ABEQ 50MG BASEA078932 003 Dec 14, 2010ABEQ 75MG BASEA078932 004 Dec 14, 2010ABEQ 100MG BASEA078932 005 Dec 14, 2010AB AMNEAL PHARMSEQ 25MG BASEA079098 001 May 11, 2010ABEQ 37.5MG BASEA079098 002 May 11, 2010ABEQ 50MG BASEA079098 003 May 11, 2010ABEQ 75MG BASEA079098 004 May 11, 2010ABEQ 100MG BASEA079098 005 May 11, 2010AB AUROBINDO PHARMAEQ 25MG BASEA090555 001 Apr 07, 2010ABEQ 37.5MG BASEA090555 002 Apr 07, 2010ABEQ 50MG BASEA090555 003 Apr 07, 2010ABEQ 75MG BASEA090555 004 Apr 07, 2010

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

TABLET;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090555 005</u>	Apr 07, 2010
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A206250 001</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A206250 002</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A206250 003</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A206250 004</u>	Nov 21, 2018
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A206250 005</u>	Nov 21, 2018
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A078301 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078301 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078301 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078301 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078301 005</u>	Jun 13, 2008
<u>AB</u>	HERITAGE PHARMS	<u>EQ 25MG BASE</u>	<u>A078554 001</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078554 002</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078554 003</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078554 004</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078554 005</u>	Jan 09, 2009
<u>AB</u>	SUN PHARM INDS INC	<u>EQ 25MG BASE</u>	<u>A078627 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078627 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078627 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078627 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078627 005</u>	Jun 13, 2008
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A076690 001</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A076690 002</u>	Aug 03, 2006
<u>AB</u>	!	<u>EQ 50MG BASE</u>	<u>A076690 003</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A076690 004</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076690 005</u>	Aug 03, 2006
<u>AB</u>	YAOPHARMA CO LTD	<u>EQ 25MG BASE</u>	<u>A202036 001</u>	May 28, 2015
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A202036 002</u>	May 28, 2015
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202036 003</u>	May 28, 2015
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A202036 004</u>	May 28, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202036 005</u>	May 28, 2015
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 25MG BASE</u>	<u>A077653 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A077653 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077653 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A077653 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077653 005</u>	Jun 13, 2008

TABLET, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 37.5MG BASE</u>	<u>A214691 001</u>	Apr 12, 2023
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214691 002</u>	Apr 12, 2023
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214691 003</u>	Apr 12, 2023
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214691 004</u>	Apr 12, 2023
<u>AB</u>	ALKEM LABS LTD	<u>EQ 37.5MG BASE</u>	<u>A214127 001</u>	Jun 10, 2021
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214127 002</u>	Jun 10, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214127 003</u>	Jun 10, 2021
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214127 004</u>	Jun 10, 2021
<u>AB</u>	APPCO	<u>EQ 150MG BASE</u>	<u>A214609 001</u>	Jun 30, 2021
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214609 002</u>	Jun 30, 2021
<u>AB</u>	ASCENT PHARMS INC	<u>EQ 37.5MG BASE</u>	<u>A214419 001</u>	Oct 21, 2020
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A214419 002</u>	Oct 21, 2020
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A214419 003</u>	Oct 21, 2020
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A214419 004</u>	Oct 21, 2020
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A211323 001</u>	Aug 29, 2019
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A211323 002</u>	Aug 29, 2019
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A211323 003</u>	Aug 29, 2019
<u>AB</u>	DEXCEL	<u>EQ 75MG BASE</u>	<u>A213927 001</u>	Jan 21, 2021
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209193 001</u>	Oct 31, 2019
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A209193 002</u>	Oct 31, 2019
<u>AB</u>	NOSTRUM LABS INC	<u>EQ 150MG BASE</u>	<u>A205468 002</u>	Mar 24, 2017
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A205468 003</u>	Mar 24, 2017
<u>AB</u>	+	<u>EQ 37.5MG BASE</u>	<u>N022104 001</u>	May 20, 2008
<u>AB</u>	+	<u>EQ 75MG BASE</u>	<u>N022104 002</u>	May 20, 2008
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N022104 003</u>	May 20, 2008
<u>AB</u>	+	<u>EQ 225MG BASE</u>	<u>N022104 004</u>	May 20, 2008
<u>AB</u>	SUN PHARM	<u>EQ 37.5MG BASE</u>	<u>A091272 001</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A091272 002</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091272 003</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A091272 004</u>	Jan 08, 2019
<u>AB</u>	UNIQUE	<u>EQ 37.5MG BASE</u>	<u>A216044 001</u>	Nov 28, 2022

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A216044 002</u>	Nov 28, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A216044 003</u>	Nov 28, 2022
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A216044 004</u>	Nov 28, 2022
<u>AB</u>	ZYDUS PHARMS	<u>EQ 37.5MG BASE</u>	<u>A215622 001</u>	Aug 30, 2022
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A215622 002</u>	Aug 30, 2022
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A215622 003</u>	Aug 30, 2022
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A215622 004</u>	Aug 30, 2022

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>120MG</u>	<u>A075138 001</u>	Apr 20, 1999
<u>AB</u>		<u>180MG</u>	<u>A075138 002</u>	Apr 20, 1999
<u>AB</u>		<u>240MG</u>	<u>A075138 003</u>	Apr 20, 1999
<u>VERELAN</u>				
<u>AB</u>	+ SOCIETAL CDMO	<u>120MG</u>	<u>N019614 001</u>	May 29, 1990
<u>AB</u>	+	<u>180MG</u>	<u>N019614 003</u>	Jan 09, 1992
<u>AB</u>	+	<u>240MG</u>	<u>N019614 002</u>	May 29, 1990
	+	360MG	N019614 004	May 10, 1996
VERELAN PM				
	+ SOCIETAL CDMO	100MG	N020943 001	Nov 25, 1998
	+	200MG	N020943 002	Nov 25, 1998
	+	300MG	N020943 003	Nov 25, 1998

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	ACIC PHARMS	<u>2.5MG/ML</u>	<u>A215829 001</u>	Mar 17, 2022
<u>AP</u>	! EUGIA PHARMA	<u>2.5MG/ML</u>	<u>A212965 001</u>	Jul 06, 2020
<u>AP</u>	GLAND PHARMA LTD	<u>2.5MG/ML</u>	<u>A214361 001</u>	Oct 15, 2020
<u>AP</u>	MANKIND PHARMA	<u>2.5MG/ML</u>	<u>A214653 001</u>	Jun 15, 2022
<u>AP</u>	ZYDUS PHARMS	<u>2.5MG/ML</u>	<u>A214215 001</u>	Oct 15, 2020

SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	AMNEAL	<u>5MG/2ML (2.5MG/ML)</u>	<u>A210994 001</u>	Jul 13, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A210994 002</u>	Jul 13, 2018
<u>AP</u>	AREVA PHARMS	<u>5MG/2ML (2.5MG/ML)</u>	<u>A213352 002</u>	Sep 16, 2020
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A213352 001</u>	Mar 17, 2020
<u>AP</u>	CAPLIN	<u>5MG/2ML (2.5MG/ML)</u>	<u>A213232 001</u>	Mar 25, 2020
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A213232 002</u>	Mar 25, 2020
<u>AP</u>	EXELA PHARMA	<u>5MG/2ML (2.5MG/ML)</u>	<u>N018925 001</u>	Mar 30, 1984
<u>AP</u>	! HOSPIRA	<u>5MG/2ML (2.5MG/ML)</u>	<u>A070738 001</u>	May 06, 1987
<u>AP</u>	!	<u>5MG/2ML (2.5MG/ML)</u>	<u>A075136 001</u>	Oct 20, 1998
<u>AP</u>	!	<u>5MG/2ML (2.5MG/ML)</u>	<u>A070737 001</u>	May 06, 1987
<u>AP</u>	!	<u>10MG/4ML (2.5MG/ML)</u>	<u>A070737 002</u>	May 06, 1987
<u>AP</u>	MICRO LABS	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211370 001</u>	Dec 28, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211370 002</u>	Dec 28, 2018
<u>AP</u>	SOMERSET	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211035 001</u>	Jun 18, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211035 002</u>	Jun 18, 2018
<u>AP</u>	SOMERSET THERAPS LLC	<u>5MG/2ML (2.5MG/ML)</u>	<u>A211015 001</u>	Jun 18, 2018
<u>AP</u>		<u>10MG/4ML (2.5MG/ML)</u>	<u>A211015 002</u>	Jun 18, 2018

TABLET; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	HERITAGE PHARMS	<u>40MG</u>	<u>A071881 002</u>	Oct 14, 2015
<u>AB</u>		<u>80MG</u>	<u>A071881 003</u>	Apr 05, 1988
<u>AB</u>	!	<u>120MG</u>	<u>A071881 001</u>	Apr 05, 1988
<u>AB</u>	WATSON LABS	<u>40MG</u>	<u>A072924 001</u>	Jun 29, 1993
<u>AB</u>		<u>80MG</u>	<u>A070995 001</u>	Oct 01, 1986
<u>AB</u>		<u>120MG</u>	<u>A070994 001</u>	Oct 01, 1986

TABLET, EXTENDED RELEASE;ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	CADILA PHARMS LTD	<u>120MG</u>	<u>A206173 003</u>	Nov 14, 2022
<u>AB</u>		<u>180MG</u>	<u>A206173 001</u>	May 05, 2017
<u>AB</u>		<u>240MG</u>	<u>A206173 002</u>	May 05, 2017
<u>AB</u>	! GLENMARK GENERICS	<u>120MG</u>	<u>A090700 001</u>	Aug 03, 2011
<u>AB</u>	!	<u>180MG</u>	<u>A090700 002</u>	Aug 03, 2011
<u>AB</u>	!	<u>240MG</u>	<u>A078906 001</u>	Sep 17, 2009
<u>AB</u>	STRIDES PHARMA	<u>120MG</u>	<u>A075072 001</u>	May 25, 1999
<u>AB</u>		<u>240MG</u>	<u>A075072 003</u>	May 25, 1999

PRESCRIPTION DRUG PRODUCT LIST

VERICIGUAT

TABLET; ORAL

VERQUVO

+	MERCK SHARP DOHME	2.5MG	N214377	001	Jan 19, 2021
+		5MG	N214377	002	Jan 19, 2021
+	!	10MG	N214377	003	Jan 19, 2021

VERTEPORFIN

INJECTABLE; INJECTION

VISUDYNE

+	!	VALEANT LUXEMBOURG	15MG/VIAL	N021119	001	Apr 12, 2000
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VIBEGRON

TABLET; ORAL

GEMTESA

+	!	UROVANT	75MG	N213006	001	Dec 23, 2020
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VIGABATRIN

FOR SOLUTION; ORAL

SABRIL

AA	+	!	LUNDBECK PHARMS LLC	500MG/PACKET	N022006	001	Aug 21, 2009
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VIGABATRIN

AA			ALKEM LABS LTD	500MG/PACKET	A213375	001	Dec 02, 2020
AA			AMNEAL PHARMS	500MG/PACKET	A210155	001	Mar 13, 2018
AA			ANNORA PHARMA	500MG/PACKET	A213519	001	Jan 26, 2021
AA			AUROBINDO PHARMA LTD	500MG/PACKET	A213899	001	Sep 29, 2021
AA			DEXCEL	500MG/PACKET	A214992	001	May 13, 2021
AA			DR REDDYS	500MG/PACKET	A211481	001	Nov 20, 2018
AA			INVAGEN PHARMS	500MG/PACKET	A211592	001	Dec 03, 2019
AA			MSN	500MG/PACKET	A215363	001	Sep 07, 2022
AA			PAR PHARM INC	500MG/PACKET	A208218	001	Apr 27, 2017
AA			PROPEL PHARMA	500MG/PACKET	A213390	001	Jul 29, 2021
AA			TEVA PHARMS USA	500MG/PACKET	A209824	001	Apr 23, 2018

VIGADRONE

AA			AUCTA	500MG/PACKET	A210196	001	Jun 21, 2018
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VIGPODER

AA			PYROS PHARMS	500MG/PACKET	A214961	001	Jun 24, 2022
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TABLET; ORAL

SABRIL

AB	+	!	LUNDBECK PHARMS LLC	500MG	N020427	001	Aug 21, 2009
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VIGABATRIN

AB			AMNEAL PHARMS	500MG	A210042	001	Jun 22, 2022
AB			AUROBINDO PHARMA LTD	500MG	A215601	001	May 10, 2022
AB			DEXCEL	500MG	A215109	001	Sep 23, 2021
AB			DR REDDYS	500MG	A211539	001	Jan 29, 2021
AB			HIKMA	500MG	A213104	001	Aug 29, 2022
AB			PROPEL PHARMA	500MG	A215519	001	Apr 28, 2023
AB			TEVA PHARMS USA	500MG	A209822	001	Jan 14, 2019
AB			UPSHER SMITH LABS	500MG	A214749	001	Jun 29, 2023
AB			ZYDUS	500MG	A215707	001	Jan 19, 2022

VILAZODONE HYDROCHLORIDE

TABLET; ORAL

VIIBRYD

AB	+	!	ABBVIE	10MG	N022567	001	Jan 21, 2011
AB	+			20MG	N022567	002	Jan 21, 2011
AB	+			40MG	N022567	003	Jan 21, 2011

VILAZODONE HYDROCHLORIDE

AB			ACCORD HLTHCARE	10MG	A208209	001	Apr 27, 2021
AB				20MG	A208209	002	Apr 27, 2021
AB				40MG	A208209	003	Apr 27, 2021
AB			ALEMBIC	10MG	A208202	001	Jan 10, 2020
AB				20MG	A208202	002	Jan 10, 2020
AB				40MG	A208202	003	Jan 10, 2020
AB			APOTEX	10MG	A208228	001	Jul 07, 2023
AB				20MG	A208228	002	Jul 07, 2023
AB				40MG	A208228	003	Jul 07, 2023
AB			INVAGEN PHARMS	10MG	A208200	001	Apr 07, 2021
AB				20MG	A208200	003	Dec 06, 2022
AB				40MG	A208200	002	Apr 07, 2021
AB			TEVA PHARMS USA	10MG	A208212	001	Sep 30, 2019
AB				20MG	A208212	002	Sep 30, 2019
AB				40MG	A208212	003	Sep 30, 2019

PRESCRIPTION DRUG PRODUCT LIST

VILOXAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

QELBREE

+	SUPERNUS PHARMS	EQ 100MG BASE	N211964 001	Apr 02, 2021
+		EQ 150MG BASE	N211964 002	Apr 02, 2021
+		EQ 200MG BASE	N211964 003	Apr 02, 2021

VILTOLARSEN

SOLUTION; INTRAVENOUS

VILTEPSO

+	NIPPON SHINYAKU	250MG/5ML (50MG/ML)	N212154 001	Aug 12, 2020
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VINBLASTINE SULFATE

INJECTABLE; INJECTION

VINBLASTINE SULFATE

!	FRESENIUS KABI USA	1MG/ML	A089515 001	Apr 29, 1987
!	HIKMA	10MG/VIAL	A089395 001	Apr 09, 1987

VINCRISTINE SULFATE

INJECTABLE; INJECTION

VINCRISTINE SULFATE PFS

!	HOSPIRA	1MG/ML	A071484 001	Apr 19, 1988
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VINORELBINE TARTRATE

INJECTABLE; INJECTION

VINORELBINE TARTRATE

<u>AP</u>	ACTAVIS TOTOWA	<u>EQ 10MG BASE/ML</u>	<u>A078011 001</u>	Jul 22, 2009	
<u>AP</u>	DR REDDYS	<u>EQ 10MG BASE/ML</u>	<u>A202017 001</u>	Sep 12, 2013	
<u>AP</u>	HIKMA	<u>EQ 10MG BASE/ML</u>	<u>A075992 001</u>	Jun 10, 2003	
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A076461 001</u>	Dec 11, 2003	
<u>AP</u>	!	JIANGSU HANSOH PHARM	<u>EQ 10MG BASE/ML</u>	<u>A091106 001</u>	Sep 26, 2012

VISMODEGIB

CAPSULE; ORAL

ERIVEDGE

+	GENENTECH	150MG	N203388 001	Jan 30, 2012
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VITAMIN A PALMITATE

INJECTABLE; INJECTION

AQUASOL A

+	CASPER PHARMA LLC	EQ 50,000 UNITS BASE/ML	N006823 001	
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VOCLOSPORIN

CAPSULE; ORAL

LUPKYNIS

+	AURINIA	7.9MG	N213716 001	Jan 22, 2021
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VONOPRAZAN FUMARATE

TABLET; ORAL

VOQUEZNA

+	PHATHOM	EQ 10MG BASE	N215151 001	Nov 01, 2023
+		EQ 20MG BASE	N215151 002	Nov 01, 2023

VORAPAXAR SULFATE

TABLET; ORAL

ZONTIVITY

+	XSPIRE PHARMA	EQ 2.08MG BASE	N204886 001	May 08, 2014
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VORICONAZOLE

FOR SUSPENSION; ORAL

VFEND

<u>AB</u>	+	PF PRISM CV	<u>200MG/5ML</u>	<u>N021630 001</u>	Dec 19, 2003
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VORICONAZOLE

<u>AB</u>		AMNEAL PHARMS	<u>200MG/5ML</u>	<u>A205034 001</u>	Apr 13, 2016
<u>AB</u>		HAINAN POLY	<u>200MG/5ML</u>	<u>A216805 001</u>	Jan 02, 2024
<u>AB</u>		NOVEL LABS INC	<u>200MG/5ML</u>	<u>A206799 001</u>	May 31, 2016

INJECTABLE; INTRAVENOUS

VFEND

<u>AP</u>	+	PF PRISM CV	<u>200MG/VIAL</u>	<u>N021267 001</u>	May 24, 2002
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VORICONAZOLE

<u>AP</u>		ALMAJECT	<u>200MG/VIAL</u>	<u>A206398 001</u>	Mar 23, 2016
<u>AP</u>		GLAND PHARMA LTD	<u>200MG/VIAL</u>	<u>A211099 001</u>	Mar 31, 2020
<u>AP</u>		MEITHEAL	<u>200MG/VIAL</u>	<u>A214516 001</u>	May 09, 2022
<u>AP</u>		SANDOZ INC	<u>200MG/VIAL</u>	<u>A090862 001</u>	May 30, 2012
<u>AP</u>		SLATE RUN PHARMA	<u>200MG/VIAL</u>	<u>A211661 001</u>	Nov 30, 2018
<u>AP</u>		UBI	<u>200MG/VIAL</u>	<u>A211264 001</u>	Mar 09, 2023
<u>AP</u>		XELLIA PHARMS APS	<u>200MG/VIAL</u>	<u>N208562 001</u>	Mar 09, 2017

PRESCRIPTION DRUG PRODUCT LIST

VORICONAZOLE

INJECTABLE; INTRAVENOUS

VORICONAZOLE

AP	ZYDUS PHARMS	200MG/VIAL	A208983 001	Jul 16, 2018
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TABLET; ORAL

VFEND

AB	+	PF PRISM CV	50MG	N021266 001	May 24, 2002
AB	+	!	200MG	N021266 002	May 24, 2002

VORICONAZOLE

AB		AJANTA PHARMA LTD	50MG	A206181 001	May 24, 2016
AB			200MG	A206181 002	May 24, 2016
AB		AUROBINDO PHARMA LTD	50MG	A206837 001	Jan 22, 2016
AB			200MG	A206837 002	Jan 22, 2016
AB		CADILA	50MG	A206747 001	May 24, 2016
AB			200MG	A206747 002	May 24, 2016
AB		CHARTWELL RX	50MG	A207371 001	May 24, 2016
AB			200MG	A207371 002	May 24, 2016
AB		EPIC PHARMA LLC	50MG	A207049 001	Sep 07, 2016
AB			200MG	A207049 002	Sep 07, 2016
AB		GLENMARK PHARMS LTD	50MG	A203503 001	Sep 02, 2015
AB			200MG	A203503 002	Sep 02, 2015
AB		MYLAN PHARMS INC	50MG	A090547 001	Apr 22, 2010
AB			200MG	A090547 002	Apr 22, 2010
AB		PRINSTON INC	50MG	A206654 001	Aug 08, 2016
AB			200MG	A206654 002	Aug 08, 2016
AB		RISING	50MG	A206762 001	May 24, 2016
AB			200MG	A206762 002	May 24, 2016
AB		SANDOZ INC	50MG	A200265 001	Dec 12, 2011
AB			200MG	A200265 002	Dec 12, 2011

VORINOSTAT

CAPSULE; ORAL

ZOLINZA

+	!	MSD SUB MERCK	100MG	N021991 001	Oct 06, 2006
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VORTIOXETINE HYDROBROMIDE

TABLET; ORAL

TRINTELLIX

+		TAKEDA PHARMS USA	EQ 5MG BASE	N204447 001	Sep 30, 2013
+			EQ 10MG BASE	N204447 002	Sep 30, 2013
+	!		EQ 20MG BASE	N204447 004	Sep 30, 2013

VOSORITIDE

POWDER; SUBCUTANEOUS

VOXZOGO

+	!	BIOMARIN PHARM	0.4MG/VIAL	N214938 001	Nov 19, 2021
+	!		0.56MG/VIAL	N214938 002	Nov 19, 2021
+	!		1.2MG/VIAL	N214938 003	Nov 19, 2021

VOXELOTOR

TABLET; ORAL

OXBRYTA

+		GLOBAL BLOOD THERAPS	300MG	N213137 002	Oct 14, 2022
+	!		500MG	N213137 001	Nov 25, 2019

TABLET, FOR SUSPENSION; ORAL

OXBRYTA

+	!	GLOBAL BLOOD THERAPS	300MG	N216157 001	Dec 17, 2021
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VUTRISIRAN SODIUM

SOLUTION; SUBCUTANEOUS

AMVUTTRA

+	!	ALNYLAM PHARMS INC	EQ 25MG BASE/0.5ML (EQ 25MG BASE/0.5ML)	N215515 001	Jun 13, 2022
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WARFARIN SODIUM

TABLET; ORAL

JANTOVEN

AB		UPSHER SMITH LABS	1MG	A040416 001	Oct 02, 2003
AB			2MG	A040416 002	Oct 02, 2003
AB			2.5MG	A040416 003	Oct 02, 2003
AB			3MG	A040416 004	Oct 02, 2003
AB			4MG	A040416 005	Oct 02, 2003
AB			5MG	A040416 006	Oct 02, 2003
AB			6MG	A040416 007	Oct 02, 2003

PRESCRIPTION DRUG PRODUCT LIST

WARFARIN SODIUM

TABLET; ORAL

JANTOVEN

<u>AB</u>		<u>7.5MG</u>	<u>A040416</u>	<u>008</u>	Oct 02, 2003
<u>AB</u>		<u>10MG</u>	<u>A040416</u>	<u>009</u>	Oct 02, 2003

WARFARIN SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>1MG</u>	<u>A202202</u>	<u>001</u>	Mar 04, 2013
<u>AB</u>		<u>2MG</u>	<u>A202202</u>	<u>002</u>	Mar 04, 2013
<u>AB</u>		<u>2.5MG</u>	<u>A202202</u>	<u>003</u>	Mar 04, 2013
<u>AB</u>		<u>3MG</u>	<u>A202202</u>	<u>004</u>	Mar 04, 2013
<u>AB</u>		<u>4MG</u>	<u>A202202</u>	<u>005</u>	Mar 04, 2013
<u>AB</u>		<u>5MG</u>	<u>A202202</u>	<u>006</u>	Mar 04, 2013
<u>AB</u>		<u>6MG</u>	<u>A202202</u>	<u>007</u>	Mar 04, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A202202</u>	<u>008</u>	Mar 04, 2013
<u>AB</u>		<u>10MG</u>	<u>A202202</u>	<u>009</u>	Mar 04, 2013
<u>AB</u>	BARR	<u>1MG</u>	<u>A040145</u>	<u>001</u>	Mar 26, 1997
<u>AB</u>		<u>2MG</u>	<u>A040145</u>	<u>002</u>	Mar 26, 1997
<u>AB</u>		<u>2.5MG</u>	<u>A040145</u>	<u>003</u>	Mar 26, 1997
<u>AB</u>		<u>3MG</u>	<u>A040145</u>	<u>008</u>	Nov 05, 1998
<u>AB</u>		<u>4MG</u>	<u>A040145</u>	<u>004</u>	Mar 26, 1997
<u>AB</u>		<u>5MG</u>	<u>A040145</u>	<u>005</u>	Mar 26, 1997
<u>AB</u>		<u>6MG</u>	<u>A040145</u>	<u>009</u>	Nov 05, 1998
<u>AB</u>		<u>7.5MG</u>	<u>A040145</u>	<u>006</u>	Mar 26, 1997
<u>AB</u>		<u>10MG</u>	<u>A040145</u>	<u>007</u>	Mar 26, 1997
<u>AB</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A090935</u>	<u>001</u>	May 25, 2011
<u>AB</u>		<u>2MG</u>	<u>A090935</u>	<u>002</u>	May 25, 2011
<u>AB</u>		<u>2.5MG</u>	<u>A090935</u>	<u>003</u>	May 25, 2011
<u>AB</u>		<u>3MG</u>	<u>A090935</u>	<u>004</u>	May 25, 2011
<u>AB</u>		<u>4MG</u>	<u>A090935</u>	<u>005</u>	May 25, 2011
<u>AB</u>		<u>5MG</u>	<u>A090935</u>	<u>006</u>	May 25, 2011
<u>AB</u>		<u>6MG</u>	<u>A090935</u>	<u>007</u>	May 25, 2011
<u>AB</u>		<u>7.5MG</u>	<u>A090935</u>	<u>008</u>	May 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A090935</u>	<u>009</u>	May 25, 2011
<u>AB</u>	IPCA LABS LTD	<u>1MG</u>	<u>A200104</u>	<u>001</u>	Jun 27, 2013
<u>AB</u>		<u>2MG</u>	<u>A200104</u>	<u>002</u>	Jun 27, 2013
<u>AB</u>		<u>2.5MG</u>	<u>A200104</u>	<u>003</u>	Jun 27, 2013
<u>AB</u>		<u>3MG</u>	<u>A200104</u>	<u>004</u>	Jun 27, 2013
<u>AB</u>		<u>4MG</u>	<u>A200104</u>	<u>005</u>	Jun 27, 2013
<u>AB</u>		<u>5MG</u>	<u>A200104</u>	<u>006</u>	Jun 27, 2013
<u>AB</u>		<u>6MG</u>	<u>A200104</u>	<u>007</u>	Jun 27, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A200104</u>	<u>008</u>	Jun 27, 2013
<u>AB</u>		<u>10MG</u>	<u>A200104</u>	<u>009</u>	Jun 27, 2013
<u>AB</u>	PLIVA	<u>1MG</u>	<u>A040616</u>	<u>009</u>	Jul 05, 2006
<u>AB</u>		<u>2MG</u>	<u>A040616</u>	<u>001</u>	Jul 05, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040616</u>	<u>002</u>	Jul 05, 2006
<u>AB</u>		<u>3MG</u>	<u>A040616</u>	<u>003</u>	Jul 05, 2006
<u>AB</u>		<u>4MG</u>	<u>A040616</u>	<u>004</u>	Jul 05, 2006
<u>AB</u>		<u>5MG</u>	<u>A040616</u>	<u>005</u>	Jul 05, 2006
<u>AB</u>		<u>6MG</u>	<u>A040616</u>	<u>006</u>	Jul 05, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040616</u>	<u>007</u>	Jul 05, 2006
<u>AB</u>		<u>10MG</u>	<u>A040616</u>	<u>008</u>	Jul 05, 2006
<u>AB</u>	TARO	<u>1MG</u>	<u>A040301</u>	<u>002</u>	Jul 15, 1999
<u>AB</u>		<u>2MG</u>	<u>A040301</u>	<u>003</u>	Jul 15, 1999
<u>AB</u>		<u>2.5MG</u>	<u>A040301</u>	<u>004</u>	Jul 15, 1999
<u>AB</u>		<u>3MG</u>	<u>A040301</u>	<u>005</u>	Jul 15, 1999
<u>AB</u>		<u>4MG</u>	<u>A040301</u>	<u>006</u>	Jul 15, 1999
<u>AB</u>		<u>5MG</u>	<u>A040301</u>	<u>007</u>	Jul 15, 1999
<u>AB</u>		<u>6MG</u>	<u>A040301</u>	<u>008</u>	Jul 15, 1999
<u>AB</u>		<u>7.5MG</u>	<u>A040301</u>	<u>009</u>	Jul 15, 1999
<u>AB</u>		<u>10MG</u>	<u>A040301</u>	<u>001</u>	Jul 15, 1999
<u>AB</u>	ZYDUS PHARMS USA	<u>1MG</u>	<u>A040663</u>	<u>001</u>	May 30, 2006
<u>AB</u>		<u>2MG</u>	<u>A040663</u>	<u>002</u>	May 30, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040663</u>	<u>003</u>	May 30, 2006
<u>AB</u>		<u>3MG</u>	<u>A040663</u>	<u>004</u>	May 30, 2006
<u>AB</u>		<u>4MG</u>	<u>A040663</u>	<u>005</u>	May 30, 2006
<u>AB</u>		<u>5MG</u>	<u>A040663</u>	<u>006</u>	May 30, 2006
<u>AB</u>		<u>6MG</u>	<u>A040663</u>	<u>007</u>	May 30, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040663</u>	<u>008</u>	May 30, 2006
<u>AB</u>		<u>10MG</u>	<u>A040663</u>	<u>009</u>	May 30, 2006

PRESCRIPTION DRUG PRODUCT LIST

XENON XE-129 HYPERPOLARIZED

GAS; INHALATION

XENOVIEW

+! POLAREAN

N/A

N214375 001 Dec 23, 2022

XENON XE-133

GAS; INHALATION

XENON XE 133

CURIUM

10mCi/VIAL

N018327 001 Mar 09, 1982

20mCi/VIAL

N018327 002 Mar 09, 1982

LANTHEUS MEDCL

10mCi/VIAL

N017284 001

20mCi/VIAL

N017284 002

ZAFIRLUKAST

TABLET; ORAL

ACCOLATEAB + STRIDES PHARMA10MGN020547 003 Sep 17, 1999AB +!20MGN020547 001 Sep 26, 1996ZAFIRLUKASTAB ANNORA PHARMA10MGA212475 001 Sep 10, 2020AB20MGA212475 002 Sep 10, 2020AB AUROBINDO PHARMA10MGA213163 001 Nov 27, 2023AB20MGA213163 002 Nov 27, 2023AB DR REDDYS LABS LTD10MGA090372 001 Nov 18, 2010AB20MGA090372 002 Nov 18, 2010AB RISING PHARMS10MGA204928 001 Aug 25, 2022AB20MGA204928 002 Aug 25, 2022ZALEPLON

CAPSULE; ORAL

SONATAAB + PFIZER5MGN020859 001 Aug 13, 1999AB +!10MGN020859 002 Aug 13, 1999ZALEPLONAB AUROBINDO PHARMA5MGA078829 001 Jun 06, 2008AB10MGA078829 002 Jun 06, 2008AB CHARTWELL MOLECULAR5MGA077505 001 Jun 20, 2008AB10MGA077505 002 Jun 20, 2008AB HIKMA5MGA077237 001 Jun 06, 2008AB10MGA077237 002 Jun 06, 2008AB ORBION PHARMS5MGA090374 001 Sep 17, 2009AB10MGA090374 002 Sep 17, 2009AB UNICHEM5MGA078989 001 Jun 06, 2008AB10MGA078989 002 Jun 06, 2008ZANAMIVIR

POWDER; INHALATION

RELENZA

+! GLAXOSMITHKLINE

5MG

N021036 001 Jul 26, 1999

ZANUBRUTINIB

CAPSULE; ORAL

BRUKINSA

+! BEIGENE

80MG

N213217 001 Nov 14, 2019

ZAVEGEPANT HYDROCHLORIDE

SPRAY, METERED; NASAL

ZAVZPRET

+! PFIZER

EQ 10MG BASE/SPRAY

N216386 001 Mar 09, 2023

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

+! TERSERA

100MCG/1ML (100MCG/ML)

N021060 002 Dec 28, 2004

+!

500MCG/20ML (25MCG/ML)

N021060 001 Dec 28, 2004

+!

500MCG/5ML (100MCG/ML)

N021060 004 Dec 28, 2004

ZIDOVUDINE

CAPSULE; ORAL

RETROVIRAB +! VIIV HLTHCARE100MGN019655 001 Mar 19, 1987ZIDOVUDINEAB AUROBINDO PHARMA100MGA078128 001 Mar 27, 2006

LTD

AB CIPLA LTD100MGA078349 001 May 23, 2007

PRESCRIPTION DRUG PRODUCT LIST

ZIDOVUDINE

INJECTABLE; INJECTION

RETROVIR

+! VIIV HLTHCARE

10MG/ML

N019951 001 Feb 02, 1990

SOLUTION; ORAL

RETROVIR**AA** +! VIIV HLTHCARE**50MG/5ML****N019910 001** Sep 28, 1989ZIDOVUDINE**AA** AUROBINDO**50MG/5ML****A077268 001** Sep 19, 2005**AA** CIPLA LTD**50MG/5ML****A077981 001** Jun 26, 2008

TABLET; ORAL

ZIDOVUDINE**AB** AUROBINDO**300MG****A077267 001** Sep 19, 2005**AB** CIPLA**300MG****A090561 001** Oct 27, 2010**AB** ! HETERO LABS LTD III**300MG****A090092 001** Apr 25, 2008**AB** MYLAN PHARMS INC**300MG****A078922 001** Feb 14, 2008ZILEUTON

TABLET; ORAL

ZYFLO

+! CHIESI

600MG

N020471 003 Dec 09, 1996

TABLET, EXTENDED RELEASE; ORAL

ZILEUTON**AB** AIZANT**600MG****A211390 001** Oct 23, 2020**AB** ANNORA PHARMA**600MG****A215742 001** Oct 11, 2022**AB** ! RISING**600MG****A204929 001** Mar 17, 2017**AB** STRIDES PHARMA**600MG****A212670 001** Dec 16, 2019ZILUCOPLAN SODIUM

SOLUTION; SUBCUTANEOUS

ZILBRYSQ

+! UCB INC

EQ 16.6MG BASE/0.416ML (EQ 16.6MG
BASE/0.416ML)

N216834 001 Oct 17, 2023

+!
EQ 23MG BASE/0.574ML (EQ 23MG
BASE/0.574ML)

N216834 002 Oct 17, 2023

+!
EQ 32.4MG BASE/0.81ML (EQ 32.4
BASE/0.81ML)

N216834 003 Oct 17, 2023

ZINC ACETATE

CAPSULE; ORAL

GALZIN

+ TEVA

EQ 25MG ZINC

N020458 001 Jan 28, 1997

+!

EQ 50MG ZINC

N020458 002 Jan 28, 1997

ZINC CHLORIDE

INJECTABLE; INJECTION

ZINC CHLORIDE**AP** EXELA PHARMA**EQ 1MG ZINC/ML****A212007 001** May 21, 2021ZINC CHLORIDE IN PLASTIC CONTAINER**AP** +! HOSPIRA**EQ 1MG ZINC/ML****N018959 001** Jun 26, 1986ZINC SULFATE

SOLUTION; INTRAVENOUS

ZINC SULFATE**AP** +! AM REGENT**EQ 10MG BASE/10ML (EQ 1MG BASE/ML)****N209377 003** Apr 15, 2020**AP** +!**EQ 25MG BASE/5ML (EQ 5MG BASE/ML)****N209377 002** Jul 18, 2019**AP** +!**EQ 30MG BASE/10ML (EQ 3MG BASE/ML)****N209377 001** Jul 18, 2019**AP** FRESENIUS KABI USA**EQ 10MG BASE/10ML (EQ 1MG BASE/ML)****A216145 001** Dec 27, 2022**AP****EQ 25MG BASE/5ML (EQ 5MG BASE/ML)****A216145 002** Dec 27, 2022**AP****EQ 30MG BASE/10ML (EQ 3MG BASE/ML)****A216145 003** Dec 27, 2022**AP** GLAND PHARMA LTD**EQ 10MG BASE/10ML (EQ 1MG BASE/ML)****A216249 003** Sep 01, 2023**AP****EQ 25MG BASE/5ML (EQ 5MG BASE/ML)****A216249 001** May 03, 2022**AP****EQ 30MG BASE/10ML (EQ 3MG BASE/ML)****A216249 002** May 03, 2022**AP** ZYDUS PHARMS**EQ 30MG BASE/10ML (EQ 3MG BASE/ML)****A217074 003** Aug 22, 2023**AP****EQ 10MG BASE/10ML (EQ 1MG BASE/ML)****A217074 001** Aug 22, 2023**AP****EQ 25MG BASE/5ML (EQ 5MG BASE/ML)****A217074 002** Aug 22, 2023ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

GEODON**AB** +! VIATRIS**EQ 20MG BASE****N020825 001** Feb 05, 2001**AB** +**EQ 40MG BASE****N020825 002** Feb 05, 2001**AB** +**EQ 60MG BASE****N020825 003** Feb 05, 2001**AB** +**EQ 80MG BASE****N020825 004** Feb 05, 2001

PRESCRIPTION DRUG PRODUCT LIST

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

ZIPRASIDONE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 20MG BASE</u>	<u>A077561 001</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077561 002</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077561 003</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077561 004</u>	Mar 02, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 20MG BASE</u>	<u>A204117 001</u>	Dec 27, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A204117 002</u>	Dec 27, 2016
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204117 003</u>	Dec 27, 2016
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204117 004</u>	Dec 27, 2016
<u>AB</u>	CADILA	<u>EQ 20MG BASE</u>	<u>A208988 001</u>	Aug 22, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208988 002</u>	Aug 22, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A208988 003</u>	Aug 22, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A208988 004</u>	Aug 22, 2017
<u>AB</u>	CHARTWELL RX	<u>EQ 20MG BASE</u>	<u>A090348 001</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090348 002</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A090348 003</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A090348 004</u>	Sep 05, 2012
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 20MG BASE</u>	<u>A077565 001</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077565 002</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077565 003</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077565 004</u>	Mar 02, 2012
<u>AB</u>	LUPIN PHARMS	<u>EQ 20MG BASE</u>	<u>A077560 001</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077560 002</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077560 003</u>	Mar 02, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077560 004</u>	Mar 02, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A204375 001</u>	Feb 17, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A204375 002</u>	Feb 17, 2017
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A204375 003</u>	Feb 17, 2017
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204375 004</u>	Feb 17, 2017
<u>AB</u>	SANDOZ INC	<u>EQ 20MG BASE</u>	<u>A077562 001</u>	Jun 01, 2012
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077562 002</u>	Jun 01, 2012
<u>AB</u>		<u>EQ 60MG BASE</u>	<u>A077562 003</u>	Jun 01, 2012
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A077562 004</u>	Jun 01, 2012

ZIPRASIDONE MESYLATE

POWDER; INTRAMUSCULAR

GEODON

<u>AP</u>	<u>+</u> VIATRIS	<u>EQ 20MG BASE/VIAL</u>	<u>N020919 001</u>	Jun 21, 2002
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ZIPRASIDONE MESYLATE

<u>AP</u>	GLAND PHARMA LTD	<u>EQ 20MG BASE/VIAL</u>	<u>A211908 001</u>	Dec 26, 2019
<u>AP</u>	MSN	<u>EQ 20MG BASE/VIAL</u>	<u>A216091 001</u>	Sep 15, 2022

ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

RECLAST

<u>AP</u>	<u>+</u> SANDOZ	<u>EQ 5MG BASE/100ML</u>	<u>N021817 001</u>	Apr 16, 2007
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ZOLEDRONIC

<u>AP</u>	<u>!</u> GLAND PHARMA LTD	<u>EQ 4MG BASE/100ML</u>	<u>A205749 001</u>	Jun 29, 2018
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ZOLEDRONIC ACID

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 4MG BASE/5ML</u>	<u>A205279 001</u>	Nov 28, 2016
<u>AP</u>	AMNEAL	<u>EQ 4MG BASE/100ML</u>	<u>A210174 001</u>	Oct 27, 2017
<u>AP</u>	APOTEX	<u>EQ 5MG BASE/100ML</u>	<u>A204367 001</u>	Dec 24, 2015
<u>AP</u>	AVET LIFESCIENCES	<u>EQ 4MG BASE/5ML</u>	<u>A201783 001</u>	Mar 12, 2013
<u>AP</u>	BPI LABS	<u>EQ 4MG BASE/5ML</u>	<u>A207341 001</u>	Dec 29, 2017
<u>AP</u>	CHARTWELL RX	<u>EQ 4MG BASE/5ML</u>	<u>A202571 001</u>	May 07, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A202163 001</u>	Aug 05, 2013
<u>AP</u>	<u>!</u> DR REDDYS LABS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A091186 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A091363 001</u>	Mar 29, 2013
<u>AP</u>	EPIC PHARMA LLC	<u>EQ 4MG BASE/5ML</u>	<u>A202548 001</u>	May 22, 2014
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A200918 001</u>	Aug 21, 2014
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 4MG BASE/5ML</u>	<u>A091516 001</u>	Apr 23, 2015
<u>AP</u>	GLAND	<u>EQ 5MG BASE/100ML</u>	<u>A204217 001</u>	Aug 18, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 4MG BASE/5ML</u>	<u>A202930 001</u>	Aug 05, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A209578 001</u>	Aug 08, 2019
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 4MG BASE/5ML</u>	<u>A202182 001</u>	Jun 03, 2013
<u>AP</u>	HOSPIRA	<u>EQ 4MG BASE/5ML</u>	<u>A090621 001</u>	Mar 19, 2015
<u>AP</u>	HOSPIRA INC	<u>EQ 5MG BASE/100ML</u>	<u>A202837 001</u>	Apr 05, 2013
<u>AP</u>	<u>+</u> INFORLIFE	<u>EQ 4MG BASE/100ML</u>	<u>N203231 001</u>	Aug 02, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A202828 001</u>	Sep 23, 2013
<u>AP</u>	MEITHEAL	<u>EQ 5MG BASE/100ML</u>	<u>A213371 001</u>	Jun 05, 2023
<u>AP</u>	MYLAN LABS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A202650 001</u>	Mar 04, 2013

PRESCRIPTION DRUG PRODUCT LIST

ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

ZOLEDRONIC ACID

<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A203841 001</u>	Feb 14, 2017
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A205254 001</u>	Oct 27, 2017
<u>AP</u>	SAGENT PHARMS INC	<u>EQ 4MG BASE/5ML</u>	<u>A091493 001</u>	Nov 24, 2014
<u>AP</u>	USV	<u>EQ 4MG BASE/5ML</u>	<u>A202923 001</u>	Sep 04, 2014
	SOLUTION; INTRAVENOUS			
	ZOLEDRONIC ACID			
	HOSPIRA	EQ 4MG BASE/100ML (EQ 0.04MG BASE/ML)	N204016 001	Dec 28, 2015

ZOLMITRIPTAN

SPRAY; NASAL

ZOLMITRIPTAN

<u>AB</u>	PADAGIS ISRAEL	<u>2.5MG/SPRAY</u>	<u>A212469 001</u>	Sep 30, 2021
<u>AB</u>		<u>5MG/SPRAY</u>	<u>A212469 002</u>	Sep 30, 2021
	<u>ZOMIG</u>			
<u>AB</u>	+ AMNEAL	<u>2.5MG/SPRAY</u>	<u>N021450 003</u>	Sep 16, 2013
<u>AB</u>	+!	<u>5MG/SPRAY</u>	<u>N021450 004</u>	Sep 30, 2003

TABLET; ORAL

ZOLMITRIPTAN

<u>AB</u>	AJANTA PHARMA LTD	<u>2.5MG</u>	<u>A204041 001</u>	May 20, 2016
<u>AB</u>	!	<u>5MG</u>	<u>A204041 002</u>	May 20, 2016
<u>AB</u>	ALEMBIC	<u>2.5MG</u>	<u>A204232 001</u>	Sep 30, 2015
<u>AB</u>		<u>5MG</u>	<u>A204232 002</u>	Sep 30, 2015
<u>AB</u>	APPCO	<u>2.5MG</u>	<u>A206973 001</u>	Jun 30, 2017
<u>AB</u>		<u>5MG</u>	<u>A206973 002</u>	Jun 30, 2017
<u>AB</u>	AUROBINDO PHARMA	<u>2.5MG</u>	<u>A207021 001</u>	May 11, 2016
<u>AB</u>		<u>5MG</u>	<u>A207021 002</u>	May 11, 2016
<u>AB</u>	GLENMARK GENERICS	<u>2.5MG</u>	<u>A201779 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A201779 002</u>	May 14, 2013
<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A204284 001</u>	Apr 09, 2014
<u>AB</u>		<u>5MG</u>	<u>A204284 002</u>	Apr 09, 2014
<u>AB</u>	JUBILANT GENERICS	<u>2.5MG</u>	<u>A202279 001</u>	Nov 20, 2014
<u>AB</u>		<u>5MG</u>	<u>A202279 002</u>	Nov 20, 2014
<u>AB</u>	ORBION PHARMS	<u>2.5MG</u>	<u>A203726 001</u>	Oct 21, 2019
<u>AB</u>		<u>5MG</u>	<u>A203726 002</u>	Oct 21, 2019
<u>AB</u>	PLD ACQUISITIONS LLC	<u>2.5MG</u>	<u>A207867 001</u>	Feb 27, 2017
<u>AB</u>		<u>5MG</u>	<u>A207867 002</u>	Feb 27, 2017
<u>AB</u>	ZYDUS PHARMS	<u>2.5MG</u>	<u>A203019 001</u>	Jul 11, 2018
<u>AB</u>		<u>5MG</u>	<u>A203019 002</u>	Jul 11, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

ZOLMITRIPTAN

<u>AB</u>	ALEMBIC	<u>2.5MG</u>	<u>A205074 001</u>	Dec 01, 2016
<u>AB</u>		<u>5MG</u>	<u>A205074 002</u>	Dec 01, 2016
<u>AB</u>	GLENMARK GENERICS	<u>2.5MG</u>	<u>A202560 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A202560 002</u>	May 14, 2013
<u>AB</u>	JUBILANT GENERICS	<u>2.5MG</u>	<u>A202956 001</u>	Sep 17, 2015
<u>AB</u>		<u>5MG</u>	<u>A202956 002</u>	Sep 17, 2015
<u>AB</u>	ZYDUS PHARMS USA INC	<u>2.5MG</u>	<u>A202890 001</u>	May 15, 2013
<u>AB</u>	!	<u>5MG</u>	<u>A202890 002</u>	May 15, 2013

ZOLPIDEM TARTRATE

CAPSULE; ORAL

ZOLPIDEM TARTRATE

+! ALMATICA

7.5MG

N215721 001 May 09, 2023

TABLET; ORAL

AMBIEN

<u>AB</u>	+ SANOFI AVENTIS US	<u>5MG</u>	<u>N019908 001</u>	Dec 16, 1992
<u>AB</u>	+!	<u>10MG</u>	<u>N019908 002</u>	Dec 16, 1992

ZOLPIDEM TARTRATE

<u>AB</u>	ACME LABS	<u>5MG</u>	<u>A077214 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077214 002</u>	Apr 23, 2007
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A077884 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077884 002</u>	Apr 23, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078413 001</u>	May 04, 2007
<u>AB</u>		<u>10MG</u>	<u>A078413 002</u>	May 04, 2007
<u>AB</u>	CHARTWELL MOLECULAR	<u>5MG</u>	<u>A077388 001</u>	Jul 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077388 002</u>	Jul 30, 2012
<u>AB</u>	INVAGEN PHARMS	<u>5MG</u>	<u>A078184 001</u>	Sep 07, 2007
<u>AB</u>		<u>10MG</u>	<u>A078184 002</u>	Sep 07, 2007
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A077322 001</u>	Apr 23, 2007

PRESCRIPTION DRUG PRODUCT LIST

ZOLPIDEM TARTRATE

TABLET; ORAL

ZOLPIDEM TARTRATE

<u>AB</u>		<u>10MG</u>	<u>A077322 002</u>	Apr 23, 2007
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076410 001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A076410 002</u>	Apr 23, 2007
<u>AB</u>	TORRENT PHARMS	<u>5MG</u>	<u>A077903 001</u>	Aug 17, 2007
<u>AB</u>		<u>10MG</u>	<u>A077903 002</u>	Aug 17, 2007

TABLET; SUBLINGUAL

EDLUAR

<u>AB</u>	+	MYLAN SPECIALITY LP	<u>5MG</u>	<u>N021997 001</u>	Mar 13, 2009
<u>AB</u>	+	!	<u>10MG</u>	<u>N021997 002</u>	Mar 13, 2009

ZOLPIDEM TARTRATE

<u>AB</u>		DR REDDYS	<u>1.75MG</u>	<u>A204503 001</u>	Nov 18, 2016
<u>AB</u>			<u>3.5MG</u>	<u>A204503 002</u>	Nov 18, 2016
<u>AB</u>		NOVEL LABS INC	<u>1.75MG</u>	<u>A204299 001</u>	Jun 03, 2015
<u>AB</u>	!		<u>3.5MG</u>	<u>A204299 002</u>	Jun 03, 2015
<u>AB</u>		PAR FORM	<u>5MG</u>	<u>A201509 001</u>	Aug 01, 2016
<u>AB</u>			<u>10MG</u>	<u>A201509 002</u>	Aug 01, 2016
<u>AB</u>		PAR PHARM INC	<u>1.75MG</u>	<u>A204229 001</u>	Sep 11, 2017
<u>AB</u>			<u>3.5MG</u>	<u>A204229 002</u>	Sep 11, 2017

TABLET, EXTENDED RELEASE; ORAL

AMBIEN CR

<u>AB</u>	+	SANOFI AVENTIS US	<u>6.25MG</u>	<u>N021774 002</u>	Sep 02, 2005
<u>AB</u>	+	!	<u>12.5MG</u>	<u>N021774 001</u>	Sep 02, 2005

ZOLPIDEM TARTRATE

<u>AB</u>		APOTEX	<u>6.25MG</u>	<u>A200266 001</u>	Sep 10, 2013
<u>AB</u>			<u>12.5MG</u>	<u>A200266 002</u>	Sep 10, 2013
<u>AB</u>		LUPIN LTD	<u>6.25MG</u>	<u>A078970 001</u>	Sep 11, 2013
<u>AB</u>			<u>12.5MG</u>	<u>A078970 002</u>	Sep 11, 2013
<u>AB</u>		SANDOZ	<u>6.25MG</u>	<u>A090107 001</u>	Jul 01, 2011
<u>AB</u>			<u>12.5MG</u>	<u>A090107 002</u>	Jul 01, 2011
<u>AB</u>		SUN PHARM	<u>6.25MG</u>	<u>A204170 001</u>	Jan 24, 2017
<u>AB</u>			<u>12.5MG</u>	<u>A204170 002</u>	Jan 24, 2017

ZONISAMIDE

CAPSULE; ORAL

ZONEGRAN

<u>AB</u>	+	CONCORDIA	<u>25MG</u>	<u>N020789 003</u>	Aug 22, 2003
<u>AB</u>	+	!	<u>100MG</u>	<u>N020789 001</u>	Mar 27, 2000

ZONISAMIDE

<u>AB</u>		APOTEX INC	<u>25MG</u>	<u>A077642 001</u>	Dec 22, 2005
<u>AB</u>			<u>50MG</u>	<u>A077642 002</u>	Dec 22, 2005
<u>AB</u>			<u>100MG</u>	<u>A077642 003</u>	Dec 22, 2005
<u>AB</u>		AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A077645 002</u>	Sep 29, 2006
<u>AB</u>			<u>50MG</u>	<u>A077645 003</u>	Sep 29, 2006
<u>AB</u>			<u>100MG</u>	<u>A077645 001</u>	Dec 22, 2005
<u>AB</u>		BIONPHARMA	<u>25MG</u>	<u>A077813 001</u>	Aug 16, 2006
<u>AB</u>			<u>50MG</u>	<u>A077813 002</u>	Aug 16, 2006
<u>AB</u>			<u>100MG</u>	<u>A077813 003</u>	Aug 16, 2006
<u>AB</u>		CADILA	<u>25MG</u>	<u>A077625 001</u>	Oct 16, 2006
<u>AB</u>			<u>50MG</u>	<u>A077625 002</u>	Oct 16, 2006
<u>AB</u>			<u>100MG</u>	<u>A077625 003</u>	Oct 16, 2006
<u>AB</u>		GLENMARK GENERICS	<u>25MG</u>	<u>A077651 001</u>	Jan 30, 2006
<u>AB</u>			<u>50MG</u>	<u>A077651 002</u>	Jan 30, 2006
<u>AB</u>			<u>100MG</u>	<u>A077651 003</u>	Jan 30, 2006
<u>AB</u>		GRANULES	<u>25MG</u>	<u>A077636 003</u>	Jul 27, 2006
<u>AB</u>			<u>50MG</u>	<u>A077636 002</u>	Jul 27, 2006
<u>AB</u>			<u>100MG</u>	<u>A077636 001</u>	Dec 22, 2005
<u>AB</u>		INVAGEN PHARMS	<u>25MG</u>	<u>A077869 001</u>	May 31, 2006
<u>AB</u>			<u>50MG</u>	<u>A077869 002</u>	May 31, 2006
<u>AB</u>			<u>100MG</u>	<u>A077869 003</u>	May 31, 2006
<u>AB</u>		SUN PHARM INDS (IN)	<u>25MG</u>	<u>A077634 001</u>	Mar 17, 2006
<u>AB</u>			<u>50MG</u>	<u>A077634 002</u>	Mar 17, 2006
<u>AB</u>			<u>100MG</u>	<u>A077634 003</u>	Mar 17, 2006
<u>AB</u>		UNICHEM	<u>25MG</u>	<u>A214492 001</u>	Jan 26, 2021
<u>AB</u>			<u>50MG</u>	<u>A214492 002</u>	Jan 26, 2021
<u>AB</u>			<u>100MG</u>	<u>A214492 003</u>	Jan 26, 2021

SUSPENSION; ORAL

ZONISAMIDE

	+	AZURITY	100MG/5ML	N214273 001	Jul 15, 2022
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PRESCRIPTION DRUG PRODUCT LISTZURANOLONE

CAPSULE; ORAL

ZURZUVAE

+ BIOGEN INC

20MG

N217369 001 Oct 31, 2023

+

25MG

N217369 002 Oct 31, 2023

+!

30MG

N217369 003 Oct 31, 2023

OTC DRUG PRODUCT LIST

ACETAMINOPHEN

SUPPOSITORY; RECTAL

ACEPHEN

COSETTE	120MG	N018060 001	
	325MG	A072344 001	Mar 27, 1992
	650MG	A072237 001	Mar 27, 1992

ACETAMINOPHEN

PERRIGO NEW YORK	120MG	A070607 001	Apr 06, 1987
	650MG	A070608 001	Dec 01, 1986
+ TARO	120MG	N018337 003	Sep 12, 1983
+	325MG	N018337 002	
+!	650MG	N018337 001	

INFANTS' FEVERALL

+ TARO	80MG	N018337 004	Aug 26, 1992
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TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

AUROBINDO PHARMA	650MG	A207229 001	Nov 09, 2016
GRANULES	650MG	A211544 001	Apr 16, 2019
HERITAGE PHARMA	650MG	A207035 001	May 31, 2018
MARKSANS PHARMA	650MG	A215486 001	Aug 25, 2021
OHM LABS	650MG	A076200 001	Mar 19, 2002
PERRIGO	650MG	A075077 001	Feb 25, 2000
SUN PHARM INDS LTD	650MG	A078569 001	Dec 14, 2011

TYLENOL

+! J AND J CONSUMER INC	650MG	N019872 001	Jun 08, 1994
+!	650MG	N019872 002	Jan 11, 2001

ACETAMINOPHEN; ASPIRIN; CAFFEINE

TABLET; ORAL

ACETAMINOPHEN, ASPIRIN AND CAFFEINE

AUROBINDO PHARMA LTD	250MG; 250MG; 65MG	A211695 001	Feb 02, 2022
GRANULES	250MG; 250MG; 65MG	A214039 001	Feb 23, 2021
PERRIGO	250MG; 250MG; 65MG	A075794 001	Nov 26, 2001

EXCEDRIN (MIGRAINE RELIEF)

+! GLAXOSMITHKLINE CONS	250MG; 250MG; 65MG	N020802 001	Jan 14, 1998
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ACETAMINOPHEN; IBUPROFEN

TABLET; ORAL

ACETAMINOPHEN AND IBUPROFEN

BIONPHARMA	250MG; 125MG	A216999 001	Aug 01, 2023
GRANULES	250MG; 125MG	A216592 001	Jul 13, 2023
L PERRIGO CO	250MG; 125MG	A214836 001	Feb 28, 2023
MARKSANS PHARMA	250MG; 125MG	A216994 001	Jul 10, 2023

ADVIL DUAL ACTION WITH ACETAMINOPHEN

+! GLAXOSMITHKLINE	250MG; 125MG	N211733 001	Feb 28, 2020
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ADAPALENE

GEL; TOPICAL

ADAPALENE

GLENMARK GENERICS	0.1%	A091314 001	Jul 01, 2010
P AND L	0.1%	A090962 001	Jun 02, 2010
TARO	0.1%	A215940 001	Jan 14, 2022

DIFFERIN

+! GALDERMA LABS LP	0.1%	N020380 002	Jul 08, 2016
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ALCAFTADINE

SOLUTION/DROPS; OPHTHALMIC

LASTACFT

+! ABBVIE	0.25%	N022134 001	Jul 28, 2010
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ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

AVAGARD

+! 3M	61%; 1%	N021074 001	Jun 07, 2001
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ASPIRIN

CAPSULE; ORAL

VAZALORE

+ PLX PHARMA	81MG	N203697 002	Feb 26, 2021
+!	325MG	N203697 001	Jan 14, 2013

OTC DRUG PRODUCT LIST

AVOBENZONE; ECAMSULE; OCTOCRYLENE

CREAM; TOPICAL

ANTHELIOS SX

+! LOREAL USA 2%;2%;10% N021502 001 Jul 21, 2006

CAPITAL SOLEIL 15

+! LOREAL USA 2%;3%;10% N021501 001 Oct 02, 2006

AVOBENZONE; ECAMSULE; OCTOCRYLENE; TITANIUM DIOXIDE

CREAM; TOPICAL

ANTHELIOS 20

+! LOREAL USA 2%;2%;10%;2% N021471 001 Oct 05, 2006

ANTHELIOS 40

+! LOREAL USA 2%;3%;10%;5% N022009 001 Mar 31, 2008

+! 2%;3%;10%;5% N022009 002 Oct 29, 2009

AZELASTINE HYDROCHLORIDE

SPRAY, METERED; NASAL

ASTEPRO ALLERGY

+! BAYER HLTHCARE 0.2055MG/SPRAY N213872 001 Jun 17, 2021

CHILDREN'S ASTEPRO ALLERGY

+! BAYER HLTHCARE 0.2055MG/SPRAY N213872 002 Jun 17, 2021

BENTOQUATAM

LOTION; TOPICAL

IVY BLOCK

+! STAND HOMEOPATH 5% N020532 001 Aug 26, 1996

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

LUMIFY

+! BAUSCH AND LOMB INC 0.025% N208144 001 Dec 22, 2017

BUDESONIDE

SPRAY, METERED; NASAL

BUDESONIDE

APOTEX INC 0.032MG/SPRAY A078949 002 Nov 20, 2015

RHINOCORT ALLERGY

+! J AND J CONSUMER 0.032MG/SPRAY N020746 003 Mar 23, 2015

INC

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

BUTENAFINE HYDROCHLORIDE

TARO 1% A205181 001 Nov 16, 2017

LOTRIMIN ULTRA

+! BAYER HEALTHCARE 1% N021307 001 Dec 07, 2001

LLC

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL

FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE

PERRIGO R AND D 800MG;10MG;165MG A077355 001 Feb 06, 2008

PEPCID COMPLETE

+! J AND J CONSUMER 800MG;10MG;165MG N020958 001 Oct 16, 2000

INC

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

AUROBINDO PHARMA 10MG A209107 001 Jul 20, 2018

LTD

+ BIONPHARMA 5MG N022429 001 Jul 23, 2009

+! 10MG N022429 004 Jul 23, 2009

CATALENT 10MG A213105 001 Sep 21, 2020

STRIDES PHARMA 10MG A205291 001 Jul 21, 2017

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AUROBINDO PHARMA 10MG A209107 002 Jul 20, 2018

LTD

+ BIONPHARMA 5MG N022429 003 Jul 23, 2009

+! 10MG N022429 002 Jul 23, 2009

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS 5MG/5ML A090765 002 Oct 07, 2009

AUROBINDO PHARMA 5MG/5ML A090750 002 Feb 02, 2010

BAJAJ 5MG/5ML A091327 001 Oct 17, 2011

CHARTWELL MOLECULAR 5MG/5ML A091130 001 Apr 22, 2011

HETERO LABS LTD III 5MG/5ML A210622 001 Mar 13, 2019

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

PERRIGO R AND D	5MG/5ML	A204226	001	Sep 09, 2013
	5MG/5ML	A090254	002	Apr 09, 2008
QUAGEN	5MG/5ML	A212266	001	May 16, 2019
TARO	5MG/5ML	A090182	002	Apr 22, 2008
	5MG/5ML	A201546	001	May 20, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS	5MG/5ML	A090765	001	Oct 07, 2009
AUROBINDO PHARMA	5MG/5ML	A090750	001	Feb 02, 2010
BAJAJ	5MG/5ML	A091327	002	Oct 17, 2011
CHARTWELL MOLECULAR	5MG/5ML	A091130	002	Apr 22, 2011
PERRIGO R AND D	5MG/5ML	A090254	001	Apr 09, 2008
TARO	5MG/5ML	A090182	001	Apr 22, 2008
	5MG/5ML	A201546	002	May 20, 2011

CHILDREN'S ZYRTEC ALLERGY

+! J AND J CONSUMER INC	5MG/5ML	N022155	002	Nov 16, 2007
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CHILDREN'S ZYRTEC HIVES RELIEF

+! J AND J CONSUMER INC	5MG/5ML	N022155	001	Nov 16, 2007
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TABLET; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS NY	5MG	A078780	001	Jan 21, 2010
	10MG	A078780	004	Jan 21, 2010
APOTEX INC	5MG	A078317	001	Dec 27, 2007
	10MG	A078317	002	Dec 27, 2007
AUROBINDO PHARMA LTD	5MG	A090760	001	Aug 05, 2015
	10MG	A090760	003	Aug 05, 2015
CONTRACT PHARMACAL	5MG	A076047	001	Dec 27, 2007
	10MG	A076047	002	Dec 27, 2007
DR REDDYS LABS LTD	5MG	A078343	004	Jan 15, 2008
	10MG	A078343	003	Jan 15, 2008
GLENMARK PHARMS INC	5MG	A078427	003	Dec 28, 2007
	10MG	A078427	004	Dec 28, 2007
GRANULES	10MG	A209274	001	Dec 22, 2017
IPCA LABS LTD	5MG	A202277	002	Mar 11, 2014
	10MG	A202277	004	Mar 11, 2014
MARKSANS PHARMA	5MG	A078933	001	Jun 15, 2010
	10MG	A078933	002	Jun 15, 2010
MYLAN	5MG	A076677	001	Dec 27, 2007
	10MG	A076677	002	Dec 27, 2007
ORBION PHARMS	5MG	A078862	001	Feb 19, 2009
	10MG	A078862	002	Feb 19, 2009
PERRIGO R AND D	5MG	A078336	001	Dec 27, 2007
	10MG	A078336	002	Dec 27, 2007
PLD ACQUISITIONS	5MG	A077946	001	Dec 27, 2007
	10MG	A077946	002	Dec 27, 2007
SUN PHARM INDS LTD	5MG	A077498	001	Dec 27, 2007
	10MG	A077498	002	Dec 27, 2007
TARO	5MG	A078072	001	Jul 22, 2009
	5MG	A078072	003	Jul 22, 2009
UNIQUE	5MG	A077829	001	Aug 26, 2009
	10MG	A077829	004	Aug 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD	5MG	A078343	001	Jan 15, 2008
	10MG	A078343	002	Jan 15, 2008
IPCA LABS LTD	5MG	A202277	001	Mar 11, 2014
	10MG	A202277	003	Mar 11, 2014
MARKSANS PHARMA	5MG	A078933	003	Jun 15, 2010
	10MG	A078933	004	Jun 15, 2010
MYLAN	5MG	A076677	004	Dec 27, 2007
	10MG	A076677	003	Dec 27, 2007
ORBION PHARMS	5MG	A078862	003	Feb 19, 2009
	10MG	A078862	004	Feb 19, 2009
PERRIGO R AND D	5MG	A078336	003	Dec 27, 2007
	10MG	A078336	004	Dec 27, 2007
SUN PHARM INDS LTD	5MG	A077498	003	Dec 27, 2007
	10MG	A077498	004	Dec 27, 2007
UNIQUE	5MG	A077829	003	Aug 26, 2009
!	10MG	A077829	002	Aug 26, 2009

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS NY	5MG	A078780	003	Jan 21, 2010
	10MG	A078780	002	Jan 21, 2010
AUROBINDO PHARMA LTD	5MG	A090760	002	Aug 05, 2015
	10MG	A090760	004	Aug 05, 2015
TARO	10MG	A078072	002	Jul 22, 2009
	10MG	A078072	004	Jul 22, 2009

ZYRTEC ALLERGY

+! J AND J CONSUMER INC	10MG	N019835	004	Nov 16, 2007
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TABLET, CHEWABLE;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

JUBILANT GENERICS	5MG	A091116	001	Feb 19, 2015
	10MG	A091116	002	Feb 19, 2015
NOVEL LABS INC	5MG	A206793	001	Mar 08, 2016
	10MG	A206793	002	Mar 08, 2016
SANDOZ	5MG	A078692	001	Feb 14, 2008
!	10MG	A078692	002	Feb 14, 2008
SUN PHARM	5MG	A090142	001	Aug 30, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

JUBILANT GENERICS	5MG	A091116	003	Feb 19, 2015
	10MG	A091116	004	Feb 19, 2015
SUN PHARM	5MG	A090142	003	Aug 30, 2011

CHILDREN'S ZYRTEC ALLERGY

+ J AND J CONSUMER INC	2.5MG	N021621	007	Nov 30, 2020
	10MG	N021621	004	Nov 16, 2007

TABLET, ORALLY DISINTEGRATING;ORAL

CETIRIZINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	10MG	A213557	001	Sep 11, 2020
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CETIRIZINE HYDROCHLORIDE ALLERGY

PERRIGO R AND D	10MG	A205490	001	Sep 02, 2015
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ZYRTEC ALLERGY

+! J AND J CONSUMER INC	10MG	N022578	001	Sep 03, 2010
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CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS	5MG;120MG	A077170	001	Feb 25, 2008
PERRIGO R AND D	5MG;120MG	A210719	001	Nov 16, 2018
PLD ACQUISITIONS	5MG;120MG	A077991	001	Mar 05, 2008
SUN PHARM INDS LTD	5MG;120MG	A090922	001	Sep 28, 2012

ZYRTEC-D 12 HOUR

+! J AND J CONSUMER INC	5MG;120MG	N021150	002	Nov 09, 2007
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CHLORHEXIDINE GLUCONATE

AEROSOL, METERED;TOPICAL

EXIDINE

+! XTTRIUM	4%	N019127	001	Dec 24, 1984
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CLOTH;TOPICAL

CHLORHEXIDINE GLUCONATE

+! SAGE PRODS	2%	N021669	001	Apr 25, 2005
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READYPREP CHG

MEDLINE INDUSTRIES	2%	N207964	001	Nov 20, 2018
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SOLUTION;TOPICAL

CHG SCRUB

ECOLAB	4%	N019258	002	Jul 22, 1986
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CHLORHEXIDINE GLUCONATE

BAJAJ	0.75%	N020111	001	Sep 11, 1997
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CIDA-STAT

ECOLAB	2%	N019258	001	Jul 22, 1986
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EXIDINE

+! XTTRIUM	2%	N019422	001	Dec 17, 1985
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	4%	N019125	001	Dec 24, 1984
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HIBICLENS

+! MOLNLYCKE HLTH	4%	N017768	001	
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OTC DRUG PRODUCT LIST

CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

HIBISTAT

+! MOLNLYCKE HLTH 0.5% N018300 001

SPONGE; TOPICAL

BIOSCRUB

GRIFFEN 4% N019822 001 Mar 31, 1989

CHLORHEXIDINE GLUCONATE

! BECTON DICKINSON 4% A072525 001 Oct 24, 1989

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SOLUTION; TOPICAL

SOLUPREP S

+! 3M HEALTH CARE 2%;70% N208288 001 Aug 08, 2018

SPONGE; TOPICAL

CHLORAPREP ONE-STEP

+! BECTON DICKINSON CO 2%;70% (3ML) N020832 001 Jul 14, 2000

+! 2%;70% (10.5ML) N020832 004 Aug 20, 2003

+! 2%;70% (26ML) N020832 006 Nov 21, 2006

+! 2%;70% (1ML) N020832 008 Oct 23, 2008

CHLORAPREP ONE-STEP FREPP

+! BECTON DICKINSON CO 2%;70% (1.5ML) N020832 003 Apr 26, 2002

CHLORAPREP WITH TINT

+! BECTON DICKINSON CO 2%;70% (26ML) N020832 002 May 03, 2005

+! 2%;70% (10.5ML) N020832 005 Apr 03, 2006

+! 2%;70% (3ML) N020832 007 Oct 10, 2006

SWAB; TOPICAL

CHLORAPREP ONE-STEP SEPP

+! BECTON DICKINSON CO 2%;70% (0.67ML) N021555 001 Oct 07, 2002

CHLORAPREP SINGLE SWABSTICK

+! BECTON DICKINSON CO 2%;70% (1.75ML) N021555 002 May 10, 2005

CHLORAPREP TRIPLE SWABSTICK

+! BECTON DICKINSON CO 2%;70% (5.25ML) N021555 003 Jun 10, 2009

PREVANTICS MAXI SWABSTICK

+! PROF DSPLS 3.15%;70% (5.1ML) N021524 003 Jun 03, 2005

PREVANTICS SWAB

+! PROF DSPLS 3.15%;70% (1ML) N021524 001 Jun 03, 2005

PREVANTICS SWABSTICK

+! PROF DSPLS 3.15%;70% (1.6ML) N021524 002 Jun 03, 2005

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE; ORAL

CHLORPHENIRAMINE MALEATE

! AVANTHI INC 12MG A040829 001 May 13, 2009

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET; ORAL

ADVIL ALLERGY AND CONGESTION RELIEF

+! GLAXOSMITHKLINE 4MG;200MG;10MG N022113 001 Dec 21, 2011

ADVIL MULTI-SYMP TOM COLD & FLU

+! GLAXOSMITHKLINE 4MG;200MG;10MG N022113 002 Apr 28, 2017

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S ADVIL ALLERGY SINUS

+! GLAXOSMITHKLINE 1MG/5ML;100MG/5ML;15MG/5ML N021587 001 Feb 24, 2004

TABLET; ORAL

ADVIL ALLERGY SINUS

+! GLAXOSMITHKLINE 2MG;200MG;30MG N021441 001 Dec 19, 2002

CIMETIDINE

TABLET; ORAL

CIMETIDINE

APOTEX 100MG A074948 001 Jun 19, 1998

200MG A074948 002 Jul 26, 2002

L PERRIGO CO 200MG A075285 001 Oct 29, 1998

TAGAMET HB

+! MEDTECH PRODUCTS 200MG N020238 002 Aug 21, 1996

OTC DRUG PRODUCT LIST

CLOTRIMAZOLE

CREAM; VAGINAL

CLOTRIMAZOLE

!	P AND L	1%	A074165	001	Jul 16, 1993
	TARO	1%	A072641	001	Dec 04, 1995

TRIVAGIZOLE 3

	TARO	2%	N021143	001	Apr 12, 2000
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CROMOLYN SODIUM

SPRAY, METERED; NASAL

CROMOLYN SODIUM

!	BAUSCH AND LOMB	5.2MG/SPRAY	A075702	001	Jul 03, 2001
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DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE

	ACTAVIS LABS FL	30MG; 600MG	A091070	001	Aug 31, 2015
		60MG; 1.2GM	A091070	002	Aug 31, 2015
	AMNEAL PHARMS	30MG; 600MG	A209692	001	Nov 01, 2018
		60MG; 1.2GM	A209692	002	Nov 01, 2018
	AUROBINDO PHARMA	30MG; 600MG	A206941	001	Mar 17, 2017
		60MG; 1.2GM	A206941	002	Mar 17, 2017
	DR REDDYS	30MG; 600MG	A217340	001	Aug 01, 2023
		60MG; 1.2GM	A217340	002	Aug 01, 2023
	PERRIGO R AND D	30MG; 600MG	A207602	002	Mar 05, 2018
		60MG; 1.2GM	A207602	001	Mar 05, 2018
	SUN PHARM	30MG; 600MG	A214781	001	Jul 01, 2021
		60MG; 1.2GM	A214781	002	Jul 01, 2021

MUCINEX DM

+	RB HLTH	30MG; 600MG	N021620	002	Apr 29, 2004
+	!	60MG; 1.2GM	N021620	001	Apr 29, 2004

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

DELSYM

+	!	RB HLTH	EQ 30MG HYDROBROMIDE/5ML	N018658	001	Oct 08, 1982
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DEXTROMETHORPHAN POLISTIREX

	AMNEAL	EQ 30MG HYDROBROMIDE/5ML	A203133	001	Jul 28, 2017
	TRIS PHARMA INC	EQ 30MG HYDROBROMIDE/5ML	A091135	001	May 25, 2012

DICLOFENAC SODIUM

GEL; TOPICAL

DICLOFENAC SODIUM

	AMNEAL PHARMS	1%	A208077	001	Mar 18, 2016
	AUROLIFE PHARMA LLC	1%	A204306	001	May 06, 2019
	ENCUBE	1%	A210986	001	Jan 27, 2020
	HIKMA	1%	A209484	001	Nov 21, 2018
	PERRIGO PHARMA INTL	1%	A211253	001	May 16, 2019

VOLTAREN ARTHRITIS PAIN

+	!	GLAXOSMITHKLINE	1%	N022122	001	Oct 17, 2007
		CONS				

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET; ORAL

ADVIL PM

+	!	GLAXOSMITHKLINE	38MG; 200MG	N021394	001	Dec 21, 2005
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IBUPROFEN AND DIPHENHYDRAMINE CITRATE

	AUROBINDO PHARMA	38MG; 200MG	A216204	001	May 31, 2022
	DR REDDYS LABS LTD	38MG; 200MG	A090619	001	Jul 08, 2009
	PERRIGO R AND D	38MG; 200MG	A079113	001	Dec 22, 2008

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN

CAPSULE; ORAL

ADVIL PM

+	!	GLAXOSMITHKLINE	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	N021393	001	Dec 21, 2005
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IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE

	AUROBINDO PHARMA	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	A210676	001	Feb 14, 2019
	LTD	SALT			
	BIONPHARMA	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	A090397	001	Nov 22, 2010
	STRIDES PHARMA	25MG; EQ 200MG FREE ACID AND POTASSIUM SALT	A200888	001	Mar 05, 2012

OTC DRUG PRODUCT LIST

DIPHENHYDRAMINE HYDROCHLORIDE; NAPROXEN SODIUM

TABLET; ORAL

ALEVE PM

+! BAYER HLTHCARE 25MG;220MG N205352 001 Jan 17, 2014

NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE

AMNEAL PHARMS CO 25MG;220MG A209726 001 Oct 23, 2018

COREPHARMA 25MG;220MG A211830 001 Aug 22, 2019

GRANULES 25MG;220MG A213663 001 Sep 24, 2020

PERRIGO R AND D 25MG;220MG A208499 001 May 10, 2019

DOCOSANOL

CREAM; TOPICAL

ABREVA

+! GLAXOSMITHKLINE 10% N020941 001 Jul 25, 2000

DOCOSANOL

ALEMBIC 10% A215839 001 May 03, 2022

ENCUBE 10% A212385 001 Oct 07, 2022

P AND L 10% A208754 001 Nov 19, 2018

TARO 10% A214454 001 Oct 26, 2023

DOXYLAMINE SUCCINATE

TABLET; ORAL

DOXYLAMINE SUCCINATE

LNK 25MG A040564 001 Aug 27, 2004

PERRIGO 25MG A040167 001 Sep 18, 1996

UNISOM

+! CHATTEM 25MG N018066 001

EPINEPHRINE

AEROSOL, METERED; INHALATION

PRIMATENE MIST

+! ARMSTRONG PHARMS 0.125MG/INH N205920 001 Nov 07, 2018

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

AMNEAL PHARMS NY EQ 20MG BASE A209716 001 Jun 05, 2019

ANDA REPOSITORY EQ 20MG BASE A216149 001 Mar 15, 2023

AUROBINDO PHARMA EQ 20MG BASE A209339 001 Oct 16, 2017

DR REDDYS EQ 20MG BASE A207673 001 May 15, 2018

GRAVITI PHARMS EQ 20MG BASE A216349 001 Jun 24, 2022

HETERO LABS LTD III EQ 20MG BASE A212507 001 Jun 02, 2020

MARKSANS PHARMA EQ 20MG BASE A217264 001 Sep 29, 2023

PERRIGO R AND D EQ 20MG BASE A207193 001 Aug 18, 2017

SUN PHARM EQ 20MG BASE A212866 001 May 04, 2020

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N204655 001 Mar 28, 2014

TABLET, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA EQ 20MG BASE A214473 001 Jul 12, 2023

LTD EQ 20MG BASE A211571 001 May 14, 2020

DR REDDYS EQ 20MG BASE A212088 001 Jun 25, 2020

MYLAN EQ 20MG BASE A212088 001 Jun 25, 2020

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N207920 001 Nov 23, 2015

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

ANNORA PHARMA 10MG A215766 001 Nov 08, 2021

20MG A215766 002 Nov 08, 2021

ASCENT PHARMS INC 10MG A216030 001 Nov 03, 2021

20MG A216030 002 Nov 03, 2021

AUROBINDO PHARMA 10MG A206531 001 Apr 26, 2016

20MG A206531 002 Apr 26, 2016

DR REDDYS LABS LTD 10MG A077367 002 Aug 17, 2001

20MG A077367 001 Sep 25, 2006

GLENMARK PHARMS INC 10MG A077146 001 Mar 07, 2005

20MG A090837 001 Aug 04, 2010

MARKSANS PHARMA 10MG A217543 001 Mar 08, 2023

20MG A217543 002 Mar 08, 2023

P AND L 10MG A075512 001 Jul 26, 2001

PERRIGO 10MG A075400 001 Mar 18, 2005

PERRIGO R AND D 20MG A077351 001 Sep 25, 2006

SUN PHARM INDS LTD 10MG A090283 001 Nov 17, 2009

OTC DRUG PRODUCT LIST

FAMOTIDINE

TABLET; ORAL
FAMOTIDINE

	20MG	A090283 002	Nov 17, 2009
VKT PHARMA	10MG	A215822 001	Jan 28, 2022
	20MG	A215822 002	Jan 28, 2022
PEPCID AC			
+ J AND J CONSUMER INC	10MG	N020325 001	Apr 28, 1995
	10MG	N020902 001	Aug 05, 1999
+	20MG	N020325 002	Sep 23, 2003

TABLET, CHEWABLE; ORAL

PEPCID AC			
+	20MG	N020801 002	Dec 17, 2007

FEXOFENADINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S ALLEGRA ALLERGY			
+	30MG/5ML	N201373 001	Jan 24, 2011
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY			
P AND L	30MG/5ML	A203330 001	Nov 18, 2014
TARO	30MG/5ML	A208123 001	Nov 09, 2017
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES			
!	30MG/5ML	A203330 002	Nov 18, 2014
TARO	30MG/5ML	A208123 002	Nov 09, 2017
FEXOFENADINE HYDROCHLORIDE			
AUROBINDO PHARMA LTD	30MG/5ML	A213466 001	May 23, 2023

TABLET; ORAL

ALLEGRA ALLERGY			
+	60MG	N020872 007	Jan 24, 2011
+	180MG	N020872 010	Jan 24, 2011
ALLEGRA HIVES			
+	180MG	N020872 009	Jan 24, 2011
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY			
AUROBINDO PHARMA LTD	30MG	A202039 001	Nov 19, 2014
DR REDDYS LABS LTD	30MG	A076502 004	Apr 12, 2011
HETERO LABS LTD V	30MG	A204097 001	Aug 19, 2016
TEVA	30MG	A076447 004	Apr 13, 2011
WOCKHARDT	30MG	A079112 002	Feb 08, 2012
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES			
DR REDDYS LABS LTD	30MG	A076502 005	Apr 12, 2011
TEVA	30MG	A076447 005	Apr 13, 2011
WOCKHARDT	30MG	A079112 001	Feb 08, 2012
FEXOFENADINE HYDROCHLORIDE			
L PERRIGO CO	60MG	A212971 001	Feb 24, 2020
	180MG	A212971 002	Feb 24, 2020
FEXOFENADINE HYDROCHLORIDE ALLERGY			
AUROBINDO PHARMA LTD	60MG	A202039 002	Nov 19, 2014
	180MG	A202039 003	Nov 19, 2014
DR REDDYS LABS LTD	60MG	A076502 006	Apr 12, 2011
	180MG	A076502 008	Apr 12, 2011
GRANULES	60MG	A211075 001	Oct 18, 2019
	180MG	A211075 002	Oct 18, 2019
HETERO LABS LTD V	60MG	A204097 002	Aug 19, 2016
	180MG	A204097 003	Aug 19, 2016
RISING	60MG	A077081 006	Jul 21, 2011
	180MG	A077081 008	Jul 21, 2011
SCIEGEN PHARMS INC	60MG	A204507 002	Sep 16, 2015
	180MG	A204507 003	Sep 16, 2015
SUN PHARM INDS	180MG	A091567 006	Feb 06, 2012
TEVA	60MG	A076447 006	Apr 13, 2011
	180MG	A076447 008	Apr 13, 2011
UNIQUE	180MG	A210137 001	Aug 13, 2018
WOCKHARDT	60MG	A079112 004	Feb 08, 2012
	180MG	A079112 006	Feb 08, 2012
FEXOFENADINE HYDROCHLORIDE HIVES			
DR REDDYS LABS LTD	60MG	A076502 007	Apr 12, 2011
	180MG	A076502 009	Apr 12, 2011
RISING	60MG	A077081 007	Jul 21, 2011
	180MG	A077081 009	Jul 21, 2011

OTC DRUG PRODUCT LIST

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

FEXOFENADINE HYDROCHLORIDE HIVES				
SCIEGEN PHARMS INC	60MG	A204507	004	Sep 16, 2015
	180MG	A204507	005	Sep 16, 2015
SUN PHARM INDS	180MG	A091567	005	Feb 06, 2012
TEVA	60MG	A076447	007	Apr 13, 2011
WOCKHARDT	60MG	A079112	003	Feb 08, 2012
	180MG	A079112	005	Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION				
+! CHATTEM SANOFI	60MG;120MG	N020786	002	Jan 24, 2011
ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION				
+! CHATTEM SANOFI	180MG;240MG	N021704	002	Jan 24, 2011
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE				
AUROBINDO PHARMA	60MG;120MG	A209116	001	Oct 30, 2017
DR REDDYS	60MG;120MG	A076667	001	Nov 18, 2014
	60MG;120MG	A215434	001	May 31, 2022
DR REDDYS LABS LTD	180MG;240MG	A079043	002	Jun 22, 2011
SUN PHARM	60MG;120MG	A090818	001	Jan 29, 2015

FLUTICASONE FUROATE

SPRAY, METERED; NASAL

FLONASE SENSIMIST ALLERGY RELIEF				
+! GLAXOSMITHKLINE	0.0275MG/SPRAY	N022051	002	Aug 02, 2016
CONS				

FLUTICASONE PROPIONATE

SPRAY, METERED; NASAL

FLONASE ALLERGY RELIEF				
+ GLAXOSMITHKLINE	0.05MG/SPRAY	N205434	001	Jul 23, 2014
CONS				
FLUTICASONE PROPIONATE				
! APOTEX	0.05MG/SPRAY	A208150	001	Feb 29, 2016
! HIKMA	0.05MG/SPRAY	A207957	001	May 26, 2016
	0.05MG/SPRAY	A208024	001	Apr 17, 2019

GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

GUAIFENESIN				
ACTAVIS LABS FL	1.2GM	A091009	002	Sep 03, 2015
AMNEAL PHARMS	600MG	A207342	001	Jul 11, 2018
	1.2GM	A207342	002	Jul 11, 2018
AUROBINDO PHARMA	600MG	A210453	001	Oct 21, 2019
	1.2GM	A210453	002	Oct 21, 2019
DR REDDYS	600MG	A215932	001	Mar 15, 2022
	1.2GM	A215932	002	Mar 15, 2022
GRANULES	600MG	A213420	001	May 08, 2020
	1.2GM	A213420	002	May 08, 2020
GUARDIAN DRUG	600MG	A209215	001	Sep 06, 2017
	1.2GM	A209215	002	Sep 06, 2017
MARKSANS PHARMA	600MG	A217780	001	Aug 21, 2023
	1.2GM	A217780	002	Aug 21, 2023
OHM LABS INC	600MG	A209254	001	Jul 16, 2018
	1.2GM	A209254	002	Jul 16, 2018
PERRIGO R AND D	600MG	A078912	001	Nov 23, 2011
	1.2GM	A078912	002	Nov 05, 2020
MUCINEX				
+ RB HLTH	600MG	N021282	001	Jul 12, 2002
+!	1.2GM	N021282	002	Dec 18, 2002

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE				
ACTAVIS LABS FL	600MG;60MG	A091071	001	May 27, 2015
	1.2GM;120MG	A091071	002	May 27, 2015
AUROBINDO PHARMA	600MG;60MG	A213203	001	Mar 25, 2020
LTD				
	1.2GM;120MG	A213203	002	Mar 25, 2020
DR REDDYS	600MG;60MG	A208369	001	Dec 29, 2017
	1.2GM;120MG	A208369	002	Dec 29, 2017
L PERRIGO CO	600MG;60MG	A214407	001	Feb 01, 2022
	1.2GM;120MG	A214407	002	Feb 01, 2022

OTC DRUG PRODUCT LIST

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE

SUN PHARM INDS INC 600MG;60MG
1.2GM;120MGA212542 001 Apr 28, 2020
A212542 002 Apr 28, 2020

MUCINEX D

+ RB HLTH 600MG;60MG
+! 1.2GM;120MGN021585 001 Jun 22, 2004
N021585 002 Jun 22, 2004IBUPROFEN

CAPSULE;ORAL

ADVIL LIQUI-GELS

+! GLAXOSMITHKLINE EQ 200MG FREE ACID AND POTASSIUM SALT

N020402 001 Apr 20, 1995

ADVIL MIGRAINE LIQUI-GELS

+! GLAXOSMITHKLINE EQ 200MG FREE ACID AND POTASSIUM SALT

N020402 002 Mar 16, 2000

IBUPROFEN

AMNEAL PHARMS EQ 200MG FREE ACID AND POTASSIUM SALT

A202300 001 Dec 23, 2011

ASCENT PHARMS INC EQ 200MG FREE ACID AND POTASSIUM SALT

A206999 001 Dec 21, 2017

AUROBINDO PHARMA EQ 200MG FREE ACID AND POTASSIUM SALT

A207753 001 Jun 29, 2018

LTD

EQ 200MG FREE ACID AND POTASSIUM SALT

A215777 001 Sep 14, 2023

BIONPHARMA EQ 200MG FREE ACID AND POTASSIUM SALT

A078682 001 Mar 24, 2009

HUMANWELL PURACAP EQ 200MG FREE ACID AND POTASSIUM SALT

A206568 001 Jun 21, 2016

MARKSANS PHARMA EQ 200MG FREE ACID AND POTASSIUM SALT

A079205 001 Jun 26, 2009

P AND L DEV LLC EQ 200MG FREE ACID AND POTASSIUM SALT

A077338 001 Jul 10, 2009

SOFGEN PHARMS EQ 200MG FREE ACID AND POTASSIUM SALT

A203599 001 Sep 07, 2016

STRIDES PHARMA EQ 200MG FREE ACID AND POTASSIUM SALT

A204469 001 Mar 28, 2018

MIDOL LIQUID GELS

+! BIONPHARMA 200MG

N021472 001 Oct 18, 2002

SUSPENSION;ORAL

CHILDREN'S ADVIL

GLAXOSMITHKLINE 100MG/5ML

N020589 001 Jun 27, 1996

CHILDREN'S ADVIL-FLAVORED

GLAXOSMITHKLINE 100MG/5ML

N020589 002 Nov 07, 1997

CHILDREN'S IBUPROFEN

PERRIGO 100MG/5ML

A074937 001 Dec 22, 1998

CHILDREN'S MOTRIN

+! J AND J CONSUMER 100MG/5ML

N020516 001 Jun 16, 1995

INC

IBUPROFEN

APTAPHARMA INC 100MG/5ML

A210602 001 Nov 23, 2018

AUROBINDO PHARMA 100MG/5ML

A209179 001 Apr 17, 2018

LTD

GUARDIAN DRUG 100MG/5ML

A210149 001 Aug 17, 2018

P AND L 100MG/5ML

A074916 001 Apr 30, 1999

TARO 100MG/5ML

A209207 001 Jun 27, 2017

SUSPENSION/DROPS;ORAL

CHILDREN'S MOTRIN

+! J AND J CONSUMER 40MG/ML

N020603 001 Jun 10, 1996

INC

IBUPROFEN

GUARDIAN DRUG 40MG/ML

A210755 001 Sep 26, 2018

L PERRIGO CO 40MG/ML

A075217 001 Dec 16, 1998

TARO 40MG/ML

A217261 001 Aug 08, 2023

TRIS PHARMA INC 40MG/ML

A079058 001 Aug 31, 2009

INFANT'S ADVIL

+! GLAXOSMITHKLINE 50MG/1.25ML

N020812 002 Jan 12, 2000

TABLET;ORAL

ADVIL

+! GLAXOSMITHKLINE 200MG

N018989 001 May 18, 1984

IBUPROFEN

AMNEAL PHARMS 200MG

A079233 001 Mar 18, 2014

AMNEAL PHARMS NY 200MG

A071333 001 Feb 17, 1987

200MG

A072199 001 May 23, 1988

AUROBINDO PHARMA 200MG

A208865 001 Nov 08, 2017

AVEMA PHARMA 200MG

A076460 001 Nov 26, 2003

CONTRACT PHARMACAL 200MG

A072299 001 Jul 01, 1988

DR REDDYS LA 200MG

A075661 001 Dec 12, 2001

DR REDDYS LABS INC 200MG

A076117 001 Nov 20, 2001

GRANULES 200MG

A202312 001 Oct 07, 2016

GRANULES INDIA 200MG

A079174 001 Dec 10, 2010

LNK 200MG

A075010 001 Mar 01, 1999

200MG

A075139 001 Mar 01, 1999

MARKSANS PHARMA 200MG

A091237 001 Feb 08, 2011

OTC DRUG PRODUCT LIST

IBUPROFEN

TABLET; ORAL

IBUPROFEN

	200MG	A091239 001	Feb 01, 2011
MCNEIL	200MG	A073019 001	Mar 30, 1994
PERRIGO	200MG	A072096 001	Dec 08, 1987
! PERRIGO R AND D	200MG	A077349 001	Jun 21, 2005
SHANDONG XINHUA	200MG	A207095 001	May 05, 2017
	200MG	A207095 002	Aug 21, 2018
STRIDES PHARMA	200MG	A070481 001	Sep 24, 1986
	200MG	A079129 001	Mar 28, 2011
	200MG	A091355 001	Apr 04, 2011
	200MG	A206989 001	Jun 29, 2018
	200MG	A207052 001	May 30, 2017
VINTAGE PHARMS	200MG	A071229 001	Apr 01, 1987
JUNIOR STRENGTH ADVIL			
GLAXOSMITHKLINE	100MG	N020267 002	Dec 13, 1996
JUNIOR STRENGTH MOTRIN			
J AND J CONSUMER	100MG	N020602 001	Jun 10, 1996
INC			
MOTRIN IB			
+ J AND J CONSUMER	200MG	N019012 003	Dec 17, 1990
INC			

TABLET, CHEWABLE; ORAL

CHILDREN'S ADVIL

GLAXOSMITHKLINE	50MG	N020944 001	Dec 18, 1998
IBUPROFEN			
! PERRIGO	100MG	A076359 002	Jan 16, 2004
JUNIOR STRENGTH ADVIL			
GLAXOSMITHKLINE	100MG	N020944 002	Dec 18, 1998

IBUPROFEN SODIUM

TABLET; ORAL

ADVIL

+! GLAXOSMITHKLINE	EQ 200MG BASE	N201803 001	Jun 12, 2012
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IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET; ORAL

ADVIL CONGESTION RELIEF

+! GLAXOSMITHKLINE	200MG; 10MG	N022565 001	May 27, 2010
IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE			
PERRIGO R AND D	200MG; 10MG	A203200 001	Jul 03, 2014

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

ADVIL COLD AND SINUS

+! GLAXOSMITHKLINE	EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG	N021374 001	May 30, 2002
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE			
AUROBINDO PHARMA	EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG	A209235 001	Dec 01, 2017

SUSPENSION; ORAL

CHILDREN'S ADVIL COLD

GLAXOSMITHKLINE	100MG/5ML; 15MG/5ML	N021373 001	Apr 18, 2002
CHILDREN'S MOTRIN COLD			
+! J AND J CONSUMER	100MG/5ML; 15MG/5ML	N021128 001	Aug 01, 2000
INC			

TABLET; ORAL

ADVIL COLD AND SINUS

+! GLAXOSMITHKLINE	200MG; 30MG	N019771 001	Sep 19, 1989
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE			
DR REDDYS LABS LTD	200MG; 30MG	A077628 001	Aug 14, 2006
IBUPROHM COLD AND SINUS			
OHM LABS	200MG; 30MG	A074567 001	Apr 17, 2001
SINE-AID IB			
J AND J CONSUMER	200MG; 30MG	N019899 001	Dec 31, 1992
INC			

IODINE POVACRYLEX; ISOPROPYL ALCOHOL

SPONGE; TOPICAL

DURAPREP

+! 3M	EQ 0.7% IODINE; 74% (6ML)	N021586 001	Sep 29, 2006
+!	EQ 0.7% IODINE; 74% (26ML)	N021586 002	Sep 29, 2006

OTC DRUG PRODUCT LIST

ISOPROPYL ALCOHOL

SOLUTION;TOPICAL

ZURAGARD

+! ZUREX PHARMA 70% N210872 001 Apr 26, 2019

IVERMECTIN

LOTION;TOPICAL

IVERMECTIN

TARO 0.5% A210720 001 May 06, 2020

SKLICE

+! ARBOR PHARMS LLC 0.5% N202736 001 Feb 07, 2012

KETOCONAZOLE

SHAMPOO;TOPICAL

NIZORAL ANTI-DANDRUFF

+! KRAMER 1% N020310 001 Oct 10, 1997

KETOTIFEN FUMARATE

SOLUTION/DROPS;OPHTHALMIC

ALAWAY

+! BAUSCH AND LOMB EQ 0.025% BASE N021996 001 Dec 01, 2006

BAUSCH AND LOMB INC EQ 0.025% BASE A208158 001 Sep 24, 2020

CHILDREN'S ALAWAY

+ BAUSCH AND LOMB EQ 0.025% BASE N021996 002 Feb 11, 2015

KETOTIFEN FUMARATE

APOTEX INC EQ 0.025% BASE A077354 001 May 09, 2006

BAYSHORE PHARMS LLC EQ 0.025% BASE A204059 001 Jun 01, 2020

SENTISS EQ 0.025% BASE A077958 001 Jul 26, 2007

ZADITOR

! ALCON PHARMS LTD EQ 0.025% BASE A077200 001 Sep 02, 2008

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

DR REDDYS LABS LTD 15MG A202194 001 May 18, 2012

GLENMARK PHARMS INC 15MG A202727 001 May 18, 2012

NATCO 15MG A203306 001 Jan 13, 2016

PERRIGO R AND D 15MG A202319 001 May 18, 2012

PREVACID 24 HR

+! PERRIGO PHARMA INTL 15MG N022327 001 May 18, 2009

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

DEXCEL 15MG N208025 001 Jun 07, 2016

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

XYZAL ALLERGY 24HR

+! CHATTEM SANOFI 2.5MG/5ML N209090 001 Jan 31, 2017

TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX 5MG A211443 001 Apr 21, 2021

DR REDDYS 5MG A210375 001 Jan 19, 2018

HETERO LABS LTD III 5MG A213513 001 Apr 29, 2020

MICRO LABS 5MG A211551 001 Nov 20, 2018

PERRIGO R AND D 5MG A211983 001 Mar 28, 2019

XYZAL ALLERGY 24HR

+! CHATTEM SANOFI 5MG N209089 001 Jan 31, 2017

LEVONORGESTREL

TABLET;ORAL

ATHENTIA NEXT

AUROBINDO PHARMA 1.5MG A206867 001 Dec 08, 2015

FALLBACK SOLO

LUPIN LTD 1.5MG A201446 001 Jun 19, 2014

HER STYLE

NOVAST LABS 1.5MG A207976 001 Mar 11, 2016

LEVONORGESTREL

GLENMARK PHARMS LTD 1.5MG A207044 001 Mar 25, 2016

LABORATOIRE HRA 1.5MG A204044 001 Jul 03, 2018

MYLAN LABS LTD 1.5MG A202739 001 Oct 31, 2014

NAARI PTE LTD 1.5MG A202380 001 May 29, 2015

NOVEL LABS INC 1.5MG A202508 001 Feb 22, 2013

PERRIGO R AND D 1.5MG A202334 001 Aug 20, 2014

XIROMED 1.5MG A205329 001 Sep 18, 2018

OTC DRUG PRODUCT LIST

LEVONORGESTREL

TABLET; ORAL

OPCICON ONE-STEP

SUN PHARM

1.5MG

A202635 001 Sep 11, 2014

PLAN B ONE-STEP

+! FDN CONSUMER

1.5MG

N021998 001 Jul 10, 2009

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

LOPERAMIDE HYDROCHLORIDE

+ BIONPHARMA

1MG

N021855 001 Aug 04, 2005

+!

2MG

N021855 002 Aug 04, 2005

STRIDES PHARMA

2MG

A213070 001 Aug 11, 2021

SOLUTION; ORAL

IMODIUM A-D

+! J AND J CONSUMER

1MG/5ML

N019487 001 Mar 01, 1988

INC

LOPERAMIDE HYDROCHLORIDE

WOCKHARDT BIO AG

1MG/5ML

A074730 001 Aug 28, 1997

SUSPENSION; ORAL

IMODIUM A-D

+! J AND J CONSUMER

1MG/7.5ML

N019487 002 Jul 08, 2004

INC

LOPERAMIDE HYDROCHLORIDE

PERRIGO R AND D

1MG/7.5ML

A091292 001 May 20, 2011

TABLET; ORAL

IMODIUM A-D

+! J AND J CONSUMER

2MG

N019860 001 Nov 22, 1989

INC

LOPERAMIDE HYDROCHLORIDE

AUROBINDO PHARMA

2MG

A206548 001 Dec 15, 2015

L PERRIGO CO

2MG

A075232 001 Jan 06, 2000

LNK

2MG

A076497 001 Jun 10, 2003

OHM LABS

2MG

A074091 001 Dec 10, 1992

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET; ORAL

IMODIUM MULTI-SYMPOM RELIEF

+! J AND J CONSUMER

2MG;125MG

N021140 001 Nov 30, 2000

INC

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

AUROBINDO PHARMA

2MG;125MG

A211059 001 Dec 14, 2020

LTD

BIONPHARMA

2MG;125MG

A213484 001 Sep 10, 2021

GUARDIAN DRUG

2MG;125MG

A214541 001 May 27, 2021

HETERO LABS LTD V

2MG;125MG

A211438 001 Jun 17, 2021

PERRIGO R AND D

2MG;125MG

A209837 001 Sep 05, 2018

SUN PHARM INDS LTD

2MG;125MG

A077500 001 Sep 06, 2006

LORATADINE

CAPSULE; ORAL

CLARITIN

+! BAYER HEALTHCARE

10MG

N021952 001 Jun 16, 2008

LLC

LORATADINE

BIONPHARMA

10MG

A202538 001 Dec 21, 2018

MARKSANS PHARMA

10MG

A206214 001 Sep 23, 2016

SUSPENSION; ORAL

LORATADINE

+! TARO

1MG/ML

N021734 001 Oct 04, 2005

SYRUP; ORAL

CLARITIN

+! BAYER HEALTHCARE

1MG/ML

N020641 002 Nov 27, 2002

LLC

LORATADINE

AUROBINDO PHARMA

1MG/ML

A208931 001 Jun 29, 2018

LTD

HETERO LABS LTD III

1MG/ML

A210409 001 May 07, 2021

LANNETT CO INC

1MG/ML

A077421 001 Jun 29, 2006

PERRIGO

1MG/ML

A075728 001 Aug 20, 2004

TARO

1MG/ML

A076805 001 Aug 20, 2004

TARO

1MG/ML

A201865 001 Jul 31, 2015

WOCKHARDT BIO AG

1MG/ML

A075815 001 Aug 20, 2004

OTC DRUG PRODUCT LIST

LORATADINE

TABLET; ORAL

CLARITIN

+	!	BAYER HEALTHCARE LLC	10MG	N019658 002	Nov 27, 2002
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CLARITIN HIVES RELIEF

+	!	BAYER HEALTHCARE LLC	10MG	N019658 003	Nov 19, 2003
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LORATADINE

		APOTEX INC	10MG	A076471 001	Feb 14, 2006
		AUROBINDO PHARMA LTD	10MG	A208314 001	Apr 16, 2018
		GRANULES	10MG	A210722 001	Dec 23, 2019
		GUARDIAN DRUG	10MG	A207569 001	Mar 12, 2019
		HETERO LABS LTD V	10MG	A211718 001	Jul 28, 2023
		MYLAN	10MG	A076154 001	Aug 20, 2003
		PERRIGO	10MG	A076301 001	Jun 25, 2004
		PLD ACQUISITIONS LLC	10MG	A075209 001	Jan 21, 2003
		SUN PHARM INDS LTD	10MG	A076134 001	Aug 18, 2003
		UNIQUE PHARM	10MG	A214684 001	Jan 07, 2021

TABLET, CHEWABLE; ORAL

CHILDREN'S CLARITIN

+		BAYER HEALTHCARE LLC	5MG	N021891 001	Aug 23, 2006
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CLARITIN

+	!	BAYER HEALTHCARE LLC	10MG	N021891 002	Nov 21, 2018
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LORATADINE

		PERRIGO PHARMA INTL	5MG	A210033 001	Jun 12, 2019
		SUN PHARM	5MG	A210088 001	Apr 16, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

ALAVERT

		FDN CONSUMER	10MG	N021375 001	Dec 19, 2002
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CLARITIN HIVES RELIEF REDITAB

+	!	BAYER HEALTHCARE LLC	10MG	N020704 003	Nov 19, 2003
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CLARITIN REDITABS

+	!	BAYER HEALTHCARE LLC	5MG	N021993 001	Dec 12, 2006
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+	!		10MG	N020704 002	Nov 27, 2002
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LORATADINE

		AUROBINDO PHARMA LTD	10MG	A208477 001	Apr 11, 2018
		PERRIGO PHARMA INTL	10MG	A076011 001	Sep 29, 2003
		RUBICON	10MG	A214280 001	Sep 10, 2020
		TENSHI	5MG	A212795 001	Sep 18, 2020
			10MG	A213294 001	Oct 30, 2020

LORATADINE REDIDOSE

		SUN PHARM INDS LTD	10MG	A077153 001	Apr 11, 2007
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LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARITIN-D

+	!	BAYER HEALTHCARE LLC	5MG;120MG	N019670 002	Nov 27, 2002
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CLARITIN-D 24 HOUR

+	!	BAYER HEALTHCARE LLC	10MG;240MG	N020470 002	Nov 27, 2002
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LORATADINE AND PSEUDOEPHEDRINE SULFATE

		P AND L	10MG;240MG	A075706 001	Feb 21, 2003
		PERRIGO PHARMA INTL	5MG;120MG	A076050 001	Jan 30, 2003
			10MG;240MG	A075989 001	Mar 04, 2004
		SUN PHARM INDS LTD	10MG;240MG	A076557 001	Sep 22, 2004

MENTHOL; METHYL SALICYLATE

PATCH; TOPICAL

SALONPAS

+	!	HISAMITSU PHARM CO	3%;10%	N022029 001	Feb 20, 2008
+			3%;10%	N022029 002	Nov 05, 2012

OTC DRUG PRODUCT LIST

MICONAZOLE NITRATE

CREAM;TOPICAL, VAGINAL

MICONAZOLE 3 COMBINATION PACK

PERRIGO 2%,4%

A076357 001 Mar 30, 2004

MONISTAT 3 COMBINATION PACK

+ MEDTECH PRODUCTS 2%,4%

N021261 003 Jun 17, 2003

MONISTAT 3 COMBINATION PACK (PREFILLED)

+! MEDTECH PRODUCTS 2%,4%

N021261 001 Feb 02, 2001

CREAM;VAGINAL

MICONAZOLE 3

TARO 4%

A076773 001 Mar 02, 2005

MICONAZOLE 7

P AND L 2%

A074164 001 Mar 29, 1996

MICONAZOLE NITRATE

COSETTE 2%

A074366 001 Feb 22, 1996

PERRIGO 2%

A074760 001 May 15, 1997

PERRIGO R AND D 4%

A091366 001 Jan 15, 2010

TARO 2%

A074444 001 Jan 13, 1997

MONISTAT 3

+! MEDTECH PRODUCTS 4%

N020827 001 Mar 30, 1998

MONISTAT 7

+! MEDTECH PRODUCTS 2%

N017450 002 Feb 15, 1991

CREAM, INSERT;TOPICAL, VAGINAL

MICONAZOLE NITRATE

PERRIGO R AND D 2%,1.2GM

A079114 001 Jun 02, 2010

MONISTAT 1 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,1.2GM

N021308 001 Jun 29, 2001

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

M-ZOLE 3 COMBINATION PACK

P AND L 2%,200MG

A074926 001 Apr 16, 1999

MICONAZOLE NITRATE COMBINATION PACK

L PERRIGO CO 2%,200MG

A075329 001 Apr 20, 1999

MONISTAT 3 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,200MG

N020670 002 Apr 16, 1996

MONISTAT 7 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,100MG

N020288 002 Apr 26, 1993

SUPPOSITORY;VAGINAL

MICONAZOLE NITRATE

COSETTE 100MG

A074414 001 Apr 30, 1997

P AND L 100MG

A073507 001 Nov 19, 1993

MONISTAT 7

+! MEDTECH PRODUCTS 100MG

N018520 002 Feb 15, 1991

MINOXIDIL

AEROSOL, FOAM;TOPICAL

MEN'S ROGAINE

+! JOHNSON AND JOHNSON 5%

N021812 001 Jan 20, 2006

MINOXIDIL

PERRIGO PHARMA INTL 5%

A091344 001 Apr 28, 2011

MINOXIDIL (FOR MEN)

P AND L 5%

A208092 001 Feb 17, 2017

TARO 5%

A209074 001 Dec 31, 2018

MINOXIDIL (FOR WOMEN)

P AND L 5%

A208092 002 Jul 27, 2017

TARO 5%

A209074 002 Apr 22, 2019

WOMEN'S ROGAINE

+! JOHNSON AND JOHNSON 5%

N021812 002 Feb 28, 2014

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

AUROBINDO PHARMA LTD 2%

A074767 001 Feb 28, 1997

L PERRIGO CO 2%

A075357 001 Jul 30, 1999

P AND L 2%

A074588 001 Apr 05, 1996

MINOXIDIL (FOR WOMEN)

L PERRIGO CO 2%

A075357 002 Jul 30, 1999

MINOXIDIL EXTRA STRENGTH (FOR MEN)

AUROBINDO PHARMA LTD 5%

A075438 001 Feb 27, 2003

P AND L 5%

A075518 001 Nov 17, 2000

PERRIGO 5%

A075598 001 Jun 13, 2001

ROGAINE (FOR MEN)

+! JOHNSON AND JOHNSON 2%

N019501 002 Feb 09, 1996

OTC DRUG PRODUCT LIST

MINOXIDIL

SOLUTION;TOPICAL

ROGAINE (FOR WOMEN)

+! JOHNSON AND JOHNSON 2% N019501 003 Feb 09, 1996

ROGAINE EXTRA STRENGTH (FOR MEN)

+! JOHNSON AND JOHNSON 5% N020834 001 Nov 14, 1997

THEROXIDIL

PURE SOURCE 2% A078176 001 Nov 09, 2007

5% A076239 001 Aug 24, 2004

MOMETASONE FUROATE

SPRAY, METERED;NASAL

NASONEX 24HR ALLERGY

+! PERRIGO PHARMA INTL 0.05MG/SPRAY N215712 001 Mar 17, 2022

NALOXONE HYDROCHLORIDE

SPRAY, METERED;NASAL

NALOXONE HYDROCHLORIDE

PADAGIS ISRAEL 4MG/SPRAY A211951 001 Jun 21, 2022

NARCAN

+! EMERGENT 4MG/SPRAY N208411 001 Nov 18, 2015

RIVIVE

+! HARM REDUCTION 3MG/SPRAY N217722 001 Jul 28, 2023

THERP

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC

NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE

ALTAIRE PHARMS INC 0.02675%;0.315% A078208 001 Sep 27, 2010

RISING 0.025%;0.3% A202795 001 Jan 24, 2013

NAPHCON-A

+! ALCON 0.025%;0.3% N020226 001 Jun 08, 1994

OPCON-A

+! BAUSCH AND LOMB 0.02675%;0.315% N020065 001 Jun 08, 1994

VISINE

+! JOHNSON AND JOHNSON 0.025%;0.3% N020485 001 Jan 31, 1996

NAPROXEN SODIUM

CAPSULE;ORAL

NAPROXEN SODIUM

+! BIONPHARMA EQ 200MG BASE N021920 001 Feb 17, 2006

CATALENT EQ 200MG BASE A202807 001 Jan 04, 2019

PATHEON SOFTGELS EQ 200MG BASE A214463 001 Jan 10, 2023

PURACAP PHARM LLC EQ 200MG BASE A208363 001 Mar 15, 2018

TABLET;ORAL

ALEVE

+! BAYER 220MG N020204 002 Jan 11, 1994

NAPROXEN SODIUM

AMNEAL PHARMS NY 220MG A079096 001 Dec 16, 2008

AUROBINDO PHARMA 220MG A205497 001 Mar 18, 2016

LTD

CONTRACT PHARMACAL 220MG A074635 001 Jan 13, 1997

DR REDDYS LABS INC 220MG A075168 001 Jul 28, 1998

GRANULES INDIA 220MG A091353 001 Sep 20, 2011

HETERO LABS LTD V 220MG A211065 001 Oct 28, 2022

LNK INTL INC 220MG A204872 001 Jan 23, 2017

MARKSANS PHARMA 220MG A090545 001 Mar 16, 2011

NOVELGENIX THERAPS 220MG A207612 001 Nov 16, 2018

PERRIGO 220MG A074661 001 Jan 13, 1997

SUN PHARM INDS LTD 220MG A091183 001 May 20, 2011

YICHANG HUMANWELL 220MG A212033 001 Aug 30, 2019

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

DR REDDYS LABS INC 220MG;120MG A077381 001 Sep 27, 2006

! PERRIGO 220MG;120MG A076518 001 Mar 17, 2004

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

HABITROL

+ DR REDDYS LABS SA 7MG/24HR N020076 004 Nov 12, 1999

+ 14MG/24HR N020076 005 Nov 12, 1999

+! 21MG/24HR N020076 006 Nov 12, 1999

OTC DRUG PRODUCT LIST

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NICODERM CQ

+ CHATTEM SANOFI	7MG/24HR	N020165 006	Aug 02, 1996
+	14MG/24HR	N020165 005	Aug 02, 1996
+!	21MG/24HR	N020165 004	Aug 02, 1996

NICOTINE

AVEVA	7MG/24HR	A074612 002	Jul 28, 2003
	14MG/24HR	A074612 003	Oct 20, 1997
	21MG/24HR	A074612 001	Oct 20, 1997

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICORETTE

+ GLAXOSMITHKLINE	EQ 2MG BASE	N018612 002	Feb 09, 1996
+	EQ 2MG BASE	N018612 004	Sep 25, 2000
+!	EQ 4MG BASE	N020066 002	Feb 09, 1996
+	EQ 4MG BASE	N020066 004	Sep 25, 2000

NICORETTE (MINT)

+ GLAXOSMITHKLINE	EQ 2MG BASE	N018612 003	Dec 23, 1998
+	EQ 4MG BASE	N020066 003	Dec 23, 1998

NICOTINE POLACRILEX

L PERRIGO CO	EQ 2MG BASE	A076775 001	Sep 16, 2004
	EQ 2MG BASE	A076777 001	Sep 16, 2004
	EQ 4MG BASE	A076779 001	Sep 16, 2004
	EQ 4MG BASE	A076789 001	Sep 16, 2004
P AND L	EQ 2MG BASE	A074507 001	Mar 15, 1999
	EQ 2MG BASE	A076569 001	Jul 29, 2004
	EQ 2MG BASE	A078699 001	Dec 29, 2008
	EQ 2MG BASE	A079044 001	Jul 08, 2009
	EQ 2MG BASE	A079216 001	Jul 08, 2009
	EQ 2MG BASE	A204794 001	May 10, 2016
	EQ 4MG BASE	A074707 001	Mar 19, 1999
	EQ 4MG BASE	A076568 002	Jul 29, 2004
	EQ 4MG BASE	A078697 001	Dec 29, 2008
	EQ 4MG BASE	A079038 001	Jul 08, 2009
	EQ 4MG BASE	A079219 001	Jul 08, 2009
	EQ 4MG BASE	A204833 001	Feb 26, 2016
PERRIGO R AND D	EQ 2MG BASE	A078325 001	Oct 30, 2006
	EQ 2MG BASE	A078547 001	May 24, 2007
	EQ 2MG BASE	A091349 001	Jul 20, 2011
	EQ 2MG BASE	A206394 001	Dec 15, 2016
	EQ 4MG BASE	A078326 001	Oct 30, 2006
	EQ 4MG BASE	A078546 001	May 24, 2007
	EQ 4MG BASE	A091354 001	Jul 20, 2011
	EQ 4MG BASE	A206393 001	Dec 15, 2016

TROCHE/LOZENGE;ORAL

NICORETTE

+ GLAXOSMITHKLINE	EQ 2MG BASE	N022360 001	May 18, 2009
+!	EQ 4MG BASE	N022360 002	May 18, 2009
+ GLAXOSMITHKLINE CONS	EQ 2MG BASE	N021330 001	Oct 31, 2002
+!	EQ 4MG BASE	N021330 002	Oct 31, 2002

NICOTINE POLACRILEX

AUROBINDO PHARMA	EQ 2MG BASE	A213266 001	Aug 03, 2021
	EQ 4MG BASE	A213266 002	Aug 03, 2021
AUROBINDO PHARMA LTD	EQ 2MG BASE	A215357 001	May 13, 2022
	EQ 4MG BASE	A215357 002	May 13, 2022
DR REDDYS LABS SA	EQ 2MG BASE	A212796 001	Jan 08, 2020
	EQ 2MG BASE	A212983 001	Feb 21, 2020
	EQ 2MG BASE	A213233 001	Aug 04, 2020
	EQ 4MG BASE	A212796 002	Jan 08, 2020
	EQ 4MG BASE	A212983 002	Feb 21, 2020
	EQ 4MG BASE	A213233 002	Aug 04, 2020
P AND L	EQ 2MG BASE	A208875 001	Oct 31, 2019
	EQ 2MG BASE	A209206 001	Jun 26, 2018
	EQ 2MG BASE	A209519 001	Jul 02, 2018
	EQ 2MG BASE	A209520 001	Oct 31, 2019
	EQ 2MG BASE	A210711 001	Oct 31, 2019
	EQ 2MG BASE	A210712 001	Sep 06, 2019
	EQ 2MG BASE	A212056 001	Jul 26, 2019
	EQ 2MG BASE	A212057 001	May 14, 2020

OTC DRUG PRODUCT LIST

NICOTINE POLACRILEX

TROCHE/LOZENGE;ORAL

NICOTINE POLACRILEX

	EQ 4MG BASE	A208875	002	Oct 31, 2019
	EQ 4MG BASE	A209206	002	Jun 26, 2018
	EQ 4MG BASE	A209519	002	Jul 02, 2018
	EQ 4MG BASE	A209520	002	Oct 31, 2019
	EQ 4MG BASE	A210711	002	Oct 31, 2019
	EQ 4MG BASE	A210712	002	Sep 06, 2019
	EQ 4MG BASE	A212056	002	Jul 26, 2019
	EQ 4MG BASE	A212057	002	May 14, 2020
PERRIGO R AND D	EQ 2MG BASE	A077007	001	Jan 31, 2006
	EQ 2MG BASE	A090711	001	Jul 10, 2009
	EQ 2MG BASE	A090821	001	Jul 10, 2009
	EQ 2MG BASE	A203690	001	Oct 09, 2012
	EQ 4MG BASE	A077007	002	Jan 31, 2006
	EQ 4MG BASE	A090711	002	Jul 10, 2009
	EQ 4MG BASE	A090821	002	Jul 10, 2009
	EQ 4MG BASE	A203690	002	Oct 09, 2012
PLD ACQUISITIONS	EQ 2MG BASE	A207868	001	Feb 07, 2019
	EQ 4MG BASE	A207868	002	Feb 07, 2019

NIZATIDINE

TABLET;ORAL

AXID AR

+! GLAXOSMITHKLINE 75MG N020555 001 May 09, 1996

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

ALEMBIC	EQ 0.1% BASE	A209919	001	Dec 07, 2018
	EQ 0.2% BASE	A209420	001	Apr 29, 2019
APOTEX	EQ 0.1% BASE	A078350	001	Dec 07, 2015
	EQ 0.2% BASE	A090918	001	Dec 05, 2017
BARR LABS INC	EQ 0.2% BASE	A090848	001	Jul 13, 2015
EUGIA PHARMA	EQ 0.1% BASE	A204812	001	Dec 18, 2015
	EQ 0.2% BASE	A209995	001	Apr 04, 2019
GLAND PHARMA LTD	EQ 0.1% BASE	A209619	001	Aug 02, 2019
	EQ 0.2% BASE	A209752	001	May 20, 2020
USV	EQ 0.1% BASE	A203152	001	Dec 07, 2015
PATADAY ONCE DAILY RELIEF				
+! ALCON LABS INC	EQ 0.2% BASE	N021545	001	Dec 22, 2004
+!	EQ 0.7% BASE	N206276	001	Jan 30, 2015
PATADAY TWICE DAILY RELIEF				
+! ALCON LABS INC	EQ 0.1% BASE	N020688	001	Dec 18, 1996

OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE

APOTEX	20MG	A210070	001	Feb 11, 2019
+! DEXCEL PHARMA	20MG	N022032	001	Dec 04, 2007
DR REDDYS	20MG	A207740	001	Nov 05, 2018
SUN PHARM	20MG	A207891	001	Oct 12, 2018

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

OMEPRAZOLE

+! DEXCEL 20MG N209400 001 Jul 05, 2017

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

! DR REDDYS LABS LTD	EQ 20MG BASE	A078878	001	Jun 05, 2009
L PERRIGO CO	EQ 20MG BASE	A216096	001	May 24, 2022
SPIL	EQ 20MG BASE	A210593	001	Jul 20, 2018

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA	EQ 20MG BASE	A206877	001	Jun 06, 2018
HETERO LABS LTD III	EQ 20MG BASE	A211732	001	Mar 25, 2020
P AND L	EQ 20MG BASE	A206582	001	Jun 01, 2020
PRILOSEC OTC				
+! ASTRAZENECA	EQ 20MG BASE	N021229	001	Jun 20, 2003

OTC DRUG PRODUCT LIST

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE; ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

AUROBINDO PHARMA 20MG; 1.1GM

A204923 001 Nov 07, 2016

PERRIGO R AND D 20MG; 1.1GM

A201361 001 Jul 15, 2016

ZYDUS 20MG; 1.1GM

A203345 001 Mar 16, 2018

ZEGERID OTC

+! RILEY CONSUMER 20MG; 1.1GM

N022281 001 Dec 01, 2009

FOR SUSPENSION; ORAL

ZEGERID OTC

+! RILEY CONSUMER 20MG/PACKET; 1.68GM/PACKET

N022283 001 Jun 17, 2013

ORLISTAT

CAPSULE; ORAL

ALLI

+! GLAXOSMITHKLINE 60MG

N021887 001 Feb 07, 2007

CONS

OXYBUTYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYTROL FOR WOMEN

+! ABBVIE 3.9MG/24HR

N202211 001 Jan 25, 2013

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

VISINE L.R.

+! JOHNSON AND JOHNSON 0.025%

N019407 001 Mar 31, 1989

PERMETHRIN

LOTION; TOPICAL

NIX

+! MEDTECH PRODUCTS 1%

N019918 001 May 02, 1990

PERMETHRIN

ACTAVIS MID 1%

A075014 001 Mar 28, 2000

ATLANTIC

PERRIGO NEW YORK 1%

A076090 001 Dec 20, 2001

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

MIRALAX

+! BAYER HEALTHCARE 17GM/SCOOPFUL

N022015 001 Oct 06, 2006

POLYETHYLENE GLYCOL 3350

ANI PHARMS 17GM/SCOOPFUL

A202850 001 Dec 15, 2015

ANNORA PHARMA 17GM/SCOOPFUL

A214990 001 Apr 14, 2021

AUROBINDO PHARMA 17GM/SCOOPFUL

A209017 001 Apr 09, 2018

LTD

ELYSIUM 17GM/SCOOPFUL

A202071 001 Dec 28, 2012

LGM PHARMA 17GM/SCOOPFUL

A090812 001 Oct 07, 2009

MYLAN 17GM/PACKET

A078915 001 Oct 06, 2009

17GM/SCOOPFUL

A078915 002 Oct 06, 2009

NOVEL LABS INC 17GM/SCOOPFUL

A091077 001 Oct 06, 2009

NUVO PHARMS INC 17GM/SCOOPFUL

A206105 001 Oct 28, 2016

PERRIGO R AND D 17GM/PACKET

A090685 001 Oct 06, 2009

17GM/SCOOPFUL

A090685 002 Oct 06, 2009

STRIDES PHARMA 17GM/SCOOPFUL

A079214 001 Jan 31, 2013

17GM/SCOOPFUL

A203928 001 Aug 24, 2016

17GM/PACKET

A203928 002 Aug 24, 2016

POTASSIUM IODIDE

SOLUTION; ORAL

POTASSIUM IODIDE

! MISSION PHARMACAL 65MG/ML

A206211 001 Mar 24, 2016

TABLET; ORAL

IOSAT

+ ANBEX 65MG

N018664 002 May 12, 2011

+! 130MG

N018664 001 Oct 14, 1982

THYROSAFE

! BTG INTL 65MG

A076350 001 Sep 10, 2002

POVIDONE-IODINE

SOLUTION; TOPICAL

POVIDONE IODINE

+! ALLEGIANCE HLTHCARE 1%

N019522 001 Mar 31, 1989

OTC DRUG PRODUCT LIST

POVIDONE-IODINE

SPONGE; TOPICAL

E-Z SCRUB 201

+! BECTON DICKINSON 20%

N019240 001 Nov 29, 1985

E-Z SCRUB 241

+! BECTON DICKINSON 10%

N019476 001 Jan 07, 1987

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA 120MG

A209008 001 Jun 09, 2017

! L PERRIGO CO 120MG

A075153 001 Feb 26, 1999

SUN PHARM INDS LTD 120MG

A077442 001 Sep 28, 2005

SUDAFED 24 HOUR

+! J AND J CONSUMER 240MG
INC

N020021 002 Dec 15, 1992

PURIFIED WATER

SOLUTION; OPHTHALMIC

PUR-WASH

+! NIAGARA PHARMS 98.3%

N022305 001 Sep 01, 2011

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

DR REDDYS LABS LTD EQ 75MG BASE

A075294 001 Mar 28, 2000

EQ 150MG BASE

A078192 001 Aug 31, 2007

THINQ PHARM-CRO PVT EQ 75MG BASE

A210250 001 Aug 30, 2019

TERBINAFINE

GEL; TOPICAL

LAMISIL AT

+! GLAXOSMITHKLINE 1%
CONS

N021958 001 Jul 24, 2006

TERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL

LAMISIL

+! GLAXOSMITHKLINE 1%

N020980 001 Mar 09, 1999

TERBINAFINE HYDROCHLORIDE

TARO 1%

A077511 001 Jul 02, 2007

SOLUTION; TOPICAL

LAMISIL AT

+! GLAXOSMITHKLINE 1%
CONS

N021124 001 Mar 17, 2000

SPRAY; TOPICAL

LAMISIL AT

+! GLAXOSMITHKLINE 1%
CONS

N021124 002 Mar 17, 2000

TIOCONAZOLE

OINTMENT; VAGINAL

TIOCONAZOLE

PERRIGO 6.5%

A075915 001 Nov 21, 2001

VAGISTAT-1

+! COMBE 6.5%

N020676 001 Feb 11, 1997

TRIAMCINOLONE ACETONIDE

SPRAY, METERED; NASAL

NASACORT ALLERGY 24 HOUR

+! CHATTEM SANOFI 0.055MG/SPRAY

N020468 002 Oct 11, 2013

TRIAMCINOLONE ACETONIDE

APOTEX 0.055MG/SPRAY

A214615 001 Jan 19, 2023

PERRIGO PHARMA INTL 0.055MG/SPRAY

A078104 002 Nov 14, 2014

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

0.9% SODIUM CHLORIDE INJECTION USP SOLUTION

INJECTABLE; INJECTION

NONE

FRESENIUS KABI AG

N125695

Sep 05, 2019

ANTICOAGULANT 4% SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

HAEMONETICS
MANUFACTURING INC

N760305

Jun 30, 1978

NONE

CSL PLASMA INC

A125750

Apr 25, 2022

ANTICOAGULANT CITRATE DEXTROSE SOLUTION (ACD)

INJECTABLE; INJECTION

NONE

CITRA LABS LLC

N020037

Aug 26, 2003

ACD-A SOLUTION

TERUMO BCT INC

A010228

Feb 25, 2002

ADSOL WITH ACD-A

FENWAL INC

N000922

Aug 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION FORMULA A

HAEMONETICS CORP

A980728

Feb 06, 2002

AS3 SOLUTION/ACD-A

TERUMO BCT INC

N001214

May 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION A

TERUMO BCT INC

A110057

May 11, 2012

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N160918

Mar 17, 1978

ANTICOAGULANT CITRATE PHOSPHATE 2X DEXTROSE SOLUTION (CP2D)

INJECTABLE; INJECTION

CITRATE PHOSPHATE DOUBLE DEXTROSE/ADDITIVE SOLUTION 3

HAEMONETICS CORP

N000127

Jan 18, 2002

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION

INJECTABLE; INJECTION

NONE

TERUMO MEDICAL CORP

N820528

Nov 03, 1982

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION (CPDA)

INJECTABLE; INJECTION

CPDA-1 BLOOD-PACK UNIT (PL 146 PLASTIC) 250, 450, 500 ML BLOOD PACK UNITS

FENWAL INC

N770420

May 12, 1978

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION USP

INJECTABLE; INJECTION

BLOOD PACK UNIT CPDA-1 IN PLASTIC CONTAINER

FENWAL INC

N940404

Jul 28, 1994

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE-1 SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS

MANUFACTURING INC

N800077

Nov 06, 1980

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION

INJECTABLE; INJECTION

ADSOL IN PLASTIC CONTAINER

FENWAL INC

N900223

Dec 27, 1991

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION (CPD)

INJECTABLE; INJECTION

CPD ANTICOAGULANT IN PL 2209 PLASTIC CONTAINER

FENWAL INC

N900224

Dec 27, 1991

MACOPRODUCTIONS SAS CPD/AS-1: MACOPHARMA LEUCOFLEX MTL1 LEUKOREDUCTION SYSTEM FOR BLOOD COMPONENTS KNOWN AS MTL1-WB

MACOPHARMA

N040083

Nov 21, 2005

NONE

TERUMO BCT INC

A070025

Jan 06, 2009

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N170401

Dec 06, 1977

N811012

Jun 28, 1983

GLOBAL LIFE SCIENCE

SOLUTIONS USA, LLC

N800222

Aug 23, 1982

TERUMO MEDICAL CORP

N781211

Jun 10, 1981

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-1:
DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

ADSOL RED BLOOD CELL PRESERVATIVE SOLUTION

FENWAL INC

N811104

May 16, 1983

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-5:
DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

OPTISOL RED BLOOD CELL PRESERVATIVE SOLUTION

TERUMO MEDICAL CORP

N880217

Oct 07, 1988

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:
AS-3: CITRIC ACID USP; MONOBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-3 NUTRICEL ADDITIVE SYSTEM

HAEMONETICS

0.042GM/100ML;0.276GM/100ML;

N820915

Oct 19, 1984

MANUFACTURING INC

0.410GM/100ML;0.30GM/100ML;

1.10GM/100ML;0.588GM/100ML

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
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ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:

AS-2: CITRIC ACID USP; DIBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-2 NUTRICEL ADDITIVE SYSTEM

MEDSEP CORP	0.042GM/100ML;0.285GM/100ML; 0.718GM/100ML;0.017GM/100ML; 0.396GM/100ML;0.588GM/100ML	N820915	Sep 22, 1983
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ANTICOAGULANT SODIUM CITRATE 4% SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS CORPORATION		N980123	Mar 03, 2000
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LABORATORIES
GRIFOIS, S.A.

A125697	Oct 25, 2019
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TERUMO BCT

A125608	Jun 26, 2018
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ANTICOAGULANT SODIUM CITRATE SOLUTION

INJECTABLE; INJECTION

TRICITRASOL

CITRA LABS, LLC

N010409	Jul 10, 2003
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ANTICOAGULANT SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N770923	Jan 20, 1978
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TERUMO MEDICAL CORP

N781214	Feb 08, 1980
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CORD BLOOD STERILE COLLECTION BAG, ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION
(CPD)

STERILE CORD BLOOD COLLECTION UNIT

NONE

MACOPHARMA

N125552	Dec 21, 2016
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DEXTRAN 1 IN SODIUM CHLORIDE 0.6%

INJECTABLE; INJECTION

PROMIT

MEDA AB

N830715	Oct 30, 1984
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DEXTRAN 40, 10% IN DEXTROSE 5%

INJECTABLE; INJECTION

LMD IN GLASS BOTTLE

HOSPIRA INC

10GM/100ML;5GM/100ML

A720563	Oct 30, 1992
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DEXTRAN 40, 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

LMD IN PLASTIC CONTAINER

HOSPIRA INC

10GM/100ML;0.9GM/100ML

A720562	Oct 30, 1992
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HETASTARCH 6% IN LACTATED ELECTROLYTE INJECTION

INJECTABLE; INJECTION

HEXTEND

BIOTIME INC

6GM/100ML

N200952	Mar 31, 1999
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**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

HETASTARCH 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

6% HETASTARCH IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
HOSPIRA INC	6GM/100ML;0.9GM/100ML	A740193 Jan 30, 1995
HESPAN IN PLASTIC CONTAINER		
B BRAUN MEDICAL INC	6GM/100ML;0.9GM/100ML	N890105 Apr 04, 1991
NONE		
TEVA PHARMACEUTICALS USA INC	6GM/100ML;0.9GM/100ML	A740592 Nov 12, 1998

ISOPLATE SOLUTION IN THE 500 ML EXCEL CONTAINERSTORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

ISOPLATE SOLUTION		
TERUMO BCT		N90067 Mar 05, 2013

LEUKOCYTE REDUCTION FILTRATION SYSTEM FOR WHOLE BLOOD WITH CPD ANTICOAGULANT AND SOLX ADDITIVE

INJECTABLE; INJECTION

LEUKOSEP HWB-600-XL		
HEMERUS MEDICAL, LLC		N110059 Apr 25, 2013

RED BLOOD CELL PROCESSING SOLUTIONSTORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

REJUVESOL		
CITRA LABS LLC		N950522 Feb 26, 1997

SODIUM CHLORIDE; SODIUM ACETATE; SODIUM CITRATE DIHYDRATE; SODIUM PHOSPHATE, DIABASIC ANHYDROUS; SODIUM PHOSPHATE MONOBASIC, MONOHYDRATESTORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

INTERSOL SOLUTION		
FRESENIUS KABI USA, LLC	2.26GM/500ML;2.21GM/500ML;1.59GM/500ML;1.53GM/500ML;0.465GM/500ML	N080041 Dec 09, 2009

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ABACAVIR SULFATE

TABLET; ORAL

ABACAVIR SULFATE

APOTEX INC

EQ 300MG BASE

A201570 001 Dec 17, 2012

ZIAGEN

+ VIIV HLTHCARE

EQ 300MG BASE

N020977 001 Dec 17, 1998

ABACAVIR SULFATE; LAMIVUDINE

TABLET; ORAL

ABACAVIR SULFATE AND LAMIVUDINE

AUROBINDO PHARMA

EQ 600MG BASE; 300MG

A206151 001 Mar 28, 2017

TEVA PHARMS USA

EQ 600MG BASE; 300MG

A079246 001 Sep 29, 2016

ZYDUS PHARMS

EQ 600MG BASE; 300MG

A208990 001 Nov 15, 2018

ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

TRIZIVIR

+ VIIV HLTHCARE

EQ 300MG BASE; 150MG; 300MG

N021205 001 Nov 14, 2000

ABAMETAPIR

LOTION; TOPICAL

XEGLYZE

+ HATCHTECH

0.74%

N206966 001 Jul 24, 2020

ABARELIX

INJECTABLE; INTRAMUSCULAR

PLENAXIS

SPECIALITY EUROPEAN

100MG/VIAL

N021320 001 Nov 25, 2003

ABIRATERONE ACETATE

TABLET; ORAL

ABIRATERONE ACETATE

TEVA PHARMS USA

125MG

A212206 001 Jun 24, 2022

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL

ACAMPROSATE CALCIUM

BARR LABS DIV TEVA

333MG

A200143 001 Nov 18, 2013

CAMPRAL

+ FOREST LABS

333MG **

N021431 001 Jul 29, 2004

ACARBOSE

TABLET; ORAL

ACARBOSE

DASH PHARMS

25MG

A091053 001 Jan 06, 2011

50MG

A091053 002 Jan 06, 2011

100MG

A091053 003 Jan 06, 2011

HANGZHOU ZHONGMEI

25MG

A213821 001 Aug 18, 2020

50MG

A213821 002 Aug 18, 2020

100MG

A213821 003 Aug 18, 2020

PRECOSE

+ BAYER HLTHCARE

25MG **

N020482 004 May 29, 1997

+

50MG **

N020482 001 Sep 06, 1995

+

100MG **

N020482 002 Sep 06, 1995

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

MYLAN

EQ 200MG BASE

A074288 001 Apr 24, 1995

EQ 400MG BASE

A074288 002 Apr 24, 1995

SECTRAL

+ PROMIUS PHARMA

EQ 200MG BASE **

N018917 001 Dec 28, 1984

+

EQ 400MG BASE **

N018917 003 Dec 28, 1984

ACETAMINOPHEN

INJECTABLE; INJECTION

INJECTAPAP

ORTHO MCNEIL PHARM

100MG/ML

N017785 001 Mar 07, 1986

POWDER; INTRAVENOUS

ACETAMINOPHEN

+ RISING

1GM/VIAL

N206610 001 Jan 15, 2021

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN

SOLUTION; INTRAVENOUS

ACETAMINOPHEN

ZYDUS PHARMS 1GM/100ML (10MG/ML) A216467 001 Oct 25, 2022

OFIRMEV

+ MALLINCKRODT HOSP 1GM/100ML (10MG/ML) N022450 001 Nov 02, 2010

SUPPOSITORY; RECTAL

ACEPHEN

COSETTE 120MG A072218 001 Mar 27, 1992

325MG N018060 003 Dec 18, 1986

650MG N018060 002

ACETAMINOPHEN

ABLE 120MG A073106 001 Feb 27, 1995

325MG A073107 001 Feb 27, 1995

650MG A073108 001 Feb 27, 1995

ACINO PRODS 120MG A071010 001 May 12, 1987

650MG A071011 001 May 12, 1987

NEOPAP

POLYMEDICA 120MG N016401 001

TYLENOL

J AND J CONSUMER INC 120MG N017756 002

650MG N017756 001

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

SUN PHARM INDS LTD 650MG A090205 001 Nov 18, 2009

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE

MIKART 150MG;180MG;15MG A081095 001 Oct 26, 1990

150MG;180MG;30MG A081096 001 Oct 26, 1990

150MG;180MG;60MG A081097 001 Oct 26, 1990

CODEINE, ASPIRIN, APAP FORMULA NO. 2

+ SCHERER LABS 150MG;180MG;15MG A085640 001

CODEINE, ASPIRIN, APAP FORMULA NO. 3

+ SCHERER LABS 150MG;180MG;30MG A085639 001

CODEINE, ASPIRIN, APAP FORMULA NO. 4

SCHERER LABS 150MG;180MG;60MG A085638 001

ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE

TABLET; ORAL

APADAZ

+ ZEVRA THERAP 325MG;EQ 4.08MG BASE N208653 002 Jan 04, 2019

+ 325MG;EQ 6.12MG BASE N208653 001 Feb 23, 2018

+ 325MG;EQ 8.16MG BASE N208653 003 Jan 04, 2019

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BANCAP

FOREST PHARMS 325MG;50MG A088889 001 Jan 16, 1986

BUCET

MALLINCKRODT 650MG;50MG A088991 001 Jun 28, 1985

PHRENILIN FORTE

VALEANT 650MG;50MG A088831 001 Jun 19, 1985

TENCON

MALLINCKRODT 650MG;50MG A089405 001 May 15, 1990

TRIAPRIN

DUNHALL 325MG;50MG A089268 001 Jul 02, 1987

TABLET; ORAL

BUTALBITAL AND ACETAMINOPHEN

ALVOGEN 300MG;50MG A207635 001 Jun 05, 2017

325MG;50MG A205120 001 Oct 30, 2015

HALSEY 325MG;50MG A089568 001 Oct 05, 1988

+ WATSON LABS 325MG;50MG A087550 001 Oct 19, 1984

BUTAPAP

MIKART 650MG;50MG A089988 001 Oct 26, 1992

PHRENILIN

+ VALEANT 325MG;50MG ** A087811 001 Jun 19, 1985

SEDAPAP

MAYRAND 650MG;50MG A088944 001 Oct 17, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

ANOQUAN

SHIRE 325MG; 50MG; 40MG A087628 001 Oct 01, 1986

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

+ GILBERT LABS 325MG; 50MG; 40MG ** A088825 001 Dec 05, 1984

GRAHAM DM 325MG; 50MG; 40MG A088743 001 Jul 18, 1985

325MG; 50MG; 40MG A088765 001 Mar 27, 1985

325MG; 50MG; 40MG A089067 001 Apr 19, 1985

HIKMA 300MG; 50MG; 40MG A215135 001 Mar 25, 2022

325MG; 50MG; 40MG A215135 002 Mar 25, 2022

500MG; 50MG; 40MG A040261 001 Oct 28, 1998

MALLINCKRODT 325MG; 50MG; 40MG A088758 001 Mar 27, 1985

ESGIC-PLUS

MIKART 500MG; 50MG; 40MG A040085 001 Mar 28, 1996

FEMCET

MALLINCKRODT 325MG; 50MG; 40MG A089102 001 Jun 19, 1985

MEDIGESIC PLUS

US CHEM 325MG; 50MG; 40MG A089115 001 Jan 14, 1986

TRIAD

MALLINCKRODT 325MG; 50MG; 40MG A089023 001 Jun 19, 1985

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

ABLE 325MG; 50MG; 40MG A040390 001 Jul 23, 2001

500MG; 50MG; 40MG A040394 001 Jul 23, 2001

ALVOGEN 325MG; 50MG; 40MG A204984 001 Jan 10, 2017

GILBERT LABS 325MG; 50MG; 40MG A087629 001 Nov 13, 1984

HIKMA 500MG; 50MG; 40MG A040336 001 Aug 18, 1999

HIKMA PHARMS 325MG; 50MG; 40MG A089718 001 Jun 12, 1995

MIKART 750MG; 50MG; 40MG A040496 001 Dec 23, 2003

MIRROR PHARMS LLC 500MG; 50MG; 40MG A040883 001 Dec 23, 2008

NESHER PHARMS 325MG; 50MG; 40MG A211543 001 Jul 17, 2020

SPECGX LLC 325MG; 50MG; 40MG A087804 001 Jan 24, 1985

SUN PHARM INDUSTRIES 325MG; 50MG; 40MG A040601 001 Jul 29, 2005

VINTAGE PHARMS 500MG; 50MG; 40MG A040513 001 Aug 25, 2003

WATSON LABS 325MG; 50MG; 40MG A089536 001 Feb 16, 1988

500MG; 50MG; 40MG A040267 001 Jul 30, 1998

ESGIC

FOREST PHARMS 325MG; 50MG; 40MG A089660 001 Dec 23, 1988

ESGIC-PLUS

MIKART 500MG; 50MG; 40MG A089451 001 May 23, 1988

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

ABLE 325MG; 50MG; 40MG; 30MG A076528 001 Aug 21, 2003

HIKMA INTL PHARMS 325MG; 50MG; 40MG; 30MG A075618 001 Mar 23, 2001

PHRENILIN WITH CAFFEINE AND CODEINE

VALEANT 325MG; 50MG; 40MG; 30MG A074911 001 Aug 22, 2001

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

MIKART 356.4MG; 30MG; 16MG A040109 001 Aug 26, 1997

WRASER PHARMS LLC 356.4MG; 30MG; 16MG A040688 001 Apr 03, 2007

DHC PLUS

PHARM RES ASSOC 356.4MG; 30MG; 16MG A088584 001 Mar 04, 1986

SYNALGOS-DC-A

LEITNER PHARMS 356.4MG; 30MG; 16MG A089166 001 May 14, 1986

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE

LARKEN LABS INC 325MG; 30MG; 16MG A204209 001 Sep 30, 2016

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

BOCA PHARMA LLC 712.8MG; 60MG; 32MG A040701 001 Apr 03, 2007

MIKART 712.8MG; 60MG; 32MG A040316 001 Apr 28, 1999

WEST-WARD PHARM CORP 712.8MG; 60MG; 32MG A040637 001 Sep 22, 2006

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET;ORAL

TAVIST ALLERGY/SINUS/HEADACHE

NOVARTIS

500MG;EQ 0.25MG BASE;30MG

N021082 001 Mar 01, 2001

ACETAMINOPHEN; CODEINE PHOSPHATE

CAPSULE;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

TEVA

300MG;15MG

A088537 001 Jun 04, 1984

300MG;30MG

A088324 001 Dec 29, 1983

300MG;60MG

A088599 001 Jun 01, 1984

PHENAPHEN W/ CODEINE NO. 2

ROBINS AH

325MG;15MG

A084444 001

PHENAPHEN W/ CODEINE NO. 3

+ ROBINS AH

325MG;30MG

A084445 001

PHENAPHEN W/ CODEINE NO. 4

+ ROBINS AH

325MG;60MG

A084446 001

PROVAL #3

SOLVAY

325MG;30MG

A085685 001

TYLENOL W/ CODEINE NO. 3

ORTHO MCNEIL PHARM

300MG;30MG

A087422 001

TYLENOL W/ CODEINE NO. 4

ORTHO MCNEIL PHARM

300MG;60MG

A087421 001

SOLUTION;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

ACI

120MG/5ML;12MG/5ML

A086366 001

+ ACTAVIS MID ATLANTIC

120MG/5ML;12MG/5ML

A085861 001

CHARTWELL MOLECULAR

120MG/5ML;12MG/5ML

A091238 001 Nov 10, 2011

DAVA PHARMS INC

120MG/5ML;12MG/5ML

A040098 001 Sep 20, 1996

WOCKHARDT BIO AG

120MG/5ML;12MG/5ML

A087006 001

TYLENOL W/ CODEINE

+ ORTHO MCNEIL PHARM

120MG/5ML;12MG/5ML

A085057 001

SUSPENSION;ORAL

CAPITAL AND CODEINE

ACTAVIS MID ATLANTIC

120MG/5ML;12MG/5ML

A085883 001

VALEANT PHARMS LLC

120MG/5ML;12MG/5ML

A086024 001

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

ABLE

300MG;30MG

A040452 001 Aug 01, 2002

300MG;60MG

A040459 001 Aug 01, 2002

AM THERAP

300MG;15MG

A089478 001 Mar 03, 1987

300MG;15MG

A089481 001 Mar 03, 1987

300MG;30MG

A089479 001 Mar 03, 1987

300MG;30MG

A089482 001 Mar 03, 1987

300MG;60MG

A089480 001 Mar 03, 1987

300MG;60MG

A089483 001 Mar 03, 1987

CHARTWELL

300MG;15MG

A089997 001 Dec 28, 1994

300MG;30MG

A089998 001 Dec 28, 1994

300MG;60MG

A089999 001 Dec 28, 1994

DURAMED PHARMS BARR

300MG;15MG

A040223 001 Nov 18, 1997

300MG;15MG

A088353 001 Feb 06, 1984

300MG;30MG

A040223 002 Nov 18, 1997

300MG;30MG

A088354 001 Feb 06, 1984

300MG;60MG

A040223 003 Nov 18, 1997

300MG;60MG

A088355 001 Feb 06, 1984

EVERYLIFE

325MG;30MG

A085217 001

FOSUN PHARMA

300MG;30MG

A081250 001 Jul 16, 1992

300MG;60MG

A081249 001 Jul 16, 1992

HALSEY

300MG;15MG

A083871 001

300MG;30MG

A083872 001

300MG;60MG

A086549 001

KV PHARM

300MG;30MG

A085288 001

300MG;60MG

A085365 001

325MG;15MG

A085364 001

325MG;45MG **

A085363 001

LEDERLE

300MG;30MG

A087141 001

MIKART

300MG;30MG

A089238 001 Feb 25, 1986

300MG;60MG

A089244 001 Feb 25, 1986

650MG;30MG

A089231 001 Mar 03, 1986

650MG;60MG

A089363 001 Sep 09, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

MUTUAL PHARM	300MG;15MG	A085795	001	
	300MG;30MG	A085794	001	
	300MG;60MG	A087653	001	Apr 13, 1982
NOSTRUM LABS INC	300MG;15MG	A088629	002	Mar 06, 1985
	300MG;30MG	A088629	003	Mar 06, 1985
+	300MG;60MG	A088629	001	Mar 06, 1985
PURACAP PHARM	300MG;30MG	A087762	001	Dec 10, 1982
+	300MG;30MG	A086681	001	
	300MG;30MG	A089080	001	Jul 17, 1986
	300MG;60MG	A086683	001	
RHODES PHARMS	300MG;15MG	A089673	002	Feb 10, 1988
	300MG;30MG	A089673	003	Feb 10, 1988
	300MG;60MG	A089673	001	Feb 10, 1988
ROXANE	300MG;15MG	A084659	001	
	300MG;30MG	A084656	001	
	300MG;60MG	A084667	001	
	500MG;15MG	A089511	001	Apr 25, 1989
	500MG;30MG	A089512	001	Apr 25, 1989
	500MG;60MG	A089513	001	Apr 25, 1989
+	300MG;15MG	A087433	001	
	300MG;30MG	A085291	002	
+	300MG;30MG	A085917	001	
	300MG;60MG	A085964	001	
	300MG;60MG	A087423	001	
STRIDES PHARMA	300MG;15MG	A089990	001	Sep 30, 1988
	300MG;30MG	A089805	001	Sep 30, 1988
	300MG;60MG	A089828	001	Sep 30, 1988
SUPERPHARM	300MG;15MG	A089183	001	Oct 18, 1985
	300MG;30MG	A089184	001	Oct 18, 1985
	300MG;30MG	A089253	001	May 19, 1986
	300MG;60MG	A089185	001	Oct 18, 1985
	300MG;60MG	A089254	001	May 19, 1986
USL PHARMA	300MG;30MG	A087919	001	Jun 22, 1982
	300MG;60MG	A087920	001	Jun 22, 1982
VALEANT PHARM INTL	300MG;30MG	A085896	001	
VITARINE	300MG;30MG	A085676	001	
WARNER CHILCOTT	300MG;15MG	A085992	001	
	300MG;30MG	A085218	002	
	300MG;60MG	A087306	001	
WATSON LABS	300MG;15MG	A087277	001	May 26, 1982
	300MG;30MG	A087276	001	May 26, 1982
	300MG;60MG	A087275	001	May 26, 1982
WATSON LABS FLORIDA	300MG;15MG	A040443	001	Jan 22, 2003
	300MG;30MG	A040443	002	Jan 22, 2003
	300MG;60MG	A040443	003	Jan 22, 2003
WHITEWORTH TOWN PLSN	300MG;30MG	A084360	001	
	300MG;60MG	A085607	001	
CAPITAL AND CODEINE				
CARNRICK	325MG;30MG	A083643	001	
CODRIX				
WATSON LABS FLORIDA	500MG;15MG	A040447	001	Feb 26, 2003
	500MG;30MG	A040441	001	Mar 27, 2003
	500MG;60MG	A040488	001	Mar 28, 2003
EMPRACET W/ CODEINE PHOSPHATE #3				
GLAXOSMITHKLINE	300MG;30MG	A083951	001	
EMPRACET W/ CODEINE PHOSPHATE #4				
GLAXOSMITHKLINE	300MG;60MG	A083951	002	
PAPA-DEINE #3				
VANGARD	300MG;30MG	A088037	001	Mar 20, 1984
PAPA-DEINE #4				
VANGARD	300MG;60MG	A088715	001	Mar 20, 1984
PHENAPHEN-650 W/ CODEINE				
ROBINS AH	650MG;30MG	A085856	001	
TYLENOL W/ CODEINE				
ORTHO MCNEIL PHARM	325MG;7.5MG **	A085056	001	
	325MG;15MG **	A085056	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

TYLENOL W/ CODEINE

325MG; 30MG **

A085056 003

325MG; 60MG **

A085056 004

TYLENOL W/ CODEINE NO. 1

JANSSEN PHARMS

300MG; 7.5MG

A085055 001

TYLENOL W/ CODEINE NO. 2

+ JANSSEN PHARMS

300MG; 15MG

A085055 002

TYLENOL W/ CODEINE NO. 3

+ JANSSEN PHARMS

300MG; 30MG

A085055 003

TYLENOL W/ CODEINE NO. 4

+ JANSSEN PHARMS

300MG; 60MG

A085055 004

ACETAMINOPHEN; DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

DRIXORAL PLUS

SCHERING PLOUGH

500MG; 3MG; 60MG

N019453 001 May 22, 1987

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN AND HYDROCODONE BITARTRATE

CENT PHARMS

500MG; 5MG

A088898 001 Mar 27, 1985

ALLAY

IVAX PHARMS

500MG; 5MG

A089907 001 Jan 13, 1989

BANCAP HC

FOREST PHARMS

500MG; 5MG

A087961 001 Mar 17, 1983

CO-GESIC

CENT PHARMS

500MG; 5MG

A089360 001 Mar 02, 1988

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

MALLINCKRODT

500MG; 5MG

A088956 001 Jul 19, 1985

500MG; 5MG

A089006 001 Aug 09, 1985

MIKART

500MG; 5MG

A081067 001 Nov 30, 1989

500MG; 5MG

A081068 001 Nov 30, 1989

500MG; 5MG

A081069 001 Nov 30, 1989

500MG; 5MG

A081070 001 Nov 30, 1989

500MG; 5MG

A089008 001 Feb 21, 1986

LORCET-HD

MALLINCKRODT

500MG; 5MG

A087336 001 Jul 08, 1982

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

GENUS

325MG/15ML; 7.5MG/15ML

A200343 001 Jan 25, 2012

MALLINCKRODT

500MG/15ML; 7.5MG/15ML

A040418 001 Jun 27, 2001

MALLINCKRODT INC

500MG/15ML; 10MG/15ML

A040508 001 Aug 29, 2003

MIKART

500MG/15ML; 5MG/15ML

A081226 001 Oct 27, 1992

500MG/15ML; 5MG/15ML

A089557 001 Apr 29, 1992

500MG/15ML; 7.5MG/15ML

A081051 001 Aug 28, 1992

NESHER PHARMS

500MG/15ML; 7.5MG/15ML

A040366 001 Jan 23, 2002

PHARM ASSOC

500MG/15ML; 7.5MG/15ML

A040182 001 Mar 13, 1998

TRIS PHARMA INC

300MG/15ML; 10MG/15ML

A201295 002 Dec 30, 2021

325MG/15ML; 7.5MG/15ML

A201295 001 Dec 30, 2021

VINTAGE PHARMS

500MG/15ML; 7.5MG/15ML

A040520 001 Oct 30, 2003

ZYFREL

CYPRESS PHARM INC

325MG/15ML; 7.5MG/15ML

A090468 001 Apr 14, 2016

TABLET; ORAL

ANEXSIA

MALLINCKRODT

500MG; 5MG

A089160 001 Apr 23, 1987

750MG; 10MG

A040468 001 Oct 31, 2002

ANEXSIA 7.5/650

MALLINCKRODT

650MG; 7.5MG

A089725 001 Sep 30, 1987

CO-GESIC

UCB INC

500MG; 5MG

A087757 001 May 03, 1982

DURADYNE DHC

FOREST PHARMS

500MG; 5MG

A087809 001 Mar 17, 1983

HY-PHEN

ASCHER

500MG; 5MG

A087677 001 May 03, 1982

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

ABLE

325MG; 5MG

A040478 001 Nov 08, 2002

325MG; 7.5MG

A040464 001 Oct 23, 2002

325MG; 10MG

A040464 002 Oct 23, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	500MG; 5MG	A040477	001	Nov 06, 2002
	500MG; 7.5MG	A040490	001	May 21, 2003
	500MG; 10MG	A040473	001	Nov 06, 2002
	650MG; 7.5MG	A040474	001	Jan 02, 2003
	650MG; 10MG	A040476	001	Oct 23, 2002
	750MG; 7.5MG	A040469	001	Oct 25, 2002
ACTAVIS LABS FL INC	300MG; 5MG	A206470	001	Jun 02, 2016
	300MG; 7.5MG	A206470	002	Jun 02, 2016
	300MG; 10MG	A206470	003	Jun 02, 2016
ALVOGEN	300MG; 5MG	A208540	001	Nov 08, 2018
	300MG; 7.5MG	A208540	002	Nov 08, 2018
	300MG; 10MG	A208540	003	Nov 08, 2018
	325MG; 2.5MG	A209958	001	Oct 24, 2018
	325MG; 5MG	A209958	002	Oct 24, 2018
	325MG; 7.5MG	A209958	003	Oct 24, 2018
	325MG; 10MG	A209958	004	Oct 24, 2018
AMNEAL PHARMS NY	500MG; 5MG	A040729	001	Aug 25, 2006
	500MG; 7.5MG	A040748	001	Aug 25, 2006
	500MG; 10MG	A040813	001	Feb 23, 2007
	650MG; 7.5MG	A040754	001	Aug 25, 2006
	650MG; 10MG	A040757	001	Aug 25, 2006
	750MG; 7.5MG	A040769	001	Aug 28, 2006
APIL	500MG; 10MG	A040148	002	Feb 14, 1997
BARR	500MG; 2.5MG	A040307	001	Jul 26, 2000
	500MG; 5MG	A040308	001	Jul 26, 2000
	500MG; 5MG	A088577	001	Dec 21, 1984
	500MG; 7.5MG	A040307	002	Jul 26, 2000
	500MG; 10MG	A040309	001	Jul 26, 2000
	650MG; 7.5MG	A040307	003	Jul 26, 2000
	650MG; 10MG	A040307	004	Jul 26, 2000
	750MG; 7.5MG	A040308	002	Jul 26, 2000
CARACO	500MG; 5MG	A090265	001	Dec 23, 2008
	500MG; 7.5MG	A090265	002	Dec 23, 2008
	500MG; 10MG	A090265	003	Dec 23, 2008
	650MG; 7.5MG	A090380	001	Dec 23, 2008
	650MG; 10MG	A090380	002	Dec 23, 2008
	660MG; 10MG	A090380	003	Dec 23, 2008
	750MG; 7.5MG	A090380	004	Dec 23, 2008
CEROVENE INC	325MG; 5MG	A211690	001	Feb 07, 2020
	325MG; 7.5MG	A211690	002	Feb 07, 2020
	325MG; 10MG	A211690	003	Feb 07, 2020
CHARTWELL	325MG; 7.5MG	A040248	001	Apr 28, 2000
	325MG; 10MG	A040248	002	Apr 28, 2000
GRANULES	325MG; 5MG	A211729	001	Jan 03, 2020
	325MG; 7.5MG	A211729	002	Jan 03, 2020
	325MG; 10MG	A211729	003	Jan 03, 2020
HALSEY	500MG; 5MG	A089554	001	Jun 12, 1987
IVAX PHARMS	500MG; 5MG	A089696	001	Apr 21, 1988
LANNETT CO INC	300MG; 5MG	A207171	001	Jun 20, 2017
	300MG; 7.5MG	A207171	002	Jun 20, 2017
	300MG; 10MG	A207171	003	Jun 20, 2017
	325MG; 5MG	A207172	001	Jun 22, 2017
	325MG; 7.5MG	A207172	002	Jun 22, 2017
	325MG; 10MG	A207172	003	Jun 22, 2017
MALLINCKRODT	500MG; 5MG	A040084	002	Jun 01, 1995
	500MG; 7.5MG	A040201	001	Feb 27, 1998
	500MG; 10MG	A040201	002	Feb 27, 1998
	650MG; 10MG	A040084	004	Oct 16, 1996
	660MG; 10MG	A040084	003	Jul 29, 1996
	750MG; 7.5MG	A040084	001	Jun 01, 1995
MIKART	325MG; 2.5MG	A040846	001	Jun 09, 2010
	500MG; 2.5MG	A089698	001	Aug 25, 1989
	500MG; 5MG	A089271	001	Jul 16, 1986
	500MG; 5MG	A089697	001	Jan 28, 1992
	500MG; 7.5MG	A089699	001	Aug 25, 1989
	650MG; 5MG	A040849	001	Jun 09, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	650MG; 7.5MG	A089689	001	Jun 29, 1988
	650MG; 10MG	A081223	001	May 29, 1992
MUTUAL PHARM	500MG; 5MG	A040236	001	Sep 25, 1997
	650MG; 7.5MG	A040240	002	Nov 26, 1997
	650MG; 10MG	A040240	001	Nov 26, 1997
	750MG; 7.5MG	A040236	002	Sep 25, 1997
NOSTRUM LABS INC	325MG; 2.5MG	A209924	001	Nov 16, 2018
	325MG; 5MG	A209924	002	Nov 16, 2018
	325MG; 7.5MG	A209924	003	Nov 16, 2018
	325MG; 10MG	A209924	004	Nov 16, 2018
NOVEL LABS INC	300MG; 5MG	A206142	001	Nov 14, 2016
	300MG; 7.5MG	A206142	002	Nov 14, 2016
	300MG; 10MG	A206142	003	Nov 14, 2016
	325MG; 5MG	A206245	001	Dec 01, 2016
	325MG; 7.5MG	A206245	002	Dec 01, 2016
	325MG; 10MG	A206245	003	Dec 01, 2016
RANBAXY	500MG; 5MG	A040825	001	Aug 16, 2007
	500MG; 10MG	A040824	001	Aug 16, 2007
RANBAXY LABS LTD	750MG; 7.5MG	A040822	001	Aug 16, 2007
SANDOZ	500MG; 5MG	A040149	001	Jan 27, 1997
	750MG; 7.5MG	A040149	002	Jan 27, 1997
STRIDES PHARMA	300MG; 5MG	A205001	001	Jul 05, 2016
	300MG; 7.5MG	A205001	002	Jul 05, 2016
	300MG; 10MG	A205001	003	Jul 05, 2016
SUN PHARM INDS INC	325MG; 5MG	A090118	001	Dec 23, 2008
	325MG; 7.5MG	A090118	002	Dec 23, 2008
	325MG; 10MG	A090118	003	Dec 23, 2008
SUN PHARM INDS LTD	325MG; 10MG	A040826	001	Aug 16, 2007
UCB INC	500MG; 10MG	A040210	001	Aug 13, 1997
	650MG; 7.5MG	A040134	001	Nov 21, 1996
UPSHER SMITH LABS	325MG; 5MG	A206484	001	Mar 24, 2017
	325MG; 7.5MG	A206484	002	Mar 24, 2017
	325MG; 10MG	A206484	003	Mar 24, 2017
USL PHARMA	500MG; 5MG	A089290	001	May 29, 1987
	500MG; 5MG	A089291	001	May 29, 1987
VINTAGE PHARMS	300MG; 5MG	A090415	001	Jan 24, 2011
	300MG; 7.5MG	A090415	002	Jan 24, 2011
	300MG; 10MG	A090415	003	Jan 24, 2011
	500MG; 2.5MG	A040144	002	Apr 25, 1997
	500MG; 5MG	A089831	001	Sep 07, 1988
	500MG; 5MG	A089971	001	Dec 02, 1988
	500MG; 7.5MG	A040144	001	Feb 22, 1996
	500MG; 10MG	A040356	001	May 31, 2000
	650MG; 7.5MG	A040155	001	Apr 14, 1997
	650MG; 10MG	A040143	001	Feb 22, 1996
	660MG; 10MG	A040358	001	May 31, 2000
	750MG; 7.5MG	A040157	001	Apr 12, 1996
VINTAGE PHARMS LLC	500MG; 5MG	A040281	001	Sep 30, 1998
	500MG; 7.5MG	A040280	001	Sep 30, 1998
	650MG; 7.5MG	A040280	002	Sep 30, 1998
	650MG; 10MG	A040280	003	Sep 30, 1998
	750MG; 7.5MG	A040281	002	Sep 30, 1998
WATSON LABS	500MG; 2.5MG	A040123	003	Mar 04, 1996
	500MG; 2.5MG	A081079	001	Aug 30, 1991
	500MG; 5MG	A040122	001	Mar 04, 1996
	500MG; 5MG	A089883	001	Dec 01, 1988
	500MG; 7.5MG	A040123	004	Mar 04, 1996
	500MG; 7.5MG	A081080	001	Aug 30, 1991
	650MG; 7.5MG	A040094	001	Sep 29, 1995
	650MG; 7.5MG	A040123	001	Mar 04, 1996
	650MG; 10MG	A040094	002	Sep 29, 1995
	650MG; 10MG	A040123	002	Mar 04, 1996
	660MG; 10MG	A040094	003	Aug 08, 2000
	750MG; 7.5MG	A040122	002	Mar 04, 1996
	750MG; 7.5MG	A081083	001	Aug 30, 1991
	750MG; 10MG	A040094	004	Mar 22, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

WATSON LABS FLORIDA	500MG;5MG	A040493	001	May 28, 2003
	660MG;10MG	A040495	001	May 28, 2003
	750MG;7.5MG	A040494	001	May 28, 2003
LORTAB				
UCB INC	500MG;5MG	A087722	001	Jul 09, 1982
	500MG;10MG	A040100	001	Jan 26, 1996
NORCET				
ABANA	500MG;5MG	A088871	001	May 15, 1986
NORCO				
APIL	325MG;2.5MG	A040148	004	Jul 07, 2014
	325MG;5MG	A040099	001	Jun 25, 1997
	325MG;5MG	A040148	005	Jul 07, 2014
	325MG;7.5MG	A040148	003	Sep 12, 2000
	325MG;10MG	A040148	001	Feb 14, 1997
TYCOLET				
ORTHO MCNEIL PHARM	500MG;5MG	A089385	001	Aug 27, 1986
VICODIN				
ABBOTT	500MG;5MG	A085667	001	
ABBVIE	500MG;5MG	A088058	001	Jan 07, 1983
VICODIN ES				
ABBVIE	750MG;7.5MG	A089736	001	Dec 09, 1988
VICODIN HP				
ABBVIE	660MG;10MG	A040117	001	Sep 23, 1996
ZYDONE				
VINTAGE PHARMS LLC	400MG;5MG	A040288	001	Nov 27, 1998
	400MG;7.5MG	A040288	002	Nov 27, 1998
	400MG;10MG	A040288	003	Nov 27, 1998

ACETAMINOPHEN; IBUPROFEN

TABLET; ORAL

COMBOGESIC

+ AFT PHARMS LTD	325MG;97.5MG	N209471	001	Mar 01, 2023
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ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

OXYCODONE AND ACETAMINOPHEN

ACTAVIS ELIZABETH	500MG;5MG	A040199	001	Dec 30, 1998
BARR	500MG;5MG	A040304	001	Oct 02, 2000
DURAMED PHARMS BARR	500MG;5MG	A040289	001	Mar 16, 1999
HALSEY	500MG;5MG	A089994	001	May 04, 1989
MALLINCKRODT	500MG;5MG	A040257	001	Aug 04, 1998
MUTUAL PHARM	500MG;5MG	A040219	001	Jan 22, 1998
VINTAGE PHARMS	500MG;5MG	A040106	001	Jul 30, 1996
VINTAGE PHARMS LLC	500MG;5MG	A040303	001	Dec 30, 1999
WATSON LABS	500MG;5MG	A040234	001	Oct 30, 1997
ROXILOX				
ROXANE	500MG;5MG	A040061	001	Jul 03, 1995
TYLOX				
JANSSEN PHARMS	500MG;5MG	A088790	001	Dec 12, 1984
TYLOX-325				
ORTHO MCNEIL PHARM	325MG;5MG	A088246	001	Nov 08, 1984
SOLUTION; ORAL				
OXYCODONE AND ACETAMINOPHEN				
SPECGX LLC	325MG/5ML; 5MG/5ML	A040680	001	Sep 29, 2006
OXYCODONE HYDROCHLORIDE AND ACETAMINOPHEN				
VINTAGE PHARMS	325MG/5ML; 5MG/5ML	A203573	001	Dec 18, 2014
ROXICET				
HIKMA	325MG/5ML; 5MG/5ML **	A089351	001	Dec 03, 1986
TABLET; ORAL				
OXYCODONE 2.5/APAP 500				
BRISTOL MYERS SQUIBB	500MG;2.5MG	A085910	001	
OXYCODONE 5/APAP 500				
BRISTOL MYERS SQUIBB	500MG;5MG	A085911	001	
OXYCODONE AND ACETAMINOPHEN				
ACTAVIS ELIZABETH	325MG;5MG	A040203	001	Mar 15, 1999
	325MG;7.5MG	A040800	001	Apr 03, 2012
	325MG;10MG	A040800	002	Apr 03, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

AMNEAL PHARMS NY	500MG;7.5MG	A040789 001	Nov 27, 2007
	650MG;10MG	A040789 002	Nov 27, 2007
BARR	325MG;5MG	A087406 001	
CEROVENE INC	325MG;5MG	A207574 001	Dec 13, 2016
DR REDDYS LABS SA	325MG;2.5MG	A090177 001	Oct 20, 2008
	325MG;5MG	A090177 002	Oct 20, 2008
	325MG;7.5MG	A090177 003	Oct 20, 2008
	325MG;10MG	A090177 004	Oct 20, 2008
	500MG;7.5MG	A090177 005	Oct 20, 2008
	650MG;10MG	A090177 006	Oct 20, 2008
DURAMED PHARMS BARR	325MG;5MG	A040272 001	Jun 30, 1998
GRANULES	325MG;2.5MG	A211708 001	Oct 31, 2019
	325MG;5MG	A211708 002	Oct 31, 2019
	325MG;7.5MG	A211708 003	Oct 31, 2019
	325MG;10MG	A211708 004	Oct 31, 2019
LANNETT CO INC	325MG;5MG	A207333 001	Sep 25, 2017
	325MG;10MG	A207333 002	Sep 25, 2017
MALLINCKRODT	500MG;7.5MG	A040550 001	Jun 30, 2004
	650MG;10MG	A040550 002	Jun 30, 2004
MIKART	400MG;2.5MG	A040679 001	May 16, 2006
	400MG;5MG	A040687 001	Apr 27, 2006
	400MG;7.5MG	A040698 001	Apr 27, 2006
	400MG;10MG	A040692 001	Apr 27, 2006
	500MG;10MG	A040676 001	Apr 19, 2006
NESHER PHARMS	325MG;2.5MG	A210079 001	Dec 28, 2017
	325MG;5MG	A210079 002	Dec 28, 2017
	325MG;7.5MG	A210079 003	Dec 28, 2017
	325MG;10MG	A210079 004	Dec 28, 2017
NOSTRUM LABS INC	325MG;5MG	A209385 001	Jul 02, 2018
	325MG;7.5MG	A209385 002	Jul 02, 2018
	325MG;10MG	A209385 003	Jul 02, 2018
SUN PHARM INDS INC	325MG;2.5MG	A090535 001	Dec 26, 2013
	325MG;5MG	A090535 002	Dec 26, 2013
	325MG;7.5MG	A090535 003	Dec 26, 2013
	325MG;10MG	A090535 004	Dec 26, 2013
VINTAGE PHARMS	325MG;2.5MG	A090733 001	Jul 11, 2013
	325MG;5MG	A040105 001	Jul 30, 1996
	325MG;7.5MG	A090734 001	Jul 11, 2013
	325MG;10MG	A090734 002	Jul 11, 2013
WATSON LABS	325MG;5MG	A040171 001	Oct 30, 1997
	325MG;7.5MG	A040535 001	Sep 05, 2003
	325MG;10MG	A040535 002	Sep 05, 2003
	500MG;7.5MG	A040371 001	Dec 29, 2000
	650MG;10MG	A040371 002	Dec 29, 2000
PERCOCET			
+ VINTAGE PHARMS LLC	325MG;5MG **	A085106 002	
	500MG;7.5MG	A040341 001	Jul 26, 1999
	650MG;10MG	A040341 002	Jul 26, 1999
ROXICET			
+ HIKMA	325MG;5MG	A087003 001	
ROXICET 5/500			
ROXANE	500MG;5MG	A089775 001	Jan 12, 1989
TABLET, EXTENDED RELEASE; ORAL			
XARTEMIS XR			
+ MALLINCKRODT INC	325MG;7.5MG **	N204031 001	Mar 11, 2014
<u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE</u>			
CAPSULE; ORAL			
TYLOX			
ORTHO MCNEIL PHARM	500MG;4.5MG;0.38MG	A085375 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE

GAVIS PHARMS	650MG;EQ 25MG BASE	A076202	001	Aug 02, 2002
WATSON LABS	650MG;EQ 25MG BASE	A074699	001	Mar 24, 2000
TALACEN				
SANOFI AVENTIS US	650MG;EQ 25MG BASE	N018458	001	Sep 23, 1982

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET; ORAL

DARVOCET

AAIPHARMA LLC	325MG;32.5MG	N016844	001	
DOLENE AP-65				
LEDERLE	650MG;65MG	A085100	001	
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN				
MYLAN	325MG;32MG	A083689	001	
	650MG;65MG	A083978	001	
SANDOZ	650MG;65MG	A089959	001	Jul 18, 1989
VINTAGE PHARMS	650MG;65MG	A040507	001	Jul 30, 2003
WATSON LABS	650MG;65MG	A040139	001	Dec 16, 1996
WYGESIC				
CARACO	650MG;65MG	A084999	001	

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

DARVOCET A500

XANODYNE PHARM	500MG;100MG	A076429	001	Sep 10, 2003
DARVOCET-N 100				
XANODYNE PHARM	650MG;100MG	N017122	002	
DARVOCET-N 50				
XANODYNE PHARM	325MG;50MG	N017122	001	
PROPACET 100				
TEVA	650MG;100MG	A070107	001	Jun 12, 1985
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN				
ABLE	650MG;100MG	A075838	001	Jul 11, 2001
ACTAVIS ELIZABETH	650MG;100MG	A070910	001	Jan 02, 1987
CORNERSTONE	325MG;100MG	A076743	001	May 07, 2004
	500MG;100MG	A076750	001	Jun 28, 2004
HALSEY	325MG;50MG	A072105	001	May 13, 1988
	650MG;100MG	A072106	001	May 13, 1988
IVAX SUB TEVA PHARMS	650MG;100MG	A070146	001	Aug 02, 1985
MALLINCKRODT	650MG;100MG	A075738	001	Feb 02, 2001
MIRROR PHARMS	650MG;100MG	A077821	001	Feb 11, 2008
MUTUAL PHARM	325MG;50MG	A070115	001	Jun 12, 1985
	650MG;100MG	A070116	001	Jun 12, 1985
	650MG;100MG	A070615	001	Mar 21, 1986
	650MG;100MG	A070771	001	Mar 21, 1986
	650MG;100MG	A070775	001	Mar 21, 1986
MYLAN	650MG;100MG	A072195	001	Feb 16, 1988
MYLAN PHARMS INC	650MG;100MG	A070145	001	Jun 12, 1985
SANDOZ	650MG;100MG	A070443	001	Jan 23, 1986
SUPERPHARM	650MG;100MG	A071319	001	Jan 06, 1987
TEVA	650MG;100MG	A070732	001	Jan 03, 1986
	650MG;100MG	A074119	001	Dec 19, 1994
VINTAGE PHARMS	325MG;50MG	A074843	002	Feb 15, 2001
	650MG;100MG	A074843	001	Feb 12, 1997
WATSON LABS	325MG;50MG	A070398	001	Dec 18, 1986
	650MG;100MG	A070399	001	Dec 18, 1986
WATSON LABS FLORIDA	500MG;100MG	A077196	001	Jun 28, 2005
	650MG;100MG	A076609	001	Nov 16, 2004
WOCKHARDT LTD	325MG;50MG	A077677	001	Mar 16, 2007
	650MG;100MG	A077677	002	Mar 16, 2007

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

CHARTWELL RX	325MG;37.5MG	A076475	001	Apr 21, 2005
GRAVITI PHARMS	325MG;37.5MG	A076914	001	Jul 26, 2006
MACLEODS PHARMS LTD	325MG;37.5MG	A206885	001	May 02, 2017
NOSTRUM LABS INC	325MG;37.5MG	A078778	001	Apr 07, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

ULTRACET

+ JANSSEN PHARMS 325MG; 37.5MG N021123 001 Aug 15, 2001

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

RISING 500MG A203917 001 Jun 18, 2019

DIAMOX

+ TEVA BRANDED PHARM 500MG ** N012945 001

TABLET; ORAL

ACETAZOLAMIDE

ALRA 250MG A083320 001

ASCOT 250MG A087686 001 Oct 20, 1982

BRECKENRIDGE 125MG A207503 001 Apr 30, 2020

250MG A207503 002 Apr 30, 2020

HERITAGE PHARMA 250MG A088882 001 Oct 22, 1985

SUN PHARM INDUSTRIES 125MG A089753 002 Jun 22, 1988

250MG A089753 001 Jun 22, 1988

VANGARD 250MG A087654 001 Feb 05, 1982

WATSON LABS 250MG A084498 002

DIAMOX

+ TEVA BRANDED PHARM 125MG ** N008943 001

+ 250MG ** N008943 002

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

HOSPIRA EQ 500MG BASE/VIAL A040108 001 Oct 30, 1995

PAR STERILE PRODUCTS EQ 500MG BASE/VIAL A205358 001 Jun 20, 2017

DIAMOX

+ TEVA WOMENS EQ 500MG BASE/VIAL ** N009388 001

ACETIC ACID, GLACIAL

SOLUTION/DROPS; OTIC

ACETASOL

ACTAVIS MID ATLANTIC 2% A087146 001

ACETIC ACID

CHARTWELL RX 2% A040166 001 Jul 26, 1996

KV PHARM 2% A085493 001

ORLEX

WARNER CHILCOTT 2% A086845 001

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS; OTIC

ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE

BAUSCH AND LOMB 2%; 0.79% A040063 001 Feb 25, 1994

BOROFAIR

PHARMAFAIR 2%; 0.79% A088606 001 Aug 21, 1985

DOMEBORO

+ BAYER PHARMS 2%; 0.79% A084476 001

ACETIC ACID, GLACIAL; DESONIDE

SOLUTION/DROPS; OTIC

TRIDESILON

BAYER PHARMS 2%; 0.05% N017914 001

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

ACETASOL HC

ACTAVIS MID ATLANTIC 2%; 1% A087143 001 Jan 13, 1982

ACETIC ACID W/ HYDROCORTISONE

KV PHARM 2%; 1% A085492 001

HYDROCORTISONE AND ACETIC ACID

BAUSCH AND LOMB 2%; 1% A040097 001 Oct 31, 1994

WOCKHARDT 2%; 1% A040168 001 Aug 30, 1996

ORLEX HC

WARNER CHILCOTT 2%; 1% A086844 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETIC ACID, GLACIAL; HYDROCORTISONE; NEOMYCIN SULFATE

SUSPENSION/DROPS;OTIC

NEO-CORT-DOME

BAYER PHARMS

2%;1%;EQ 0.35% BASE

N050238 001

ACETOHEXAMIDE

TABLET;ORAL

ACETOHEXAMIDE

ANI PHARMS

250MG

A070870 002 Feb 09, 1987

500MG

A070870 001 Feb 09, 1987

USL PHARMA

250MG

A070753 001 Nov 03, 1986

500MG

A070754 001 Nov 03, 1986

WATSON LABS TEVA

250MG

A071893 001 Nov 25, 1987

500MG

A071894 001 Nov 25, 1987

DYMELOR

LILLY

250MG

N013378 002

500MG

N013378 001

ACETOPHENAZINE MALEATE

TABLET;ORAL

TINDAL

SCHERING

20MG

N012254 002

ACETRIZOATE SODIUM

SOLUTION;INTRAUTERINE

SALPIX

ORTHO MCNEIL PHARM

53%

N009008 001

ACETYLCHOLINE CHLORIDE

FOR SOLUTION;OPHTHALMIC

MIOCHOL

+ NOVARTIS

20MG/VIAL **

N016211 001

ACETYLCYSTEINE

INJECTABLE;INTRAVENOUS

ACETYLCYSTEINE

EXELA PHARMA

6GM/30ML (200MG/ML)

A204797 001 Apr 15, 2021

SOLUTION;INHALATION, ORAL

ACETYLCYSTEINE

HOSPIRA

10%

A071364 001 May 01, 1989

20%

A071365 001 May 01, 1989

ROXANE

10%

A072323 001 Apr 30, 1992

10%

A072621 001 Sep 30, 1992

20%

A072324 001 Apr 30, 1992

20%

A072622 001 Sep 30, 1992

MUCOMYST

+ APOTHECON

10% **

N013601 002

+

20% **

N013601 001

MUCOSIL-10

DEY

10%

A070575 001 Oct 14, 1986

MUCOSIL-20

DEY

20%

A070576 001 Oct 14, 1986

TABLET, EFFERVESCENT;ORAL

CETYLEV

+ ARBOR PHARMS LLC

500MG

N207916 001 Jan 29, 2016

+

2.5GM

N207916 002 Jan 29, 2016

ACETYLCYSTEINE; ISOPROTERENOL HYDROCHLORIDE

SOLUTION;INHALATION

MUCOMYST W/ ISOPROTERENOL

MEAD JOHNSON

10%;0.05%

N017366 001

ACETYLDIGITOXIN

TABLET;ORAL

ACYLANID

NOVARTIS

0.1MG

N009436 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACITRETIN

CAPSULE; ORAL

ACITRETIN

MYLAN

17.5MG

A203707 001 Sep 10, 2015

22.5MG

A203707 002 Sep 10, 2015

SORIATANE

+ STIEFEL LABS INC

10MG **

N019821 001 Oct 28, 1996

+

17.5MG **

N019821 003 Aug 06, 2009

+

22.5MG **

N019821 004 Aug 06, 2009

+

25MG **

N019821 002 Oct 28, 1996

ACRISORCIN

CREAM; TOPICAL

AKRINOL

SCHERING

2MG/GM

N012470 001

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

SEMPREX-D

+ AUXILIUM PHARMS LLC

8MG; 60MG

N019806 001 Mar 25, 1994

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

ACTAVIS ELIZABETH

200MG

A074906 001 Aug 26, 1997

AUROBINDO PHARMA USA

200MG

A074727 001 Apr 22, 1997

CHARTWELL MOLECULES

200MG

A074872 001 Apr 22, 1997

IVAX SUB TEVA PHARMS

200MG

A074674 001 Apr 22, 1997

LEK PHARM

200MG

A074750 001 Apr 22, 1997

MYLAN

200MG

A074977 001 Apr 13, 1998

RANBAXY

200MG

A074975 001 Sep 30, 1998

ROXANE

200MG

A074570 002 Apr 22, 1997

STRIDES PHARMA

200MG

A074833 001 Apr 22, 1997

TEVA

200MG

A074828 001 Apr 22, 1997

TEVA PHARMS

200MG

A074914 001 Nov 26, 1997

WATSON LABS

200MG

A075101 001 Apr 15, 1998

ZOVIRAX

+ MYLAN

200MG **

N018828 001 Jan 25, 1985

OINTMENT; OPHTHALMIC

AVACLYR

+ FERA PHARMS LLC

3% **

N202408 001 Mar 29, 2019

OINTMENT; TOPICAL

ACYCLOVIR

ANDA REPOSITORY

5%

A206437 001 Jul 31, 2017

PADAGIS ISRAEL

5%

A205659 001 Feb 20, 2019

PRINSTON INC

5%

A212202 001 Nov 15, 2021

SUSPENSION; ORAL

ACYCLOVIR

VISTAPHARM

200MG/5ML

A213951 001 Jan 11, 2021

ZOVIRAX

+ MYLAN

200MG/5ML

N019909 001 Dec 22, 1989

TABLET; ORAL

ACYCLOVIR

ACTAVIS ELIZABETH

400MG

A074870 001 Jun 05, 1997

800MG

A074870 002 Jun 05, 1997

AUROBINDO PHARMA USA

400MG

A075211 001 Sep 28, 1998

800MG

A075211 002 Sep 28, 1998

CHARTWELL MOLECULES

400MG

A074834 001 Apr 24, 1997

800MG

A074834 002 Apr 24, 1997

IVAX SUB TEVA PHARMS

400MG

A074836 001 Apr 22, 1997

800MG

A074836 002 Apr 22, 1997

LEK PHARM

400MG

A074658 001 Apr 22, 1997

800MG

A074658 002 Apr 22, 1997

MYLAN

400MG

A074976 001 Apr 13, 1998

800MG

A074976 002 Apr 13, 1998

SUN PHARM INDS LTD

400MG

A074980 001 Sep 30, 1998

800MG

A074980 002 Sep 30, 1998

TEVA

200MG **

A074556 001 Apr 22, 1997

TEVA PHARMS

400MG

A075021 001 Mar 18, 1998

800MG

A075021 002 Mar 18, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACYCLOVIR

TABLET; ORAL

ZOVIRAX

+	MYLAN	400MG **	N020089 001	Apr 30, 1991
+		800MG **	N020089 002	Apr 30, 1991

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR

	ABBVIE	EQ 50MG BASE/ML	A075114 001	Jul 26, 1999
	ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE			
	EUROHLTH INTL SARL	EQ 500MG BASE/VIAL	A074885 001	Dec 19, 1997
		EQ 1GM BASE/VIAL	A074885 002	Dec 19, 1997

ACYCLOVIR SODIUM

	APOTHECON	EQ 500MG BASE/VIAL	A074897 001	Feb 27, 1998
		EQ 1GM BASE/VIAL	A074897 002	Feb 27, 1998
	CHARTWELL INJECTABLE	EQ 500MG BASE/VIAL	A074596 002	Apr 22, 1997
		EQ 1GM BASE/VIAL	A074596 001	Apr 22, 1997
	DR REDDYS	EQ 50MG BASE/ML	A207919 001	Jun 17, 2020
	EUROHLTH INTL SARL	EQ 500MG BASE/VIAL	A074913 001	Oct 15, 1997
		EQ 1GM BASE/VIAL	A074913 002	Oct 15, 1997
	FRESENIUS KABI USA	EQ 500MG BASE/VIAL	A075015 001	Apr 30, 1998
	HIKMA	EQ 500MG BASE/VIAL	A205771 001	Feb 29, 2016
		EQ 1GM BASE/VIAL	A205771 002	Feb 29, 2016
	HOSPIRA	EQ 25MG BASE/ML	A074720 001	Apr 22, 1997
		EQ 50MG BASE/ML	A075065 001	Feb 25, 1999
		EQ 500MG BASE/VIAL	A074663 001	Apr 22, 1997
		EQ 500MG BASE/VIAL	A074758 001	Apr 22, 1997
		EQ 1GM BASE/VIAL	A074663 002	Apr 22, 1997
		EQ 1GM BASE/VIAL	A074758 002	Apr 22, 1997
	MYLAN LABS LTD	EQ 500MG BASE/VIAL	A203927 001	Mar 29, 2017
		EQ 1GM BASE/VIAL	A203927 002	Mar 29, 2017
	TEVA PARENTERAL	EQ 50MG BASE/ML	A075627 001	Mar 28, 2001
		EQ 500MG BASE/VIAL	A074969 001	Aug 26, 1997
		EQ 1GM BASE/VIAL	A074969 002	Aug 26, 1997
	ZYDUS PHARMS	EQ 500MG BASE/VIAL	A206606 001	Jun 13, 2017
		EQ 1GM BASE/VIAL	A206606 002	Jun 13, 2017
	ZOVIRAX			
+	GLAXOSMITHKLINE	EQ 250MG BASE/VIAL **	N018603 003	Aug 30, 1983
+		EQ 500MG BASE/VIAL **	N018603 001	Oct 22, 1982
+		EQ 1GM BASE/VIAL **	N018603 002	Jun 29, 1989

ADAPALENE

SOLUTION; TOPICAL

DIFFERIN

+	GALDERMA LABS LP	0.1% **	N020338 001	May 31, 1996
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ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

	ACTAVIS MID ATLANTIC	0.1%; 2.5%	A203790 001	Sep 30, 2015
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ADENOSINE

INJECTABLE; INJECTION

ADENOCARD

+	ASTELLAS	3MG/ML **	N019937 002	Oct 30, 1989
	ADENOSINE			
	AM REGENT	3MG/ML	A090010 001	Apr 28, 2009
	HIKMA	3MG/ML	A076501 001	Jun 16, 2004
	MYLAN LABS LTD	3MG/ML	A078640 001	Mar 21, 2014
	TEVA PHARMS USA	3MG/ML	A076564 001	Jun 16, 2004
		3MG/ML	A078676 001	Jul 31, 2008
	WOCKHARDT	3MG/ML	A090220 001	Jul 20, 2009

SOLUTION; INTRAVENOUS

ADENOSCAN

+	ASTELLAS	60MG/20ML (3MG/ML) **	N020059 001	May 18, 1995
+		90MG/30ML (3MG/ML) **	N020059 002	May 18, 1995

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALATROFLOXACIN MESYLATE

INJECTABLE; INJECTION

TROVAN PRESERVATIVE FREE

PFIZER

EQ 200MG BASE/VIAL

N020760 001 Dec 18, 1997

EQ 300MG BASE/VIAL

N020760 002 Dec 18, 1997

ALBENDAZOLE

TABLET; ORAL

ALBENDAZOLE

CHARTWELL RX

200MG

A211636 001 Jun 10, 2020

CIPLA LTD

200MG

A210434 001 Sep 21, 2018

ALBENZA

+

IMPAX LABS INC

200MG

N020666 001 Jun 11, 1996

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

ARMSTRONG PHARMS

0.09MG/INH

A072273 001 Aug 14, 1996

GENPHARM

0.09MG/INH

A073045 001 Aug 19, 1997

IVAX SUB TEVA PHARMS

0.09MG/INH

A073272 001 Dec 28, 1995

PLIVA

0.09MG/INH

A074072 001 Aug 01, 1996

PROVENTIL

SCHERING

0.09MG/INH

N017559 001

VENTOLIN

GLAXOSMITHKLINE

0.09MG/INH

N018473 001

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

ALBUTEROL SULFATE

PADAGIS US

EQ 0.09MG BASE/INH

A203760 001 Feb 24, 2020

CAPSULE; INHALATION

VENTOLIN ROTACAPS

GLAXOSMITHKLINE

EQ 0.2MG BASE

N019489 001 May 04, 1988

SOLUTION; INHALATION

ACCUNEB

+

MYLAN SPECIALITY LP

EQ 0.021% BASE **

N020949 002 Apr 30, 2001

+

EQ 0.042% BASE **

N020949 001 Apr 30, 2001

ALBUTEROL SULFATE

ACTAVIS MID ATLANTIC

EQ 0.083% BASE

A073533 001 Sep 26, 1995

APOTEX INC

EQ 0.021% BASE

A078623 001 Apr 05, 2010

EQ 0.042% BASE

A078623 002 Apr 05, 2010

EQ 0.083% BASE

A075717 001 Feb 02, 2007

EQ 0.5% BASE

A076391 001 Apr 01, 2003

BAUSCH

EQ 0.083% BASE

A075358 001 Mar 29, 2000

EQ 0.5% BASE

A075050 001 Jun 18, 1998

COPLEY PHARM

EQ 0.083% BASE

A073495 001 May 28, 1993

EQ 0.5% BASE

A073307 001 Nov 27, 1991

EPIC PHARMA LLC

EQ 0.083% BASE

A075063 001 Feb 09, 1999

LANDELA PHARM

EQ 0.083% BASE

A077569 001 Apr 04, 2006

LUOXIN AUROVITAS

EQ 0.021% BASE

A211888 001 Apr 20, 2020

EQ 0.042% BASE

A211888 002 Apr 20, 2020

MYLAN SPECLT

EQ 0.083% BASE **

A072652 001 Feb 21, 1992

ROXANE

EQ 0.083% BASE

A075129 001 Feb 13, 2001

TEVA PHARMS

EQ 0.083% BASE

A075343 001 Nov 09, 1999

WATSON LABS INC

EQ 0.083% BASE

A076370 001 Nov 24, 2003

WOCKHARDT EU OPERATN

EQ 0.083% BASE

A075394 001 Nov 22, 1999

PROVENTIL

+

SCHERING

EQ 0.083% BASE **

N019243 002 Jan 14, 1987

+

EQ 0.5% BASE **

N019243 001 Jan 14, 1987

VENTOLIN

+

GLAXOSMITHKLINE

EQ 0.083% BASE **

N019773 001 Apr 23, 1992

EQ 0.5% BASE **

N019269 002 Jan 16, 1987

SYRUP; ORAL

ALBUTEROL SULFATE

ACTAVIS MID ATLANTIC

EQ 2MG BASE/5ML

A075262 001 Mar 30, 1999

MOVA

EQ 2MG BASE/5ML

A074302 001 Sep 30, 1994

WATSON LABS

EQ 2MG BASE/5ML

A073165 001 Apr 29, 1993

PROVENTIL

+

SCHERING

EQ 2MG BASE/5ML **

N018062 001 Jan 19, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALBUTEROL SULFATE

SYRUP;ORAL

VENTOLIN

GLAXOSMITHKLINE EQ 2MG BASE/5ML ** N019621 001 Jun 10, 1987

TABLET;ORAL

ALBUTEROL SULFATE

AM THERAP EQ 2MG BASE A072449 001 Dec 05, 1989

EQ 4MG BASE A072450 001 Dec 05, 1989

AUROBINDO PHARMA LTD EQ 2MG BASE A213657 001 May 14, 2020

EQ 4MG BASE A213657 002 May 14, 2020

CHARTWELL RX EQ 2MG BASE A072151 001 Dec 05, 1989

EQ 4MG BASE A072151 002 Dec 05, 1989

COPLEY PHARM EQ 2MG BASE A072966 001 Nov 22, 1991

EQ 4MG BASE A072967 001 Nov 22, 1991

EYWA EQ 2MG BASE A213524 001 Oct 08, 2020

EQ 4MG BASE A213524 002 Oct 08, 2020

PLIVA EQ 2MG BASE A072316 001 Dec 05, 1989

EQ 4MG BASE A072317 001 Dec 05, 1989

STRIDES PHARMA EQ 2MG BASE A072860 002 Dec 20, 1989

EQ 4MG BASE A072860 001 Dec 20, 1989

TEVA EQ 2MG BASE A072619 001 Dec 05, 1989

EQ 2MG BASE A072779 001 Jun 25, 1993

EQ 2MG BASE A072938 001 Mar 30, 1990

EQ 4MG BASE A072620 001 Dec 05, 1989

EQ 4MG BASE A072780 001 Jun 25, 1993

EQ 4MG BASE A072939 001 Mar 30, 1990

UCB INC EQ 2MG BASE A073120 001 Sep 29, 1992

EQ 4MG BASE A073121 001 Sep 29, 1992

WARNER CHILCOTT EQ 2MG BASE A072817 001 Jan 09, 1990

EQ 4MG BASE A072818 001 Jan 09, 1990

WATSON LABS EQ 2MG BASE A072629 001 Jan 31, 1991

EQ 2MG BASE A072764 001 Aug 28, 1991

EQ 4MG BASE A072630 001 Jan 31, 1991

EQ 4MG BASE A072765 001 Aug 28, 1991

PROVENTIL

+ SCHERING EQ 2MG BASE ** N017853 001 May 07, 1982

+ EQ 4MG BASE ** N017853 002 May 07, 1982

VENTOLIN

GLAXOSMITHKLINE EQ 2MG BASE N019112 001 Jul 10, 1986

EQ 4MG BASE N019112 002 Jul 10, 1986

TABLET, EXTENDED RELEASE;ORAL

ALBUTEROL SULFATE

RISING EQ 4MG BASE A078092 002 Jan 29, 2007

EQ 8MG BASE A078092 001 Jan 29, 2007

PROVENTIL

SCHERING EQ 4MG BASE N019383 001 Jul 13, 1987

VOLMAX

+ MURO EQ 4MG BASE N019604 002 Dec 23, 1992

+ EQ 8MG BASE N019604 001 Dec 23, 1992

VOSPIRE ER

STRIDES PHARMA EQ 4MG BASE A076130 002 Sep 26, 2002

EQ 8MG BASE A076130 003 Sep 26, 2002

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

COMBIVENT

BOEHRINGER INGELHEIM EQ 0.09MG BASE/INH;0.018MG/INH N020291 001 Oct 24, 1996

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

APOTEX INC EQ 0.083% BASE;0.017% A077117 001 Dec 31, 2007

FOSUN PHARMA EQ 0.083% BASE;0.017% A076867 001 Dec 21, 2006

LUOXIN AUROVITAS EQ 0.083% BASE;0.017% A206532 001 Jul 08, 2020

TEVA PHARMS EQ 0.083% BASE;0.017% A076724 001 Dec 31, 2007

WATSON LABS TEVA EQ 0.083% BASE;0.017% A077063 001 Dec 31, 2007

DUONEB

+ MYLAN SPECIALITY LP EQ 0.083% BASE;0.017% ** N020950 001 Mar 21, 2001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALCAFTADINE

SOLUTION/DROPS;OPHTHALMIC

ALCAFTADINE

EUGIA PHARMA

0.25%

A210659 001 Jun 23, 2023

ALCLOMETASONE DIPROPIONATE

CREAM;TOPICAL

ACLOVATE

+ FOUGERA PHARMS

0.05% **

N018707 001 Dec 14, 1982

OINTMENT;TOPICAL

ACLOVATE

+ FOUGERA PHARMS

0.05% **

N018702 001 Dec 14, 1982

ALENDRONATE SODIUM

SOLUTION;ORAL

FOSAMAX

+ MERCK

EQ 70MG BASE/75ML **

N021575 001 Sep 17, 2003

TABLET;ORAL

ALENDRONATE SODIUM

CHARTWELL RX

EQ 5MG BASE

A075871 001 Apr 22, 2009

EQ 5MG BASE

A079049 003 Aug 04, 2008

EQ 10MG BASE

A075871 002 Apr 22, 2009

EQ 10MG BASE

A079049 004 Aug 04, 2008

EQ 35MG BASE

A075871 004 Apr 22, 2009

EQ 35MG BASE

A079049 001 Aug 04, 2008

EQ 40MG BASE

A075871 003 Apr 22, 2009

EQ 70MG BASE

A075871 005 Apr 22, 2009

EQ 70MG BASE

A079049 002 Aug 04, 2008

IMPAX LABS INC

EQ 5MG BASE

A075710 001 Feb 06, 2008

EQ 10MG BASE

A075710 002 Feb 06, 2008

EQ 35MG BASE

A075710 003 Feb 06, 2008

EQ 40MG BASE

A075710 004 Feb 06, 2008

EQ 70MG BASE

A075710 005 Feb 06, 2008

JUBILANT CADISTA

EQ 5MG BASE

A090557 001 Feb 18, 2010

EQ 10MG BASE

A090557 002 Feb 18, 2010

EQ 35MG BASE

A090557 003 Feb 18, 2010

EQ 70MG BASE

A090557 004 Feb 18, 2010

MYLAN

EQ 35MG BASE

A078638 001 Aug 04, 2008

EQ 70MG BASE

A078638 002 Aug 04, 2008

RISING

EQ 5MG BASE

A076584 001 Aug 04, 2008

EQ 10MG BASE

A076584 002 Aug 04, 2008

EQ 35MG BASE

A076584 003 Aug 04, 2008

EQ 70MG BASE

A076584 004 Aug 04, 2008

TEVA PHARMS

EQ 35MG BASE

A076184 002 Aug 04, 2008

EQ 70MG BASE

A076184 001 Feb 06, 2008

FOSAMAX

+ ORGANON

EQ 5MG BASE **

N020560 003 Apr 25, 1997

+

EQ 10MG BASE **

N020560 001 Sep 29, 1995

+

EQ 35MG BASE **

N020560 004 Oct 20, 2000

+

EQ 40MG BASE **

N020560 002 Sep 29, 1995

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALFUZOSIN HYDROCHLORIDE

CHARTWELL RX

10MG

A079056 001 Jul 18, 2011

MYLAN

10MG

A079014 001 Jul 18, 2011

TORRENT PHARMS

10MG

A079054 001 Jul 18, 2011

WOCKHARDT

10MG

A090221 001 Aug 10, 2012

ALISKIREN HEMIFUMARATE

CAPSULE, PELLET;ORAL

TEKTURNA

+ NODEN PHARMA

EQ 37.5MG BASE

N210709 001 Nov 14, 2017

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE

TABLET;ORAL

TEKAMLO

NOVARTIS

EQ 150MG BASE;EQ 5MG BASE

N022545 001 Aug 26, 2010

EQ 150MG BASE;EQ 10MG BASE

N022545 002 Aug 26, 2010

EQ 300MG BASE;EQ 5MG BASE

N022545 003 Aug 26, 2010

EQ 300MG BASE;EQ 10MG BASE

N022545 004 Aug 26, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMTURNIDE

NOVARTIS	EQ 150MG BASE;EQ 5MG BASE;12.5MG	N200045 001	Dec 21, 2010
	EQ 300MG BASE;EQ 5MG BASE;12.5MG	N200045 002	Dec 21, 2010
	EQ 300MG BASE;EQ 5MG BASE;25MG	N200045 003	Dec 21, 2010
	EQ 300MG BASE;EQ 10MG BASE;12.5MG	N200045 004	Dec 21, 2010
	EQ 300MG BASE;EQ 10MG BASE;25MG	N200045 005	Dec 21, 2010

ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEKTURNA HCT

+ NODEN PHARMA	EQ 150MG BASE;12.5MG **	N022107 001	Jan 18, 2008
+	EQ 150MG BASE;25MG **	N022107 002	Jan 18, 2008
+	EQ 300MG BASE;12.5MG **	N022107 003	Jan 18, 2008
+	EQ 300MG BASE;25MG **	N022107 004	Jan 18, 2008

ALISKIREN HEMIFUMARATE; VALSARTAN

TABLET; ORAL

VALTURNA

NOVARTIS	EQ 150MG BASE;160MG	N022217 001	Sep 16, 2009
	EQ 300MG BASE;320MG	N022217 002	Sep 16, 2009

ALKAVERVIR

TABLET; ORAL

VERILOID

3M	2MG	N007336 002	
	3MG	N007336 003	

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

AIPING PHARM INC	100MG	A070268 001	Dec 31, 1985
MUTUAL PHARM	100MG	A070466 001	Dec 24, 1985
	300MG	A070467 001	Dec 24, 1985
PURACAP PHARM	100MG	A070150 001	Dec 10, 1985
	300MG	A070147 001	Dec 10, 1985
PUREPAC PHARM	100MG	A070579 001	Apr 14, 1986
	300MG	A070580 001	Apr 14, 1986
SANDOZ	300MG	A070269 001	Dec 31, 1985
SUN PHARM INDS INC	100MG	A078390 001	Aug 30, 2007
	300MG	A078390 002	Aug 30, 2007
SUPERPHARM	100MG	A070950 001	Nov 30, 1988
	300MG	A070951 001	Nov 30, 1988
WATSON LABS	100MG	N018241 001	Nov 16, 1984
	100MG	N018785 001	Sep 28, 1984
	300MG	N018241 002	Nov 16, 1984
	300MG	N018785 002	Sep 28, 1984
LOPURIN			
ABBOTT	100MG	N018297 001	
	300MG	N018297 002	

ALLOPURINOL; LESINURAD

TABLET; ORAL

DUZALLO

+ IRONWOOD PHARMS INC	200MG;200MG	N209203 001	Aug 18, 2017
+	300MG;200MG	N209203 002	Aug 18, 2017

ALMOTRIPTAN MALATE

TABLET; ORAL

AXERT

+ JANSSEN PHARMS	EQ 6.25MG BASE **	N021001 001	May 07, 2001
+	EQ 12.5MG BASE **	N021001 002	May 07, 2001

ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

OSENİ

+ TAKEDA PHARMS USA	EQ 12.5MG BASE;EQ 15MG BASE **	N022426 004	Jan 25, 2013
+	EQ 12.5MG BASE;EQ 45MG BASE **	N022426 006	Jan 25, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALOSETRON HYDROCHLORIDE

TABLET;ORAL

ALOSETRON HYDROCHLORIDE

EYWA PHARMA	EQ 0.5MG BASE	A211621 001	Sep 16, 2019
	EQ 1MG BASE	A211621 002	Sep 16, 2019
HIKMA	EQ 0.5MG BASE	A200652 001	May 04, 2015
	EQ 1MG BASE	A200652 002	May 04, 2015
PAR PHARM INC	EQ 0.5MG BASE	A206113 001	Feb 23, 2018
	EQ 1MG BASE	A206113 002	Feb 23, 2018

ALPRAZOLAM

SOLUTION;ORAL

ALPRAZOLAM

ROXANE	0.5MG/5ML	A074314 001	Oct 31, 1993
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TABLET;ORAL

ALPRAZOLAM

ANI PHARMS	0.25MG	A074085 001	Feb 16, 1994
	0.5MG	A074085 002	Feb 16, 1994
	1MG	A074085 003	Feb 16, 1994
	2MG	A074085 004	Feb 26, 1996
IVAX SUB TEVA PHARMS	0.25MG	A074294 001	Jul 29, 1994
	0.5MG	A074294 002	Jul 29, 1994
	1MG	A074294 003	Jul 29, 1994
	2MG	A074294 004	Jul 29, 1994
MYLAN	0.25MG	A074215 001	Jan 27, 1994
	0.5MG	A074215 002	Jan 27, 1994
	1MG	A074215 003	Jan 27, 1994
	2MG	A074215 004	Jan 27, 1994
MYLAN PHARMS INC	0.25MG	A074046 001	Oct 19, 1993
	0.5MG	A074046 002	Oct 19, 1993
	1MG	A074046 003	Oct 19, 1993
	2MG	A074046 004	May 07, 1997
OXFORD PHARMS	0.25MG	A078491 001	Sep 25, 2008
	0.5MG	A078491 002	Sep 25, 2008
	1MG	A078491 003	Sep 25, 2008
	2MG	A078491 004	Dec 12, 2008
ROXANE	0.25MG	A074199 001	Oct 19, 1993
	0.5MG	A074199 002	Oct 19, 1993
	1MG	A074199 003	Oct 19, 1993
WATSON LABS	0.25MG	A074456 001	Aug 31, 1995
	0.25MG	A074479 001	Jan 21, 1997
	0.5MG	A074456 002	Aug 31, 1995
	0.5MG	A074479 002	Jan 21, 1997
	1MG	A074456 003	Aug 31, 1995
	1MG	A074479 003	Jan 21, 1997

TABLET, EXTENDED RELEASE;ORAL

ALPRAZOLAM

ACTAVIS LABS FL INC	0.5MG	A077198 001	May 13, 2010
	1MG	A077198 002	May 13, 2010
	2MG	A077198 003	May 13, 2010
	3MG	A077198 004	May 13, 2010
ANI PHARMS	0.5MG	A077725 001	Jul 31, 2006
	0.5MG	A077979 001	Feb 28, 2007
	1MG	A077725 002	Jul 31, 2006
	1MG	A077979 002	Feb 28, 2007
	2MG	A077725 004	Jul 31, 2006
	2MG	A077979 003	Feb 28, 2007
	3MG	A077725 003	Jul 31, 2006
	3MG	A077979 004	Feb 28, 2007
HERITAGE PHARMS INC	0.5MG	A078489 001	Oct 17, 2008
	1MG	A078489 002	Oct 17, 2008
	2MG	A078489 003	Oct 17, 2008
	3MG	A078489 004	Oct 17, 2008
IMPAX LABS	0.5MG	A077968 004	May 24, 2007
	1MG	A077968 003	May 24, 2007
	2MG	A077968 002	May 24, 2007
	3MG	A077968 001	May 24, 2007
IMPAX LABS INC	0.5MG	A077996 001	Jan 31, 2007
	1MG	A077996 002	Jan 31, 2007
	2MG	A077996 003	Jan 31, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALPRAZOLAMTABLET, EXTENDED RELEASE;ORAL
ALPRAZOLAM

	3MG	A077996 004	Jan 31, 2007
MYLAN	0.5MG	A077391 002	Jan 26, 2006
	1MG	A077391 003	Jan 26, 2006
	2MG	A077391 004	Jan 26, 2006
	3MG	A077391 001	Jan 26, 2006
SANDOZ INC	0.5MG	A077777 001	Jun 30, 2006
	1MG	A077777 002	Jun 30, 2006
	2MG	A077777 003	Jun 30, 2006
	3MG	A077777 004	Jun 30, 2006
VINTAGE PHARMS	0.5MG	A078442 001	Oct 15, 2007
	1MG	A078442 002	Oct 15, 2007
	2MG	A078442 003	Oct 15, 2007
	3MG	A078442 004	Oct 15, 2007

TABLET, ORALLY DISINTEGRATING;ORAL

NIRAVAM

+	UCB INC	0.25MG **	N021726 001	Jan 19, 2005
+		0.5MG **	N021726 002	Jan 19, 2005
+		1MG **	N021726 003	Jan 19, 2005
+		2MG **	N021726 004	Jan 19, 2005

ALPROSTADIL

INJECTABLE;INJECTION

CAVERJECT

+	PFIZER	0.005MG/VIAL	N020379 003	Jun 27, 1996
		0.005MG/ML	N020755 001	Oct 31, 1997
		0.01MG/ML	N020755 002	Oct 01, 1997
		0.02MG/ML	N020755 003	Oct 01, 1997

EDEX

	AUXILIUM PHARMS LLC	0.005MG/VIAL	N020649 001	Jun 12, 1997
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SUPPOSITORY;URETHRAL

MUSE

+	MYLAN SPECIALITY LP	0.125MG	N020700 001	Nov 19, 1996
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ALSEROXYLON

TABLET;ORAL

RAUTENSIN

	NOVARTIS	2MG	N009215 001	
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RAUWILOID

	3M	2MG	N008867 001	
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ALTRETAMINE

CAPSULE;ORAL

HEXALEN

+	EISAI INC	50MG	N019926 001	Dec 26, 1990
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ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE;ORAL

ALUMINUM HYDROXIDE AND MAGNESIUM TRISILICATE

	PENNEK	80MG;20MG	A089449 001	Nov 27, 1987
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FOAMCOAT

	GUARDIAN DRUG	80MG;20MG	A071793 001	Sep 04, 1987
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FOAMICON

	NOVARTIS	80MG;20MG	A072687 001	Jun 28, 1989
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GAVISCON

+	CHATTEM SANOFI	80MG;20MG **	N018685 001	Dec 09, 1983
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+		160MG;40MG **	N018685 002	Dec 09, 1983
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ALVIMOPAN

CAPSULE;ORAL

ENTEREG

+	CUBIST PHARMS	12MG	N021775 001	May 20, 2008
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AMANTADINE HYDROCHLORIDE

CAPSULE;ORAL

AMANTADINE HYDROCHLORIDE

	ACTAVIS ELIZABETH	100MG	A077659 001	Feb 23, 2006
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	CHARTWELL MOLECULAR	100MG	A209221 001	Jun 15, 2017
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	INVAGEN PHARMS	100MG	A207570 001	Sep 30, 2016
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	WATSON LABS	100MG	A071382 001	Jan 21, 1987
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

WATSON LABS INC

100MG

A208107 001 Dec 06, 2016

SYMADINE

SOLVAY

100MG

A071000 001 Sep 04, 1986

SYMMETREL

+ ENDO PHARMS

100MG **

N016020 001

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

ANDA REPOSITORY

50MG/5ML

A076352 001 Sep 10, 2004

G AND W LABS INC

50MG/5ML

A072655 001 Oct 30, 1990

TEVA PHARMS

50MG/5ML

A073115 001 Aug 23, 1991

VINTAGE

50MG/5ML

A077992 001 Dec 12, 2006

XTTRIUM LABS INC

50MG/5ML

A075060 001 Dec 24, 1998

SYMMETREL

+ ENDO PHARMS

50MG/5ML **

N016023 002

TABLET; ORAL

AMANTADINE HYDROCHLORIDE

EDENBRIDGE PHARMS

100MG

A212407 001 May 27, 2022

INVAGEN PHARMS

100MG

A207571 001 Jan 31, 2017

JUBILANT GENERICS

100MG

A210403 001 Feb 07, 2018

SYMMETREL

+ ENDO PHARMS

100MG **

N018101 001

TABLET, EXTENDED RELEASE; ORAL

OSMOLEX ER

+ ADAMAS OPERATIONS

EQ 161MG BASE

N209410 004 Apr 22, 2020

+

EQ 258MG BASE

N209410 003 Feb 16, 2018

AMBENONIUM CHLORIDE

TABLET; ORAL

MYTELASE

SANOFI AVENTIS US

10MG

N010155 002

AMCINONIDE

CREAM; TOPICAL

CYCLOCORT

+ ASTELLAS

0.025% **

N018116 001

+

0.1% **

N018116 002

LOTION; TOPICAL

CYCLOCORT

+ ASTELLAS

0.1%

N019729 001 Jun 13, 1988

OINTMENT; TOPICAL

CYCLOCORT

+ ASTELLAS

0.1% **

N018498 001

AMDINOCILLIN

INJECTABLE; INJECTION

COACTIN

ROCHE

250MG/VIAL

N050565 001 Dec 21, 1984

500MG/VIAL

N050565 002 Dec 21, 1984

1GM/VIAL

N050565 003 Dec 21, 1984

AMIFOSTINE

INJECTABLE; INJECTION

AMIFOSTINE

EUGIA PHARMA SPECLTS

500MG/VIAL

A204363 001 Jul 17, 2017

SUN PHARM

500MG/VIAL

A077126 001 Mar 14, 2008

ETHYOL

CLINIGEN

375MG/VIAL

N020221 002 Sep 10, 1999

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

ABBOTT

EQ 250MG BASE/ML

A063265 001 Nov 30, 1994

EQ 250MG BASE/ML

A063266 001 Oct 31, 1994

FRESENIUS KABI USA

EQ 50MG BASE/ML

A205605 001 Dec 09, 2015

HIKMA

EQ 50MG BASE/ML

A063274 001 May 18, 1992

EQ 250MG BASE/ML

A063275 001 May 18, 1992

HOSPIRA

EQ 50MG BASE/ML

A063263 001 Nov 30, 1994

EQ 50MG BASE/ML

A063350 001 Jul 30, 1993

EQ 62.5MG BASE/ML

A063283 001 Oct 31, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

	EQ 250MG BASE/ML	A063264	001	Nov 30, 1994
	EQ 250MG BASE/ML	A063350	002	Jul 30, 1993
	EQ 250MG BASE/ML	A064098	001	Jun 26, 1995
	EQ 250MG BASE/ML	A064099	001	Jun 20, 1995
IGI LABS INC	EQ 50MG BASE/ML	A063167	001	Dec 14, 1995
	EQ 250MG BASE/ML	A063169	001	Dec 14, 1995
MEITHEAL	EQ 50MG BASE/ML	A064045	001	Sep 28, 1993
AMIKACIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
HOSPIRA	EQ 500MG BASE/100ML	A064146	001	Apr 02, 1997
AMIKIN				
APOTHECON	EQ 50MG BASE/ML	A062311	001	
	EQ 50MG BASE/ML	A062562	001	Sep 20, 1984
+	EQ 50MG BASE/ML **	N050495	001	
	EQ 250MG BASE/ML	A062311	002	
	EQ 250MG BASE/ML	A062562	002	Sep 20, 1984
+	EQ 250MG BASE/ML **	N050495	002	
AMIKIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
APOTHECON	EQ 5MG BASE/ML	N050618	002	Nov 30, 1987
	EQ 10MG BASE/ML	N050618	001	Nov 30, 1987

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

CHARTWELL RX	5MG	A204180	001	Aug 07, 2015
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AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	EQ 5MG ANHYDROUS; 50MG	A073357	001	Nov 27, 1991
TEVA	EQ 5MG ANHYDROUS; 50MG	A070795	001	Apr 17, 1988
WATSON LABS	EQ 5MG ANHYDROUS; 50MG	A073334	001	Jul 19, 1991
HYDRO-RIDE				
PAR PHARM	EQ 5MG ANHYDROUS; 50MG	A070347	001	Dec 25, 1990
MODURETIC 5-50				
+	MERCK	EQ 5MG ANHYDROUS; 50MG **	N018201	001

AMINO ACIDS

INJECTABLE; INJECTION

AMINESS 5.2% ESSENTIAL AMINO ACIDS W/ HISTADINE

HOSPIRA	5.2% (5.2GM/100ML)	N018901	001	Apr 06, 1984
AMINOSYN 10%				
ICU MEDICAL INC	10% (10GM/100ML)	N017673	003	
AMINOSYN 10% (PH6)				
ICU MEDICAL INC	10% (10GM/100ML)	N017673	008	Nov 18, 1985
AMINOSYN 3.5%				
ICU MEDICAL INC	3.5% (3.5GM/100ML)	N017789	004	
AMINOSYN 3.5% IN PLASTIC CONTAINER				
ABBOTT	3.5% (3.5GM/100ML)	N018804	001	May 15, 1984
	3.5% (3.5GM/100ML)	N018875	001	Aug 08, 1984
AMINOSYN 5%				
ICU MEDICAL INC	5% (5GM/100ML)	N017673	001	
AMINOSYN 7%				
ICU MEDICAL INC	7% (7GM/100ML)	N017673	002	
AMINOSYN 7% (PH6)				
ICU MEDICAL INC	7% (7GM/100ML)	N017673	006	Nov 18, 1985
AMINOSYN 8.5%				
ICU MEDICAL INC	8.5% (8.5GM/100ML)	N017673	004	
AMINOSYN 8.5% (PH6)				
ICU MEDICAL INC	8.5% (8.5GM/100ML)	N017673	007	Nov 18, 1985
AMINOSYN II 10%				
ICU MEDICAL INC	10% (10GM/100ML)	N019438	005	Apr 03, 1986
AMINOSYN II 3.5%				
ICU MEDICAL INC	3.5% (3.5GM/100ML)	N019438	001	Apr 03, 1986
AMINOSYN II 3.5% IN PLASTIC CONTAINER				
ABBOTT	3.5% (3.5GM/100ML)	N019491	001	Oct 10, 1986
AMINOSYN II 5%				
ICU MEDICAL INC	5% (5GM/100ML)	N019438	002	Apr 03, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS

INJECTABLE; INJECTION

AMINOSYN II 7%			
ICU MEDICAL INC	7% (7GM/100ML)	N019438 003	Apr 03, 1986
AMINOSYN II 8.5%			
ICU MEDICAL INC	8.5% (8.5GM/100ML)	N019438 004	Apr 03, 1986
AMINOSYN-HBC 7%			
ICU MEDICAL INC	7% (7GM/100ML)	N019374 001	Jul 12, 1985
AMINOSYN-HBC 7% IN PLASTIC CONTAINER			
ABBOTT	7% (7GM/100ML)	N019400 001	Jul 23, 1986
AMINOSYN-HF 8%			
ICU MEDICAL INC	8% (8GM/100ML)	A020345 001	Apr 04, 1996
AMINOSYN-RF 5.2%			
ICU MEDICAL INC	5.2% (5.2GM/100ML)	N018429 001	
BRANCHAMIN 4%			
BAXTER HLTHCARE	4% (4GM/100ML)	N018678 001	Sep 28, 1984
BRANCHAMIN 4% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	4% (4GM/100ML)	N018684 001	Sep 28, 1984
FREAMINE 8.5%			
B BRAUN	8.5% (8.5GM/100ML)	N016822 001	
FREAMINE HBC 6.9%			
B BRAUN	6.9% (6.9GM/100ML)	N016822 006	May 17, 1983
FREAMINE II 8.5%			
B BRAUN	8.5% (8.5GM/100ML)	N016822 002	
FREAMINE III 10%			
B BRAUN	10% (10GM/100ML)	N016822 005	
FREAMINE III 8.5%			
B BRAUN	8.5% (8.5GM/100ML)	N016822 004	
HEPATAMINE 8%			
B BRAUN	8% (8GM/100ML)	N018676 001	Aug 03, 1982
HEPATASOL 8%			
BAXTER HLTHCARE	8% (8GM/100ML)	A020360 001	Apr 04, 1996
NEOPHAM 6.4%			
HOSPIRA	6.4% (6.4GM/100ML)	N018792 001	Jan 17, 1984
NEPHRAMINE 5.4%			
B BRAUN	5.4% (5.4GM/100ML)	N017766 001	
NOVAMINE 11.4%			
HOSPIRA INC	11.4% (11.4GM/100ML)	N017957 003	Aug 09, 1982
NOVAMINE 15%			
HOSPIRA INC	15% (75GM/500ML)	N017957 004	Nov 28, 1986
NOVAMINE 15% SULFITE FREE IN PLASTIC CONTAINER			
BAXTER HLTHCARE	15% (15GM/100ML) **	N020107 001	Feb 05, 1993
NOVAMINE 8.5%			
HOSPIRA INC	8.5% (8.5GM/100ML)	N017957 002	Aug 09, 1982
RENAMIN W/O ELECTROLYTES			
BAXTER HLTHCARE	6.5% (6.5GM/100ML)	N017493 007	Oct 15, 1982
TRAVASOL 10% W/O ELECTROLYTES			
BAXTER HLTHCARE	10% (10GM/100ML)	N017493 006	
TRAVASOL 5.5% W/O ELECTROLYTES			
BAXTER HLTHCARE	5.5% (5.5GM/100ML)	N017493 004	
TRAVASOL 8.5% W/O ELECTROLYTES			
BAXTER HLTHCARE	8.5% (8.5GM/100ML)	N017493 005	

AMINO ACIDS; CALCIUM ACETATE; GLYCERIN; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PROCALAMINE			
B BRAUN	3%; 26MG/100ML; 3GM/100ML; 54MG/100ML; 41MG /100ML; 150MG/100ML; 200MG/100ML; 120MG/100ML	N018582 001	May 08, 1982

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER			
ABBOTT	3.5%; 36.8MG/100ML; 25GM/100ML; 51MG/100ML ; 22.4MG/100ML; 261MG/100ML; 205MG/100ML	N019714 001	Sep 12, 1988
HOSPIRA INC	3.5%; 36.8MG/100ML; 25GM/100ML; 51MG/100ML ; 22.4MG/100ML; 261MG/100ML; 205MG/100ML	N019683 001	Nov 07, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	4.25%;36.8MG/100ML;20GM/100ML;51MG/100ML	N019714 002	Sep 12, 1988	
	L;22.4MG/100ML;261MG/100ML;205MG/100ML			
HOSPIRA INC				
	4.25%;36.8MG/100ML;20GM/100ML;51MG/100ML	N019683 002	Nov 07, 1988	
	L;22.4MG/100ML;261MG/100ML;205MG/100ML			
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	4.25%;36.8MG/100ML;25GM/100ML;51MG/100ML	N019714 004	Sep 12, 1988	
	L;22.4MG/100ML;261MG/100ML;205MG/100ML			
HOSPIRA INC				
	4.25%;36.8MG/100ML;25GM/100ML;51MG/100ML	N019683 003	Nov 07, 1988	
	L;22.4MG/100ML;261MG/100ML;205MG/100ML			
AMINOSYN II 5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2	N019714 003	Sep 12, 1988	
	2.4MG/100ML;261MG/100ML;205MG/100ML			
HOSPIRA INC				
	5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2	N019683 004	Nov 07, 1988	
	2.4MG/100ML;261MG/100ML;205MG/100ML			

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

AMINOSYN 3.5% W/ DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	3.5%;25GM/100ML	N019118 001	Oct 11, 1984	
AMINOSYN 3.5% W/ DEXTROSE 5% IN PLASTIC CONTAINER				
ABBOTT	3.5%;5GM/100ML	N019120 001	Oct 11, 1984	
AMINOSYN 4.25% W/ DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	4.25%;25GM/100ML	N019119 001	Oct 11, 1984	
AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	3.5%;25GM/100ML	N019505 002	Nov 07, 1986	
	3.5%;25GM/100ML	N019713 006	Sep 09, 1988	
HOSPIRA				
	3.5%;25GM/100ML	N019681 001	Nov 01, 1988	
AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER				
ABBOTT	3.5%;5GM/100ML	N019506 001	Nov 07, 1986	
	3.5%;5GM/100ML	N019713 002	Sep 09, 1988	
HOSPIRA				
	3.5%;5GM/100ML	N019681 002	Nov 01, 1988	
AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER				
ABBOTT	4.25%;10GM/100ML	N019713 001	Sep 09, 1988	
HOSPIRA				
	4.25%;10GM/100ML	N019681 004	Nov 01, 1988	
AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER				
ABBOTT	4.25%;20GM/100ML	N019713 004	Sep 09, 1988	
HOSPIRA				
	4.25%;20GM/100ML	N019681 005	Nov 01, 1988	
AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	4.25%;25GM/100ML	N019504 002	Nov 07, 1986	
	4.25%;25GM/100ML	N019713 005	Sep 09, 1988	
HOSPIRA				
	4.25%;25GM/100ML	N019681 003	Nov 01, 1988	
AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	5%;25GM/100ML	N019565 001	Dec 17, 1986	
	5%;25GM/100ML	N019713 003	Sep 09, 1988	
HOSPIRA				
	5%;25GM/100ML	N019681 006	Nov 01, 1988	
TRAVASOL 2.75% IN DEXTROSE 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;10GM/100ML	N019520 002	Sep 23, 1988	
TRAVASOL 2.75% IN DEXTROSE 15% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;15GM/100ML	N019520 003	Sep 23, 1988	
TRAVASOL 2.75% IN DEXTROSE 20% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;20GM/100ML	N019520 004	Sep 23, 1988	
TRAVASOL 2.75% IN DEXTROSE 25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;25GM/100ML	N019520 005	Sep 23, 1988	
TRAVASOL 2.75% IN DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;5GM/100ML	N019520 001	Sep 23, 1988	
TRAVASOL 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;10GM/100ML	N019520 007	Sep 23, 1988	
TRAVASOL 4.25% IN DEXTROSE 15% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;15GM/100ML	N019520 008	Sep 23, 1988	
TRAVASOL 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;20GM/100ML	N019520 009	Sep 23, 1988	
TRAVASOL 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;25GM/100ML	N019520 010	Sep 23, 1988	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

TRAVASOL 4.25% IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%; 5GM/100ML

N019520 006 Sep 23, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECT AND ADJUSTED PHOSPHATE IN DEXTROSE 10% IN PLASTIC CONTAINER

ABBOTT 4.25%; 10GM/100ML; 51MG/100ML; 176.5MG/100
ML; 22.4MG/100ML; 104.5MG/100ML; 205MG/100
ML

N019712 002 Sep 08, 1988

HOSPIRA INC

4.25%; 10GM/100ML; 51MG/100ML; 176.5MG/100
ML; 22.4MG/100ML; 104.5MG/100ML; 205MG/100
ML

N019682 003 Nov 01, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT 3.5%; 25GM/100ML; 51MG/100ML; 22.4MG/100ML
; 261MG/100ML; 205MG/100ML

N019564 002 Dec 16, 1986

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT 4.25%; 25GM/100ML; 51MG/100ML; 22.4MG/100M
L; 261MG/100ML; 205MG/100ML

N019564 004 Dec 16, 1986

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT 3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 12
0MG/100ML; 49.3MG/100ML

N019564 001 Dec 16, 1986

3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 12
0MG/100ML; 49.3MG/100ML

N019712 001 Sep 08, 1988

HOSPIRA INC

3.5%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 12
0MG/100ML; 49.3MG/100ML

N019682 001 Nov 01, 1988

AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER

ABBOTT 4.25%; 10GM/100ML; 30MG/100ML; 97MG/100ML;
120MG/100ML; 49.3MG/100ML

N019564 003 Dec 16, 1986

HOSPIRA INC

4.25%; 5GM/100ML; 30MG/100ML; 97MG/100ML; 1
20MG/100ML; 49.3MG/100ML

N019682 002 Nov 01, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%; 10GM/100ML; 51MG/100ML; 261MG/100ML
; 216MG/100ML; 112MG/100ML

N020147 002 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%; 15GM/100ML; 51MG/100ML; 261MG/100ML
; 216MG/100ML; 112MG/100ML

N020147 003 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%; 20GM/100ML; 51MG/100ML; 261MG/100ML
; 216MG/100ML; 112MG/100ML

N020147 004 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%; 25GM/100ML; 51MG/100ML; 261MG/100ML
; 216MG/100ML; 112MG/100ML

N020147 005 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%; 5GM/100ML; 51MG/100ML; 261MG/100ML;
216MG/100ML; 112MG/100ML

N020147 001 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%; 10GM/100ML; 51MG/100ML; 261MG/100ML
; 297MG/100ML; 77MG/100ML

N020147 007 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%; 15GM/100ML; 51MG/100ML; 261MG/100ML
; 297MG/100ML; 77MG/100ML

N020147 008 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%; 20GM/100ML; 51MG/100ML; 261MG/100ML
; 297MG/100ML; 77MG/100ML

N020147 009 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%; 25GM/100ML; 51MG/100ML; 261MG/100ML
; 297MG/100ML; 77MG/100ML

N020147 010 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%; 5GM/100ML; 51MG/100ML; 261MG/100ML;
297MG/100ML; 77MG/100ML

N020147 006 Oct 23, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

FREAMINE III 8.5% W/ ELECTROLYTES

B BRAUN

8.5%; 110MG/100ML; 230MG/100ML; 10MG/100ML
; 440MG/100ML; 690MG/100ML

N016822 007 Jul 01, 1988

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

ICU MEDICAL INC

3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML;
234MG/100ML

N017789 003

AMINOSYN 3.5% M IN PLASTIC CONTAINER

ABBOTT

3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML;
234MG/100ML

N018804 002 May 15, 1984

3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML;
234MG/100ML

N018875 002 Aug 08, 1984

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

FREAMINE III 3% W/ ELECTROLYTES

B BRAUN

3%; 54MG/100ML; 40MG/100ML; 150MG/100ML; 20
0MG/100ML; 120MG/100ML

N016822 003

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

ICU MEDICAL INC

3.5%; 21MG/100ML; 128MG/100ML; 234MG/100ML

N017789 005

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN PLASTIC CONTAINER

ABBOTT

3.5%; 32MG/100ML; 128MG/100ML; 222MG/100ML
; 49MG/100ML

N019493 001 Oct 16, 1986

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

VEINAMINE 8%

HOSPIRA INC

8%; 61MG/100ML; 211MG/100ML; 56MG/100ML; 38
8MG/100ML

N017957 001

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 10% W/ ELECTROLYTES

ICU MEDICAL INC

10%; 102MG/100ML; 45MG/100ML; 522MG/100ML;
410MG/100ML

N019437 004 Apr 03, 1986

AMINOSYN II 7% W/ ELECTROLYTES

ICU MEDICAL INC

7%; 102MG/100ML; 45MG/100ML; 522MG/100ML; 4
10MG/100ML

N019437 006 Apr 03, 1986

AMINOSYN II 8.5% W/ ELECTROLYTES

ICU MEDICAL INC

8.5%; 102MG/100ML; 45MG/100ML; 522MG/100ML
; 410MG/100ML

N019437 005 Apr 03, 1986

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC

INJECTABLE; INJECTION

AMINOSYN 8.5% W/ELECTROLYTES

ICU MEDICAL INC

8.5%; 102MG/100ML; 487MG/100ML; 28MG/100ML
; 425MG/100ML

N017673 009 Oct 25, 2002

AMINOSYN II 8.5% W/ELECTROLYTES

ICU MEDICAL INC

8.5%; 102MG/100ML; 492MG/100ML; 60MG/100ML
; 425MG/100ML

N019437 008 Oct 25, 2002

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M

ICU MEDICAL INC

3.5%; 30MG/100ML; 97MG/100ML; 120MG/100ML;
49MG/100ML

N019437 007 Apr 03, 1986

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER					
BAXTER HLTHCARE	3.5%;51MG/100ML;131MG/100ML;218MG/100ML;35MG/100ML	N020177	001	Oct 23, 1995	
TRAVASOL 3.5% W/ ELECTROLYTES					
BAXTER HLTHCARE	3.5%;51MG/100ML;131MG/100ML;218MG/100ML;35MG/100ML	N017493	003		
TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER					
BAXTER HLTHCARE	5.5%;102MG/100ML;522MG/100ML;431MG/100ML;224MG/100ML	N020173	001	Oct 27, 1995	
TRAVASOL 5.5% W/ ELECTROLYTES					
BAXTER HLTHCARE	5.5%;102MG/100ML;522MG/100ML;431MG/100ML;224MG/100ML	N017493	001		
TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER					
BAXTER HLTHCARE	8.5%;102MG/100ML;522MG/100ML;594MG/100ML;154MG/100ML	N020173	002	Oct 27, 1995	
TRAVASOL 8.5% W/ ELECTROLYTES					
BAXTER HLTHCARE	8.5%;102MG/100ML;522MG/100ML;594MG/100ML;154MG/100ML	N017493	002		

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 7% W/ ELECTROLYTES					
ICU MEDICAL INC	7%;102MG/100ML;522MG/100ML;410MG/100ML	N017789	002		
AMINOSYN 8.5% W/ ELECTROLYTES					
ICU MEDICAL INC	8.5%;102MG/100ML;522MG/100ML;410MG/100ML	N017673	005		

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMICAR					
+ EPIC PHARMA LLC	250MG/ML **				
N015229		002			
AMINOCAPROIC ACID					
ABRAXIS PHARM	250MG/ML	A070522	001	Jun 17, 1986	
BAXTER HLTHCARE	250MG/ML	N018590	001	Oct 29, 1982	
HOSPIRA	250MG/ML	A070888	001	Jun 16, 1988	

SOLUTION; ORAL

AMINOCAPROIC ACID					
EPIC PHARMA LLC	0.25GM/ML	A074759	001	Sep 02, 1998	
VISTAPHARM	0.25GM/ML	A212814	001	Feb 26, 2020	

TABLET; ORAL

AMINOCAPROIC					
HIKMA	500MG	A075602	001	May 24, 2001	
AMINOCAPROIC ACID					
EDENBRIDGE PHARMS	500MG	A212110	001	Jun 14, 2021	

AMINOGLUTETHIMIDE

TABLET; ORAL

CYTADREN					
NOVARTIS	250MG	N018202	001		

AMINOHIPPURATE SODIUM

INJECTABLE; INJECTION

AMINOHIPPURATE SODIUM					
MERCK	20%	N005619	001		

AMINOPHYLLINE

ENEMA; RECTAL

SOMOPHYLLIN					
FISONS	300MG/5ML	N018232	001	Apr 02, 1982	

INJECTABLE; INJECTION

AMINOPHYLLIN					
+ GD SEARLE LLC	25MG/ML	A087243	001	May 24, 1982	
+ AMINOPHYLLINE	25MG/ML	A087621	001	May 24, 1982	
AMINOPHYLLINE					
ABRAXIS PHARM	25MG/ML	A084568	001		
	25MG/ML	A087200	001		
	25MG/ML	A087250	001	Jan 06, 1982	
	25MG/ML	A087886	001	Aug 30, 1983	
	25MG/ML	A088407	001	Jan 25, 1984	
AM REGENT	25MG/ML	A087600	001		
ELKINS SINN	25MG/ML	A087239	001		

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

HOSPIRA	25MG/ML	A087601	001	Jul 23, 1982
INTL MEDICATION	25MG/ML	A087209	001	Feb 01, 1982
	25MG/ML	A087867	001	Nov 10, 1983
	25MG/ML	A087868	001	Nov 10, 1983
KING PHARMS	25MG/ML	A086606	001	
LUITPOLD	25MG/ML	A087240	001	
LYPHOMED	25MG/ML	A087431	001	
PHARMA SERVE NY	25MG/ML	A087387	001	Jun 03, 1983
	25MG/ML	A087392	001	Dec 15, 1983
SMITH AND NEPHEW	25MG/ML	A088429	001	May 30, 1985
	25MG/ML	A088749	001	May 30, 1985
TEVA PARENTERAL	25MG/ML	A081142	001	Sep 25, 1991
AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%				
HOSPIRA	100MG/100ML	A088147	002	May 03, 1983
	200MG/100ML	A088147	003	May 03, 1983
AMINOPHYLLINE IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
HOSPIRA	100MG/100ML	N018924	001	Dec 12, 1984
	200MG/100ML	N018924	002	Dec 12, 1984
	400MG/100ML	N018924	003	Dec 12, 1984
	500MG/100ML	N018924	004	Dec 12, 1984

SOLUTION; ORAL

AMINOPHYLLINE

MORTON GROVE	105MG/5ML	A088156	001	Dec 05, 1983
ROXANE	105MG/5ML	A088126	001	Aug 19, 1983
AMINOPHYLLINE DYE FREE				
ACTAVIS MID ATLANTIC	105MG/5ML	A087727	001	Apr 16, 1982
SOMOPHYLLIN				
FISONS	105MG/5ML	A086466	001	
SOMOPHYLLIN-DF				
FISONS	105MG/5ML	A087045	001	

SUPPOSITORY; RECTAL

TRUPHYLLINE

COSETTE	250MG	A085498	001	Mar 23, 1983
	500MG	A085498	002	Jan 03, 1983

TABLET; ORAL

AMINOPHYLLIN

GD SEARLE LLC	100MG	N002386	002	
	200MG	N002386	003	
AMINOPHYLLINE				
ANI PHARMS	100MG	A085261	004	
	200MG	A085261	002	
ASCOT	100MG	A087522	001	Feb 12, 1982
	200MG	A087523	001	Feb 12, 1982
BARR	100MG	A088297	001	Aug 19, 1983
	200MG	A088298	001	Aug 19, 1983
CHARTWELL MOLECULAR	100MG	A084588	001	
	200MG	A084588	002	
DURAMED PHARMS BARR	100MG	A088182	001	Mar 31, 1983
	200MG	A088183	001	Mar 31, 1983
HALSEY	100MG	A084674	001	
HIKMA INTL PHARMS	100MG	A084540	001	
	200MG	A085003	001	
IMPAX LABS	100MG	A084574	001	
	200MG	A084576	001	
KV PHARM	100MG	A085284	001	
	200MG	A085289	001	
PAL PAK	100MG	A084533	001	
PANRAY	100MG	A084552	001	
	200MG	A084552	002	
PUREPAC PHARM	100MG	A084699	001	
	200MG	A085333	001	
ROXANE	100MG	A087500	001	Feb 09, 1982
	200MG	A087501	001	Feb 09, 1982
VALEANT PHARM INTL	200MG	A084563	001	
VANGARD	100MG	A088314	001	Oct 03, 1983
	200MG	A088319	001	Oct 03, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOPHYLLINE

TABLET; ORAL

AMINOPHYLLINE

VINTAGE PHARMS	100MG	A085409	001
	200MG	A085410	001
WATSON LABS	100MG	A085567	001
	200MG	A085564	001

TABLET, DELAYED RELEASE; ORAL

AMINOPHYLLINE

IMPAX LABS	100MG	A084577	001
	200MG	A084575	001
TABLICAPS	100MG	A084632	002
VALE	100MG	A084531	001
	200MG	A084530	001

TABLET, EXTENDED RELEASE; ORAL

PHYLLOCONTIN

PHARM RES ASSOC	225MG	A086760	001
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AMINOSALICYLATE SODIUM

POWDER; ORAL

P.A.S. SODIUM

CENTURY PHARMS	4GM/PACKET	A080947	001
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SODIUM AMINOSALICYLATE

HEXCEL	100%	A080097	001
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TABLET; ORAL

PARASAL SODIUM

PANRAY	500MG	N006811	006
	1GM	N006811	011

SODIUM P.A.S.

LANNETT	500MG	A080138	002
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TEEBACIN

CONSOLIDATED MIDLAND	500MG	N007320	002
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AMINOSALICYLATE SODIUM; AMINOSALICYLIC ACID

TABLET; ORAL

NEOPASALATE

MEDPOINTE PHARM HLC	846MG;112MG	A080059	002
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AMINOSALICYLIC ACID

TABLET; ORAL

PARASAL

PANRAY	500MG	N006811	001
	1GM	N006811	002

AMINOSALICYLIC ACID RESIN COMPLEX

POWDER; ORAL

REZIPAS

BRISTOL MYERS SQUIBB	EQ 500MG BASE/GM	N009052	001
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AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

BEDFORD	50MG/ML	A076018	001	Oct 15, 2002
BEDFORD LABS	50MG/ML	A076299	001	Oct 24, 2002
BEN VENUE	50MG/ML	A076088	001	Oct 15, 2002
DR REDDYS	50MG/ML	A076163	001	Sep 05, 2003
EPIC PHARMA LLC	50MG/ML	A076232	001	Jul 05, 2006
EUGIA PHARMA	50MG/ML	A204550	001	Oct 25, 2017
HOSPIRA	50MG/ML	A075955	001	Oct 18, 2002
	50MG/ML	A076108	001	Oct 15, 2002
	50MG/ML	A203885	001	Nov 25, 2013
INTL MEDICATION SYS	50MG/ML	N021594	001	Feb 04, 2004
PAR STERILE PRODUCTS	50MG/ML	A076394	001	Apr 25, 2003

CORDARONE

+ WYETH PHARMS INC 50MG/ML **

N020377 001 Aug 03, 1995

NEXTERONE

+ BAXTER HLTHCARE 50MG/ML **

N022325 001 Dec 24, 2008

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

MYLAN	200MG	A075188	001	Feb 24, 1999
TEVA	200MG	A074895	001	Apr 16, 1999
UPSHER SMITH LABS	200MG	A075315	001	Dec 23, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMIODARONE HYDROCHLORIDE

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

400MG

A075315 002 Jun 30, 2000

CORDARONE

+ WYETH PHARMS

200MG **

N018972 001 Dec 24, 1985

AMITRIPTYLINE HYDROCHLORIDE

CONCENTRATE; ORAL

ENDEP

+ ROCHE

40MG/ML **

A085749 001

INJECTABLE; INJECTION

AMITRIPTYLINE HYDROCHLORIDE

WATSON LABS

10MG/ML

A085594 001

ELAVIL

ASTRAZENECA

10MG/ML

N012704 001

TABLET; ORAL

AMITID

BRISTOL MYERS SQUIBB

10MG

A086454 001

25MG

A086454 002

50MG

A086454 003

75MG

A086454 004

100MG

A086454 005

AMITRIL

WARNER CHILCOTT

10MG

A083939 001

25MG

A083937 001

50MG

A083938 002

75MG

A084957 001

100MG

A085093 001

150MG

A086295 001

AMITRIPTYLINE HYDROCHLORIDE

AIPING PHARM INC

10MG

A212654 002 Sep 29, 2021

25MG

A212654 001 Apr 07, 2020

50MG

A212654 003 Sep 29, 2021

75MG

A212654 004 Sep 29, 2021

100MG

A212654 005 Sep 29, 2021

150MG

A212654 006 Sep 29, 2021

AM THERAP

25MG

A088672 001 Nov 20, 1984

50MG

A088673 001 Nov 20, 1984

75MG

A088674 001 Nov 20, 1984

100MG

A088675 001 Nov 20, 1984

ANI PHARMS

10MG

A085031 002

25MG

A085031 001

50MG

A085031 003

75MG

A085031 004

AUROBINDO PHARMA USA

10MG

A086009 002

25MG

A086009 003

50MG

A086009 001

75MG

A086009 004

100MG

A086009 005

150MG

A086009 006

COPLEY PHARM

10MG

A088421 001 Apr 30, 1984

25MG

A088422 001 Apr 30, 1984

50MG

A088423 001 Apr 30, 1984

75MG

A088424 001 Apr 30, 1984

100MG

A088425 001 Apr 30, 1984

150MG

A088426 001 Apr 30, 1984

HALSEY

10MG

A085923 001

25MG

A085922 001

50MG

A085925 001

50MG

A087557 001 Mar 05, 1982

75MG

A085926 001 May 20, 1983

100MG

A085927 001 May 20, 1983

LEDERLE

10MG

A086744 001

10MG

A087366 001 Jan 04, 1982

25MG

A086746 001

25MG

A087367 001 May 03, 1982

50MG

A086743 001

50MG

A087181 001 Jan 04, 1982

75MG

A086745 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

	75MG	A087369 001	Jan 04, 1982
	100MG	A086747 001	
	100MG	A087368 001	May 03, 1982
	150MG	A087370 001	Jan 04, 1982
MUTUAL PHARM	10MG	A085744 001	
	25MG	A085627 001	
	50MG	A085745 001	
	75MG	A085743 001	
	100MG	A085742 002	May 11, 1982
	150MG	A089423 001	Feb 17, 1987
PAR PHARM	10MG	A088697 001	Sep 25, 1984
	25MG	A088698 001	Sep 25, 1984
	50MG	A088699 001	Sep 25, 1984
	75MG	A088700 001	Sep 25, 1984
	100MG	A088701 001	Sep 25, 1984
	150MG	A088702 001	Sep 25, 1984
PLIVA	10MG	A088883 001	Sep 26, 1984
	25MG	A088884 001	Sep 26, 1984
	50MG	A088885 001	Sep 26, 1984
	75MG	A088886 001	Sep 26, 1984
	100MG	A088887 001	Sep 26, 1984
	150MG	A088888 001	Sep 26, 1984
PUREPAC PHARM	10MG	A088075 001	Sep 16, 1983
	10MG	A088084 001	Jul 18, 1983
	25MG	A088076 001	May 20, 1983
	25MG	A088085 001	Jul 18, 1983
	50MG	A088077 001	Sep 16, 1983
	50MG	A088105 001	Jul 18, 1983
	75MG	A088078 001	Sep 16, 1983
	75MG	A088106 001	Jul 18, 1983
	100MG	A088079 001	Sep 16, 1983
	100MG	A088107 001	Jul 18, 1983
ROXANE	10MG	A086002 001	
	10MG	A086144 001	
	25MG	A085944 001	
	25MG	A086145 001	
	50MG	A085945 001	
	50MG	A086143 001	
	75MG	A086004 001	
	75MG	A086147 001	
	100MG	A086003 001	
	100MG	A086146 001	
	150MG	A086090 001	
	150MG	A086148 001	
SUN PHARM INDS INC	10MG	A040816 002	Jun 27, 2008
	25MG	A040816 001	Jun 27, 2008
	50MG	A040816 003	Jun 27, 2008
	75MG	A040816 004	Jun 27, 2008
	100MG	A040816 005	Jun 27, 2008
	150MG	A040816 006	Jun 27, 2008
SUPERPHARM	10MG	A088853 001	Nov 13, 1984
	25MG	A088854 001	Nov 13, 1984
	50MG	A088855 001	Nov 13, 1984
	75MG	A088856 001	Nov 13, 1984
	100MG	A088857 001	Nov 13, 1984
TEVA	10MG	A086610 001	
	25MG	A086859 001	
	50MG	A086857 001	
	75MG	A086860 001	
	100MG	A085836 001	
	100MG	A086854 001	
	150MG	A086853 001	
UCB INC	10MG	A085864 001	
	25MG	A085935 001	
	50MG	A085936 001	
	75MG	A086337 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

	100MG	A086336	001	
	150MG	A086335	001	
USL PHARMA	25MG	A087775	001	Feb 10, 1982
VANGARD	10MG	A087632	001	Feb 01, 1982
	50MG	A087616	001	Feb 08, 1982
	75MG	A087617	001	Feb 05, 1982
	100MG	A087639	001	Feb 08, 1982
VINTAGE PHARMS	10MG	A040218	001	Sep 11, 1997
	25MG	A040218	002	Sep 11, 1997
	50MG	A040218	003	Sep 11, 1997
	75MG	A040218	004	Sep 11, 1997
	100MG	A040218	005	Sep 11, 1997
	150MG	A040218	006	Sep 11, 1997
WATSON LABS	10MG	A085816	001	
	10MG	A088620	001	Mar 02, 1984
	25MG	A085817	001	
	25MG	A088621	001	Mar 02, 1984
	50MG	A085815	001	
	50MG	A088622	001	Mar 02, 1984
	75MG	A085819	001	
	75MG	A088633	001	Mar 02, 1984
	100MG	A085820	001	
	100MG	A088634	001	Mar 02, 1984
	150MG	A085821	001	
	150MG	A088635	001	Mar 02, 1984
WEST WARD	10MG	A087647	001	Mar 05, 1982
	25MG	A087278	001	
ELAVIL				
+ ASTRAZENECA	10MG **	N012703	001	
+	25MG **	N012703	003	
+	50MG **	N012703	004	
+	75MG **	N012703	005	
+	100MG **	N012703	006	
+	150MG **	N012703	007	
ENDEP				
ROCHE	10MG	A083639	001	
	25MG	A083639	002	
	50MG	A083639	003	
	75MG	A083639	004	
	100MG	A083639	005	
	150MG	A085303	001	

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET;ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

ANDA REPOSITORY	EQ 12.5MG BASE;5MG	A070765	001	Dec 10, 1986
	EQ 25MG BASE;10MG	A070766	001	Dec 10, 1986
CHARTWELL RX	EQ 12.5MG BASE;5MG	A072277	001	May 09, 1988
	EQ 25MG BASE;10MG	A072278	001	May 09, 1988
HERITAGE PHARMA	EQ 12.5MG BASE;5MG	A072052	001	Dec 16, 1988
	EQ 25MG BASE;10MG	A072053	001	Dec 16, 1988
USL PHARMA	EQ 12.5MG BASE;5MG	A070477	001	Jan 12, 1988
	EQ 25MG BASE;10MG	A070478	001	Jan 12, 1988
LIMBITROL				
+ CHARTWELL RX	EQ 12.5MG BASE;5MG **	N016949	001	
LIMBITROL DS				
+ CHARTWELL RX	EQ 25MG BASE;10MG **	N016949	002	

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET;ORAL

ETRAFON 2-10

SCHERING	10MG;2MG **	N014713	007	
ETRAFON 2-25				
SCHERING	25MG;2MG **	N014713	004	
ETRAFON-A				
SCHERING	10MG;4MG **	N014713	002	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

ETRAFON-FORTE

SCHERING 25MG; 4MG ** N014713 006

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

CHARTWELL RX	10MG; 2MG	A073007	001	Oct 17, 1991
	10MG; 4MG	A073009	001	Oct 17, 1991
	25MG; 2MG	A073008	001	Oct 17, 1991
	25MG; 4MG	A073010	001	Oct 17, 1991
FOSUN PHARMA	10MG; 2MG	A071062	001	Nov 27, 1987
	10MG; 4MG	A071862	001	Dec 21, 1987
	25MG; 2MG	A071063	001	Nov 27, 1987
	25MG; 4MG	A071064	001	Nov 27, 1987
	50MG; 4MG	A071863	001	Dec 21, 1987
IVAX SUB TEVA PHARMS	10MG; 2MG	A070935	001	Sep 11, 1986
	10MG; 4MG	A070937	001	Sep 11, 1986
	25MG; 2MG	A070936	001	Sep 11, 1986
	25MG; 4MG	A070938	001	Sep 11, 1986
	50MG; 4MG	A070939	001	Sep 12, 1986
PAR PHARM	10MG; 2MG	A070565	001	Sep 11, 1986
	10MG; 4MG	A070620	001	Sep 11, 1986
	25MG; 2MG	A070621	001	Sep 11, 1986
	25MG; 4MG	A070595	001	Sep 11, 1986
	50MG; 4MG	A070574	001	Sep 11, 1986
SUN PHARM INDUSTRIES	10MG; 2MG	A071077	001	Nov 12, 1986
	10MG; 4MG	A071078	001	Nov 12, 1986
	25MG; 2MG	A070297	001	Nov 12, 1986
	25MG; 4MG	A071079	001	Nov 12, 1986
WATSON LABS	10MG; 2MG	A070373	001	Aug 25, 1986
	10MG; 2MG	A072539	001	Feb 15, 1989
	10MG; 4MG	A070375	001	Aug 25, 1986
	10MG; 4MG	A072540	001	Feb 15, 1989
	25MG; 2MG	A070374	001	Aug 25, 1986
	25MG; 2MG	A072541	001	Feb 15, 1989
	25MG; 4MG	A070376	001	Aug 25, 1986
	25MG; 4MG	A072134	001	Feb 15, 1989
	50MG; 4MG	A070377	001	Nov 04, 1986
	50MG; 4MG	A071558	001	Mar 02, 1987
	50MG; 4MG	A072135	001	Feb 15, 1989

TRIAVIL 2-10

NEW RIVER 10MG; 2MG ** N014715 004

TRIAVIL 2-25

NEW RIVER 25MG; 2MG ** N014715 002

TRIAVIL 4-10

NEW RIVER 10MG; 4MG ** N014715 003

TRIAVIL 4-25

NEW RIVER 25MG; 4MG ** N014715 005

TRIAVIL 4-50

NEW RIVER 50MG; 4MG ** N014715 006

AMLEXANOX

PASTE; DENTAL

APHTHASOL

ULURU 5% N020511 001 Dec 17, 1996

PATCH; TOPICAL

AMLEXANOX

ULURU 2MG N021727 001 Sep 29, 2004

AMLODIPINE BENZOATE

SUSPENSION; ORAL

AMLODIPINE BENZOATE

AMNEAL EQ 1MG BASE/ML A215035 001 Jun 13, 2023

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

AMNEAL PHARMS NY EQ 2.5MG BASE A078477 001 Jan 16, 2008

EQ 5MG BASE A078477 002 Jan 16, 2008

EQ 10MG BASE A078477 003 Jan 16, 2008

CHARTWELL RX EQ 2.5MG BASE A076859 001 Sep 10, 2007

EQ 5MG BASE A076859 002 Sep 10, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE

TABLET;ORAL

AMLODIPINE BESYLATE

	EQ 10MG BASE	A076859 003	Sep 10, 2007
GEDEON RICHTER USA	EQ 2.5MG BASE	A077333 001	Jul 17, 2007
	EQ 5MG BASE	A077333 002	Jul 17, 2007
	EQ 10MG BASE	A077333 003	Jul 17, 2007
GENPHARM	EQ 2.5MG BASE	A077362 001	Jul 09, 2007
	EQ 5MG BASE	A077362 002	Jul 09, 2007
	EQ 10MG BASE	A077362 003	Jul 09, 2007
HIKMA	EQ 2.5MG BASE	A077262 001	Jul 09, 2007
	EQ 5MG BASE	A077262 002	Jul 09, 2007
	EQ 10MG BASE	A077262 003	Jul 09, 2007
HIKMA PHARMS	EQ 2.5MG BASE	A077771 001	Apr 12, 2011
	EQ 5MG BASE	A077771 002	Apr 12, 2011
	EQ 10MG BASE	A077771 003	Apr 12, 2011
MACLEODS PHARMS LTD	EQ 5MG BASE	A201380 001	Apr 13, 2012
	EQ 10MG BASE	A201380 002	Apr 13, 2012
MYLAN	EQ 2.5MG BASE	A076418 001	Oct 03, 2005
	EQ 2.5MG BASE	A078224 001	Feb 27, 2008
	EQ 5MG BASE	A076418 002	Oct 03, 2005
	EQ 5MG BASE	A078224 002	Feb 27, 2008
	EQ 10MG BASE	A076418 003	Oct 03, 2005
	EQ 10MG BASE	A078224 003	Feb 27, 2008
PURACAP PHARM	EQ 2.5MG BASE	A078131 001	Sep 04, 2007
	EQ 5MG BASE	A078131 002	Sep 04, 2007
	EQ 10MG BASE	A078131 003	Sep 04, 2007
SOVEREIGN PHARMS	EQ 2.5MG BASE	A204900 001	Jul 23, 2015
	EQ 5MG BASE	A204900 002	Jul 23, 2015
	EQ 10MG BASE	A204900 003	Jul 23, 2015
SUN PHARM INDS INC	EQ 2.5MG BASE	A078231 001	Nov 30, 2007
	EQ 5MG BASE	A078231 002	Nov 30, 2007
	EQ 10MG BASE	A078231 003	Nov 30, 2007
SUN PHARM INDUSTRIES	EQ 2.5MG BASE	A078081 001	Jan 31, 2008
	EQ 5MG BASE	A078081 002	Jan 31, 2008
	EQ 10MG BASE	A078081 003	Jan 31, 2008
SUNSHINE	EQ 2.5MG BASE	A206524 001	May 04, 2018
	EQ 5MG BASE	A206524 002	May 04, 2018
	EQ 10MG BASE	A206524 003	May 04, 2018
SYNTHON PHARMS	EQ 2.5MG BASE	A077080 001	Jun 27, 2007
	EQ 5MG BASE	A077080 002	Jun 27, 2007
	EQ 10MG BASE	A077080 003	Jun 27, 2007
TORRENT PHARMS	EQ 2.5MG BASE	A078573 001	Sep 22, 2008
	EQ 5MG BASE	A078573 002	Sep 22, 2008
	EQ 10MG BASE	A078573 003	Sep 22, 2008
UPSHER SMITH LABS	EQ 2.5MG BASE	A077759 001	Jul 09, 2007
	EQ 5MG BASE	A077759 002	Jul 09, 2007
	EQ 10MG BASE	A077759 003	Jul 09, 2007
WATSON LABS	EQ 2.5MG BASE	A077671 001	Jul 19, 2007
	EQ 5MG BASE	A077671 002	Jul 19, 2007
	EQ 10MG BASE	A077671 003	Jul 19, 2007
WOCKHARDT	EQ 2.5MG BASE	A078500 001	Sep 06, 2007
	EQ 5MG BASE	A078500 002	Sep 06, 2007
	EQ 10MG BASE	A078500 003	Sep 06, 2007

TABLET, ORALLY DISINTEGRATING;ORAL

AMLODIPINE BESYLATE

+	SYNTHON PHARMS	EQ 2.5MG BASE	N022026 001	Sep 27, 2007
+		EQ 5MG BASE	N022026 002	Sep 27, 2007
+		EQ 10MG BASE	N022026 003	Sep 27, 2007

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET;ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

MYLAN	EQ 2.5MG BASE;EQ 10MG BASE	A200465 001	Nov 29, 2013
	EQ 2.5MG BASE;EQ 20MG BASE	A200465 002	Nov 29, 2013
	EQ 2.5MG BASE;EQ 40MG BASE	A200465 003	Nov 29, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

CIPLA	EQ 2.5MG BASE;10MG	A077215 001	Dec 07, 2018
	EQ 5MG BASE;10MG	A077215 002	Dec 07, 2018
	EQ 5MG BASE;20MG	A077215 003	Dec 07, 2018
	EQ 10MG BASE;20MG	A077215 004	Dec 07, 2018
MYLAN	EQ 2.5MG BASE;10MG	A077375 001	May 21, 2010
	EQ 5MG BASE;10MG	A077375 002	May 21, 2010
	EQ 5MG BASE;20MG	A077375 003	May 21, 2010
	EQ 5MG BASE;40MG	A079047 001	Jul 05, 2011
	EQ 10MG BASE;20MG	A077375 004	May 21, 2010
	EQ 10MG BASE;40MG	A079047 002	Jul 05, 2011
STRIDES PHARMA	EQ 2.5MG BASE;10MG	A078381 001	Jul 29, 2010
	EQ 5MG BASE;10MG	A078381 002	Jul 29, 2010
	EQ 5MG BASE;20MG	A078381 003	Jul 29, 2010
	EQ 5MG BASE;40MG	A078381 005	Jul 29, 2010
	EQ 10MG BASE;20MG	A078381 004	Jul 29, 2010
	EQ 10MG BASE;40MG	A078381 006	Jul 29, 2010
TEVA PHARMS	EQ 2.5MG BASE;10MG	A077179 001	May 18, 2007
	EQ 5MG BASE;10MG	A077179 002	May 18, 2007
	EQ 5MG BASE;20MG	A077179 003	May 18, 2007
	EQ 5MG BASE;40MG	A077179 005	Jul 05, 2011
	EQ 10MG BASE;20MG	A077179 004	May 18, 2007
	EQ 10MG BASE;40MG	A077179 006	Jul 05, 2011

AMLODIPINE BESYLATE; CELECOXIB

TABLET;ORAL

CONSENSI

+	PURPLE BIOTECH	EQ 2.5MG BASE;200MG	N210045 001	May 31, 2018
+		EQ 5MG BASE;200MG	N210045 002	May 31, 2018
+		EQ 10MG BASE;200MG	N210045 003	May 31, 2018

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET;ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

TEVA PHARMS	EQ 5MG BASE;12.5MG;160MG	A200435 001	Sep 25, 2012
	EQ 5MG BASE;25MG;160MG	A200435 002	Sep 25, 2012
	EQ 10MG BASE;12.5MG;160MG	A200435 005	Sep 25, 2012
	EQ 10MG BASE;25MG;160MG	A200435 003	Sep 25, 2012
	EQ 10MG BASE;25MG;320MG	A200435 004	Sep 25, 2012
TORRENT	EQ 5MG BASE;12.5MG;160MG	A201593 001	Jun 03, 2015
	EQ 5MG BASE;25MG;160MG	A201593 002	Jun 03, 2015
	EQ 10MG BASE;12.5MG;160MG	A201593 003	Jun 03, 2015
	EQ 10MG BASE;25MG;160MG	A201593 004	Jun 03, 2015
	EQ 10MG BASE;25MG;320MG	A201593 005	Jun 03, 2015

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

ACCORD HLTHCARE INC	EQ 5MG BASE;20MG	A209600 001	Aug 30, 2018
	EQ 5MG BASE;40MG	A209600 003	Aug 30, 2018
	EQ 10MG BASE;20MG	A209600 002	Aug 30, 2018
JUBILANT GENERICS	EQ 10MG BASE;40MG	A209600 004	Aug 30, 2018
	EQ 5MG BASE;20MG	A207450 001	May 15, 2017
	EQ 5MG BASE;40MG	A207450 002	May 15, 2017
	EQ 10MG BASE;20MG	A207450 003	May 15, 2017
	EQ 10MG BASE;40MG	A207450 004	May 15, 2017
SCIEGEN PHARMS INC	EQ 5MG BASE;20MG	A209010 001	Dec 03, 2018
	EQ 5MG BASE;40MG	A209010 002	Dec 03, 2018
	EQ 10MG BASE;20MG	A209010 003	Dec 03, 2018
	EQ 10MG BASE;40MG	A209010 004	Dec 03, 2018
TEVA PHARMS USA	EQ 5MG BASE;20MG	A091154 001	Oct 26, 2016
	EQ 5MG BASE;40MG	A091154 002	Oct 26, 2016
	EQ 10MG BASE;20MG	A091154 003	Oct 26, 2016
	EQ 10MG BASE;40MG	A091154 004	Oct 26, 2016
TORRENT	EQ 5MG BASE;20MG	A202933 001	Nov 25, 2016
	EQ 5MG BASE;40MG	A202933 002	Nov 25, 2016
	EQ 10MG BASE;20MG	A202933 003	Nov 25, 2016
	EQ 10MG BASE;40MG	A202933 004	Nov 25, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMLODIPINE BESYLATE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND AMLODIPINE

ALEMBIC	EQ 5MG BASE;40MG	A205234 001	Nov 17, 2016
	EQ 5MG BASE;80MG	A205234 003	Nov 17, 2016
	EQ 10MG BASE;40MG	A205234 002	Nov 17, 2016
	EQ 10MG BASE;80MG	A205234 004	Nov 17, 2016
TORRENT	EQ 5MG BASE;40MG	A202517 001	Jan 08, 2014
	EQ 5MG BASE;80MG	A202517 003	Jan 08, 2014
	EQ 10MG BASE;40MG	A202517 002	Jan 08, 2014
	EQ 10MG BASE;80MG	A202517 004	Jan 08, 2014
TWYNSTA			
+	BOEHRINGER INGELHEIM EQ 5MG BASE;40MG **	N022401 001	Oct 16, 2009
+	EQ 5MG BASE;80MG **	N022401 003	Oct 16, 2009
+	EQ 10MG BASE;40MG **	N022401 002	Oct 16, 2009
+	EQ 10MG BASE;80MG **	N022401 004	Oct 16, 2009

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

TEVA PHARMS USA	EQ 5MG BASE;160MG	A091235 001	Mar 30, 2015
	EQ 5MG BASE;320MG	A091235 003	Mar 30, 2015
	EQ 10MG BASE;160MG	A091235 002	Mar 30, 2015
	EQ 10MG BASE;320MG	A091235 004	Mar 30, 2015
TORRENT	EQ 5MG BASE;160MG	A202377 001	Mar 30, 2015
	EQ 5MG BASE;320MG	A202377 002	Mar 30, 2015
	EQ 10MG BASE;160MG	A202377 003	Mar 30, 2015
	EQ 10MG BASE;320MG	A202377 004	Mar 30, 2015

AMLODIPINE MALEATE

TABLET; ORAL

AMVAZ

DR REDDYS LABS INC	2.5MG	N021435 001	Oct 31, 2003
	5MG	N021435 002	Oct 31, 2003
	10MG	N021435 003	Oct 31, 2003

AMMONIA N-13

INJECTABLE; INTRAVENOUS

AMMONIA N 13

CENTRAL RADIOPHARM	3.75-260mCi/ML	A204539 001	Jun 23, 2015
ESSENTIAL ISOTOPES	3.75mCi-260mCi/ML	A205687 001	Dec 17, 2015
SHERTECH LABS LLC	3.75mCi-260mCi/ML	A204366 001	Sep 19, 2014
SOFIE	18.8mCi-188mCi/5ML (3.75-37.5mCi/ML)	A204667 001	Apr 22, 2015
UNIV TX MD ANDERSON	30mCi-300mCi/8ML (3.75-37.5mCi/ML)	A203933 001	Jun 27, 2014
WISCONSIN	3.75mCi-260mCi/ML	A204356 001	Dec 18, 2014

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE

ABBOTT	5MEQ/ML	A083130 001	
GD SEARLE LLC	3MEQ/ML	A086205 001	
AMMONIUM CHLORIDE 0.9% IN NORMAL SALINE			
MCGAW	900MG/100ML	N006580 001	
AMMONIUM CHLORIDE 2.14%			
B BRAUN	40MEQ/100ML	A085734 001	

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

WATSON LABS INC	EQ 12% BASE	A076829 001	Feb 07, 2006
LAC-HYDRIN			
+	SUN PHARM INDS INC EQ 12% BASE **	N020508 001	Aug 29, 1996
LOTION; TOPICAL			
AMMONIUM LACTATE			
WATSON LABS INC	EQ 12% BASE	A075575 001	Jun 11, 2002
LAC-HYDRIN			
+	SUN PHARM INDS INC EQ 12% BASE **	N019155 001	Apr 24, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMODIAQUINE HYDROCHLORIDE

TABLET; ORAL

CAMOQUIN HYDROCHLORIDE

PARKE DAVIS

EQ 200MG BASE

N006441 001

AMOXAPINE

TABLET; ORAL

AMOXAPINE

CHARTWELL RX

25MG

A072943 001 Jun 28, 1991

50MG

A072944 001 Jun 28, 1991

100MG

A072878 001 Jun 28, 1991

150MG

A072879 001 Jun 28, 1991

WATSON PHARMS TEVA

25MG

A072418 001 Aug 01, 1989

50MG

A072419 001 Aug 01, 1989

100MG

A072420 001 May 11, 1989

150MG

A072421 001 May 11, 1989

ASENDIN

LEDERLE

25MG

N018021 001

50MG

N018021 002

100MG

N018021 003

150MG

N018021 004

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

LABS ATRAL

250MG

A062528 001 Aug 07, 1985

500MG

A062528 002 Aug 07, 1985

MYLAN

250MG

A062067 001

500MG

A062067 002

STRIDES PHARMA

250MG

A062884 001 Feb 25, 1988

500MG

A062881 001 Feb 25, 1988

SUN PHARM INDS LTD

250MG

A065016 001 Apr 08, 1999

500MG

A065016 002 Apr 08, 1999

TEVA

250MG

A062853 001 Dec 22, 1987

250MG

A063030 001 Feb 28, 1989

500MG

A062854 001 Dec 22, 1987

500MG

A063031 001 Feb 28, 1989

AMOXIL

+ GLAXOSMITHKLINE

250MG **

N050459 001

+

500MG **

N050459 002

TRIMOX

APOTHECON

250MG

A061885 001

250MG

A062098 001

250MG

A062152 001

250MG

A063099 001 Mar 20, 1992

500MG

A061885 002

500MG

A062098 002

500MG

A062152 002

500MG

A063099 002 Mar 20, 1992

UTIMOX

PARKE DAVIS

250MG

A062107 001

500MG

A062107 002

WYMOX

WYETH AYERST

250MG

A062120 001

500MG

A062120 002

FOR SUSPENSION; ORAL

AMOXICILLIN

CHARTWELL RX

125MG/5ML

A062059 001

250MG/5ML

A062059 002

MYLAN

125MG/5ML

A062090 001

250MG/5ML

A062090 002

SUN PHARM INDS LTD

200MG/5ML

A065113 001 Nov 29, 2002

400MG/5ML

A065113 002 Nov 29, 2002

TEVA

125MG/5ML

A061931 001

125MG/5ML

A062946 001 Nov 01, 1988

250MG/5ML

A063001 001 Jan 06, 1989

AMOXIL

+ GLAXOSMITHKLINE

50MG/ML **

N050460 005

+

125MG/5ML **

N050460 001

+

250MG/5ML **

N050460 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN

FOR SUSPENSION;ORAL

AMOXIL					
+	US ANTIBIOTICS	200MG/5ML **	N050760 001	Apr 15, 1999	
+		400MG/5ML **	N050760 002	Apr 15, 1999	
LAROTID					
+	GLAXOSMITHKLINE	50MG/ML **	N050460 006		
POLYMOX					
	APOTHECON	125MG/5ML	A061851 001		
		125MG/5ML	A062323 001		
		250MG/5ML	A061851 002		
		250MG/5ML	A062323 002		
TRIMOX					
	APOTHECON	50MG/ML	A061886 001		
		125MG/5ML	A061886 002		
		125MG/5ML	A062099 001		
		125MG/5ML	A062154 001		
		125MG/5ML	A062885 001	Mar 08, 1988	
		250MG/5ML	A061886 003		
		250MG/5ML	A062099 002		
		250MG/5ML	A062154 002		
		250MG/5ML	A062885 002	Mar 08, 1988	
UTIMOX					
	PARKE DAVIS	125MG/5ML	A062127 001		
		250MG/5ML	A062127 002		
WYMOX					
	WYETH AYERST	125MG/5ML	A062131 001		
		250MG/5ML	A062131 002		

TABLET;ORAL

AMOXICILLIN					
	CHARTWELL RX	875MG	A065344 001	Jan 15, 2009	
	SUN PHARM INDS LTD	500MG	A065059 001	Nov 24, 2000	
		875MG	A065059 002	Nov 24, 2000	
AMOXIL					
+	US ANTIBIOTICS	500MG **	N050754 002	Jul 10, 1998	
+		875MG **	N050754 001	Jul 10, 1998	
TABLET, CHEWABLE;ORAL					
AMOXICILLIN					
	APOTHECON	125MG	A064131 001	May 06, 1996	
		250MG	A064131 002	May 06, 1996	
	CHARTWELL RX	125MG	A064139 001	Jan 29, 1996	
		250MG	A064139 002	Jan 29, 1996	
	HIKMA	125MG	A205274 001	Jun 25, 2020	
		250MG	A205274 002	Jun 25, 2020	
	SUN PHARM INDS LTD	125MG	A065021 001	Dec 23, 1999	
		200MG	A065060 001	Nov 29, 2000	
		250MG	A065021 002	Dec 23, 1999	
		400MG	A065060 002	Nov 29, 2000	
	TEVA	125MG	A064031 001	Dec 19, 1996	
		250MG	A064031 002	Dec 19, 1996	
AMOXIL					
+	US ANTIBIOTICS	125MG **	N050542 002		
		200MG	N050761 001	Apr 15, 1999	
+		250MG **	N050542 001		
		400MG	N050761 002	Apr 15, 1999	
TABLET, EXTENDED RELEASE;ORAL					
MOXATAG					
+	PRAGMA	775MG	N050813 001	Jan 23, 2008	
TABLET, FOR SUSPENSION;ORAL					
AMOXICILLIN					
	AUROBINDO PHARMA LTD	200MG	A065324 001	Jan 17, 2007	
		400MG	A065324 002	Jan 17, 2007	
DISPERMOX					
	RANBAXY LABS LTD	200MG	A065080 002	Aug 11, 2003	
		400MG	A065080 001	Aug 11, 2003	
		600MG	A065159 001	Dec 04, 2003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED REL PELLETS;ORAL
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED)
 ANI PHARMS 500MG;500MG;30MG A200218 001 Aug 30, 2013
 PREVPAC (COPACKAGED)
 + TAKEDA PHARMS USA 500MG;500MG;30MG ** N050757 001 Dec 02, 1997

AMOXICILLIN; CLARITHROMYCIN; OMEPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED RELEASE;ORAL
 OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN
 + CUMBERLAND 500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,20M N050824 001 Feb 08, 2011
 G **

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM
 SUN PHARM INDS LTD 200MG/5ML;EQ 28.5MG BASE/5ML A065132 001 Mar 19, 2003
 400MG/5ML;EQ 57MG BASE/5ML A065132 002 Mar 19, 2003
 600MG/5ML;EQ 42.9MG BASE/5ML A065207 002 Jan 30, 2007
 AUGMENTIN '200'
 + US ANTIBIOTICS 200MG/5ML;EQ 28.5MG BASE/5ML N050725 001 May 31, 1996
 AUGMENTIN '400'
 + US ANTIBIOTICS 400MG/5ML;EQ 57MG BASE/5ML N050725 002 May 31, 1996

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM
 APOTEX INC 250MG;EQ 125MG BASE A065333 001 Feb 24, 2009
 500MG;EQ 125MG BASE A065333 002 Feb 24, 2009
 875MG;EQ 125MG BASE A065317 003 Oct 20, 2008
 SUN PHARM INDS LTD 500MG;EQ 125MG BASE A065109 001 Nov 04, 2002
 875MG;EQ 125MG BASE A065102 001 Sep 17, 2002
 AUGMENTIN '250'
 + US ANTIBIOTICS 250MG;EQ 125MG BASE ** N050564 001 Aug 06, 1984
 AUGMENTIN '500'
 + US ANTIBIOTICS 500MG;EQ 125MG BASE ** N050564 002 Aug 06, 1984

TABLET, CHEWABLE;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM
 SANDOZ 200MG;EQ 28.5MG BASE A065065 001 Apr 18, 2002
 400MG;EQ 57MG BASE A065065 002 Apr 18, 2002
 SUN PHARM INDS LTD 200MG;EQ 28.5MG BASE A065161 001 Dec 03, 2003
 400MG;EQ 57MG BASE A065161 002 Dec 03, 2003
 AUGMENTIN '125'
 + US ANTIBIOTICS 125MG;EQ 31.25MG BASE ** N050597 001 Jul 22, 1985
 AUGMENTIN '200'
 + US ANTIBIOTICS 200MG;EQ 28.5MG BASE N050726 001 May 31, 1996
 AUGMENTIN '250'
 + US ANTIBIOTICS 250MG;EQ 62.5MG BASE ** N050597 002 Jul 22, 1985
 AUGMENTIN '400'
 + US ANTIBIOTICS 400MG;EQ 57MG BASE N050726 002 May 31, 1996

TABLET, EXTENDED RELEASE;ORAL

AUGMENTIN XR
 + US ANTIBIOTICS 1GM;EQ 62.5MG BASE N050785 001 Sep 25, 2002

AMPHETAMINE

SUSPENSION, EXTENDED RELEASE;ORAL

ADZENYS ER
 + NEOS THERAPS INC EQ 1.25MG BASE/ML N204325 001 Sep 15, 2017

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL

AMPHETAMINE
 ACTAVIS LABS FL INC EQ 3.1MG BASE A209253 001 Jun 22, 2023
 EQ 6.3MG BASE A209253 002 Jun 22, 2023
 EQ 9.4MG BASE A209253 003 Jun 22, 2023
 EQ 12.5MG BASE A209253 004 Jun 22, 2023
 EQ 15.7MG BASE A209253 005 Jun 22, 2023
 EQ 18.8MG BASE A209253 006 Jun 22, 2023

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHETAMINE ADIPATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE ADIPATE; DEXTROAMPHETAMINE SULFATE

CAPSULE; ORAL

DELCOBESE

TEVA

1.25MG; 1.25MG; 1.25MG; 1.25MG **	A083564	001
2.5MG; 2.5MG; 2.5MG; 2.5MG **	A083564	002
3.75MG; 3.75MG; 3.75MG; 3.75MG **	A083564	003
5MG; 5MG; 5MG; 5MG **	A083564	004

TABLET; ORAL

DELCOBESE

TEVA

1.25MG; 1.25MG; 1.25MG; 1.25MG	A083563	004
2.5MG; 2.5MG; 2.5MG; 2.5MG	A083563	003
3.75MG; 3.75MG; 3.75MG; 3.75MG	A083563	002
5MG; 5MG; 5MG; 5MG	A083563	001

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AUROLIFE PHARMA LLC

1.25MG; 1.25MG; 1.25MG; 1.25MG	A211876	001	Jun 24, 2020
2.5MG; 2.5MG; 2.5MG; 2.5MG	A211876	002	Jun 24, 2020
3.75MG; 3.75MG; 3.75MG; 3.75MG	A211876	003	Jun 24, 2020
5MG; 5MG; 5MG; 5MG	A211876	004	Jun 24, 2020
6.25MG; 6.25MG; 6.25MG; 6.25MG	A211876	005	Jun 24, 2020
7.5MG; 7.5MG; 7.5MG; 7.5MG	A211876	006	Jun 24, 2020

NESHER PHARMS

1.25MG; 1.25MG; 1.25MG; 1.25MG	A210080	001	Jul 17, 2019
2.5MG; 2.5MG; 2.5MG; 2.5MG	A210080	002	Jul 17, 2019
3.75MG; 3.75MG; 3.75MG; 3.75MG	A210080	003	Jul 17, 2019
5MG; 5MG; 5MG; 5MG	A210080	004	Jul 17, 2019
6.25MG; 6.25MG; 6.25MG; 6.25MG	A210080	005	Jul 17, 2019
7.5MG; 7.5MG; 7.5MG; 7.5MG	A210080	006	Jul 17, 2019

PAR PHARM INC

1.25MG; 1.25MG; 1.25MG; 1.25MG	A206159	001	May 31, 2019
2.5MG; 2.5MG; 2.5MG; 2.5MG	A206159	002	May 31, 2019
3.75MG; 3.75MG; 3.75MG; 3.75MG	A206159	003	May 31, 2019
5MG; 5MG; 5MG; 5MG	A206159	004	May 31, 2019
6.25MG; 6.25MG; 6.25MG; 6.25MG	A206159	005	May 31, 2019
7.5MG; 7.5MG; 7.5MG; 7.5MG	A206159	006	May 31, 2019

TEVA

1.25MG; 1.25MG; 1.25MG; 1.25MG	A077488	001	Apr 29, 2013
2.5MG; 2.5MG; 2.5MG; 2.5MG	A077488	002	Apr 29, 2013
3.75MG; 3.75MG; 3.75MG; 3.75MG	A077488	003	Apr 29, 2013
5MG; 5MG; 5MG; 5MG	A077488	004	Apr 29, 2013
6.25MG; 6.25MG; 6.25MG; 6.25MG	A077488	005	Apr 29, 2013
7.5MG; 7.5MG; 7.5MG; 7.5MG	A077488	006	Apr 29, 2013

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

BARR LABS INC

1.25MG; 1.25MG; 1.25MG; 1.25MG	A076536	001	Feb 12, 2013
2.5MG; 2.5MG; 2.5MG; 2.5MG	A076536	002	Feb 12, 2013
3.75MG; 3.75MG; 3.75MG; 3.75MG	A076536	003	Feb 12, 2013
5MG; 5MG; 5MG; 5MG	A076536	004	Feb 12, 2013
6.25MG; 6.25MG; 6.25MG; 6.25MG	A076536	005	Feb 12, 2013
7.5MG; 7.5MG; 7.5MG; 7.5MG	A076536	006	Feb 12, 2013

TABLET; ORAL

ADDERALL 10

+ TEVA WOMENS

2.5MG; 2.5MG; 2.5MG; 2.5MG **	N011522	007	Feb 13, 1996
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ADDERALL 12.5

+ TEVA WOMENS

3.125MG; 3.125MG; 3.125MG; 3.125MG **	N011522	012	Aug 31, 2000
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ADDERALL 15

+ TEVA WOMENS

3.75MG; 3.75MG; 3.75MG; 3.75MG **	N011522	013	Aug 31, 2000
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ADDERALL 20

+ TEVA WOMENS

5MG; 5MG; 5MG; 5MG **	N011522	008	Feb 13, 1996
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ADDERALL 30

+ TEVA WOMENS

7.5MG; 7.5MG; 7.5MG; 7.5MG **	N011522	010	May 12, 1997
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ADDERALL 5

+ TEVA WOMENS

1.25MG; 1.25MG; 1.25MG; 1.25MG **	N011522	009	May 12, 1997
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ADDERALL 7.5

+ TEVA WOMENS

1.875MG; 1.875MG; 1.875MG; 1.875MG **	N011522	011	Aug 31, 2000
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DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

ACCORD HLTHCARE

1.25MG; 1.25MG; 1.25MG; 1.25MG	A214347	001	Nov 22, 2021
1.875MG; 1.875MG; 1.875MG; 1.875MG	A214347	002	Nov 22, 2021
2.5MG; 2.5MG; 2.5MG; 2.5MG	A214347	003	Nov 22, 2021
3.125MG; 3.125MG; 3.125MG; 3.125MG	A214347	004	Nov 22, 2021
3.75MG; 3.75MG; 3.75MG; 3.75MG	A214347	005	Nov 22, 2021

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET;ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

	5MG;5MG;5MG;5MG	A214347	006	Nov 22, 2021
	7.5MG;7.5MG;7.5MG;7.5MG	A214347	007	Nov 22, 2021
ACTAVIS ELIZABETH	1.25MG;1.25MG;1.25MG;1.25MG	A040456	001	May 06, 2003
	2.5MG;2.5MG;2.5MG;2.5MG	A040456	002	May 06, 2003
	5MG;5MG;5MG;5MG	A040456	003	May 06, 2003
	7.5MG;7.5MG;7.5MG;7.5MG	A040456	004	May 06, 2003
CEDIPROF INC	1.25MG;1.25MG;1.25MG;1.25MG	A210754	001	Jul 05, 2022
	2.5MG;2.5MG;2.5MG;2.5MG	A210754	002	Jul 05, 2022
	3.75MG;3.75MG;3.75MG;3.75MG	A210754	003	Jul 05, 2022
	5MG;5MG;5MG;5MG	A210754	004	Jul 05, 2022
	7.5MG;7.5MG;7.5MG;7.5MG	A210754	005	Jul 05, 2022
MYLAN	1.25MG;1.25MG;1.25MG;1.25MG	A206721	001	Nov 10, 2015
	1.875MG;1.875MG;1.875MG;1.875MG	A206721	002	Nov 10, 2015
	2.5MG;2.5MG;2.5MG;2.5MG	A206721	003	Nov 10, 2015
	3.125MG;3.125MG;3.125MG;3.125MG	A206721	004	Nov 10, 2015
	3.75MG;3.75MG;3.75MG;3.75MG	A206721	005	Nov 10, 2015
	5MG;5MG;5MG;5MG	A206721	006	Nov 10, 2015
	7.5MG;7.5MG;7.5MG;7.5MG	A206721	007	Nov 10, 2015
TEVA PHARMS	1.25MG;1.25MG;1.25MG;1.25MG	A040472	001	Sep 30, 2003
	2.5MG;2.5MG;2.5MG;2.5MG	A040472	002	Sep 30, 2003
	5MG;5MG;5MG;5MG	A040472	003	Sep 30, 2003
	7.5MG;7.5MG;7.5MG;7.5MG	A040472	004	Sep 30, 2003

AMPHETAMINE RESIN COMPLEX; DEXTROAMPHETAMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE;ORAL

BIPHETAMINE 12.5

UCB INC EQ 6.25MG BASE;EQ 6.25MG BASE N010093 007

BIPHETAMINE 20

UCB INC EQ 10MG BASE;EQ 10MG BASE N010093 003

BIPHETAMINE 7.5

UCB INC EQ 3.75MG BASE;EQ 3.75MG BASE N010093 009

AMPHETAMINE SULFATE

TABLET;ORAL

AMPHETAMINE SULFATE

DR REDDYS LABS SA	5MG	A213898	001	Jul 14, 2020
	10MG	A213898	002	Jul 14, 2020
GLENMARK PHARMS LTD	5MG	A212186	001	Jan 27, 2021
	10MG	A212186	002	Jan 27, 2021
+ LANNETT	5MG	A083901	001	Aug 31, 1984
+	10MG	A083901	002	Aug 31, 1984
NOVAST LABS	5MG	A213763	001	Aug 24, 2020
	10MG	A213763	002	Aug 24, 2020

TABLET, ORALLY DISINTEGRATING;ORAL

EVEKEO ODT

+ AZURITY 2.5MG ** N209905 005 Apr 16, 2021

AMPHOTERICIN B

CREAM;TOPICAL

FUNGIZONE

APOTHECON 3% N050314 001

INJECTABLE;INJECTION

AMPHOTERICIN B

ABBOT	50MG/VIAL	A064141	001	Dec 23, 1996
ABRAXIS PHARM	50MG/VIAL	A062728	001	Apr 13, 1987
TEVA PARENTERAL	50MG/VIAL	A064062	001	Mar 31, 1995

FUNGIZONE

APOTHECON 50MG/VIAL A060517 001

INJECTABLE, LIPID COMPLEX;INJECTION

AMPHOTEC

ALKOPHARMA USA	50MG/VIAL	N050729	001	Nov 22, 1996
	100MG/VIAL	N050729	002	Nov 22, 1996

LOTION;TOPICAL

FUNGIZONE

APOTHECON 3% A060570 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHOTERICIN BOINTMENT; TOPICAL
FUNGIZONE

APOTHECON 3% N050313 001

SUSPENSION; ORAL
FUNGIZONE

BRISTOL MYERS SQUIBB 100MG/ML N050341 003

AMPICILLIN SODIUMINJECTABLE; INJECTION
AMPICILLIN SODIUM

ACS DOBFAR SPA	EQ 500MG BASE/VIAL	A090884 001	Apr 03, 2013
	EQ 1GM BASE/VIAL	A090884 002	Apr 03, 2013
	EQ 2GM BASE/VIAL	A090884 003	Apr 03, 2013
APOTHECON	EQ 125MG BASE/VIAL	A062860 001	Feb 05, 1988
	EQ 250MG BASE/VIAL	A062860 002	Feb 05, 1988
	EQ 500MG BASE/VIAL	A062860 003	Feb 05, 1988
	EQ 1GM BASE/VIAL	A062860 004	Feb 05, 1988
	EQ 2GM BASE/VIAL	A062860 005	Feb 05, 1988
CONSOLIDATED PHARM	EQ 125MG BASE/VIAL	A061936 005	
	EQ 250MG BASE/VIAL	A061936 001	
	EQ 500MG BASE/VIAL	A061936 002	
	EQ 1GM BASE/VIAL	A061936 003	
	EQ 2GM BASE/VIAL	A061936 004	
EUGIA PHARMA SPECLTS	EQ 125MG BASE/VIAL	A065499 001	Aug 17, 2010
HIKMA	EQ 125MG BASE/VIAL	A062692 001	Jun 24, 1986
	EQ 250MG BASE/VIAL	A062692 002	Jun 24, 1986
	EQ 500MG BASE/VIAL	A062692 003	Jun 24, 1986
	EQ 1GM BASE/VIAL	A062692 004	Jun 24, 1986
	EQ 2GM BASE/VIAL	A062692 005	Jun 24, 1986
	EQ 10GM BASE/VIAL	A062692 006	Jun 24, 1986
HOSPIRA	EQ 250MG BASE/VIAL	A202864 001	Sep 04, 2015
	EQ 500MG BASE/VIAL	A202864 002	Sep 04, 2015
	EQ 1GM BASE/VIAL	A202864 003	Sep 04, 2015
	EQ 2GM BASE/VIAL	A202864 004	Sep 04, 2015
	EQ 10GM BASE/VIAL	A202865 001	Sep 04, 2015
HQ SPECLT PHARMA	EQ 125MG BASE/VIAL	A062772 005	Apr 15, 1993
	EQ 500MG BASE/VIAL	A062772 008	Apr 15, 1993
	EQ 1GM BASE/VIAL	A062772 002	Apr 15, 1993
	EQ 2GM BASE/VIAL	A062772 004	Apr 15, 1993
INTL MEDICATION	EQ 1GM BASE/VIAL	A062634 002	Jan 09, 1987
	EQ 2GM BASE/VIAL	A062634 003	Jan 09, 1987
ISTITUTO BIO ITA SPA	EQ 125MG BASE/VIAL	A062797 001	Jul 12, 1993
LILLY	EQ 500MG BASE/VIAL	A062565 001	Apr 04, 1985
	EQ 1GM BASE/VIAL	A062565 002	Apr 04, 1985
	EQ 2GM BASE/VIAL	A062565 003	Jun 24, 1986
STERISCIENCE	EQ 250MG BASE/VIAL	A201025 001	Apr 09, 2014
	EQ 500MG BASE/VIAL	A201025 002	Apr 09, 2014
WATSON LABS INC	EQ 125MG BASE/VIAL	A062816 001	Oct 24, 1988
	EQ 250MG BASE/VIAL	A062816 002	Oct 24, 1988
	EQ 500MG BASE/VIAL	A062816 003	Oct 24, 1988
	EQ 1GM BASE/VIAL	A062816 004	Oct 24, 1988
	EQ 2GM BASE/VIAL	A062816 005	Oct 24, 1988
	EQ 10GM BASE/VIAL	A062994 001	Sep 15, 1988
OMNIPEN-N			
WYETH AYERST	EQ 125MG BASE/VIAL	A060626 001	
	EQ 125MG BASE/VIAL	A062718 001	Dec 16, 1986
	EQ 250MG BASE/VIAL	A060626 002	
	EQ 250MG BASE/VIAL	A062718 002	Dec 16, 1986
	EQ 500MG BASE/VIAL	A060626 003	
	EQ 500MG BASE/VIAL	A062718 003	Dec 16, 1986
	EQ 1GM BASE/VIAL	A060626 004	
	EQ 1GM BASE/VIAL	A062718 004	Dec 16, 1986
	EQ 2GM BASE/VIAL	A060626 005	
	EQ 2GM BASE/VIAL	A062718 005	Dec 16, 1986
PENBRITIN-S			
+ WYETH AYERST	EQ 125MG BASE/VIAL **	N050072 001	
+	EQ 250MG BASE/VIAL **	N050072 002	
+	EQ 500MG BASE/VIAL **	N050072 003	
+	EQ 1GM BASE/VIAL **	N050072 004	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN SODIUM

INJECTABLE; INJECTION

PENBRITIN-S

+

EQ 2GM BASE/VIAL **

N050072 005

+

EQ 4GM BASE/VIAL **

N050072 006

POLYCILLIN-N

BRISTOL

EQ 125MG BASE/VIAL **

N050309 001

EQ 250MG BASE/VIAL **

N050309 002

EQ 500MG BASE/VIAL **

N050309 003

EQ 1GM BASE/VIAL **

N050309 004

EQ 2GM BASE/VIAL **

N050309 005

TOTACILLIN-N

GLAXOSMITHKLINE

EQ 125MG BASE/VIAL

A060677 001

EQ 250MG BASE/VIAL

A060677 002

EQ 500MG BASE/VIAL

A060677 003

EQ 1GM BASE/VIAL

A060677 004

EQ 1GM BASE/VIAL

A062727 001 Dec 19, 1986

EQ 2GM BASE/VIAL

A060677 005

EQ 2GM BASE/VIAL

A062727 002 Dec 19, 1986

EQ 10GM BASE/VIAL

A060677 006

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

EUGIA PHARMA SPECLTS

EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL

A090340 001 Sep 20, 2010

EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL

A090340 002 Sep 20, 2010

HOSPIRA INC

EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL

A090375 001 Dec 21, 2011

EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL

A090653 001 Dec 21, 2011

EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL

A090375 002 Dec 21, 2011

EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL

A090653 002 Dec 21, 2011

EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL

A090646 001 Dec 21, 2011

PHARM ASSOC

EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL

A065316 001 Jun 29, 2007

EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL

A065316 002 Jun 29, 2007

UNASYN

PFIZER

EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL

A062901 002 Feb 27, 1992

EQ 500MG BASE/VIAL;EQ 250MG BASE/VIAL

N050608 003 Dec 31, 1986

EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL

A062901 001 Nov 23, 1988

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMCILL

PARKE DAVIS

EQ 250MG BASE

A062041 001

EQ 500MG BASE

A062041 002

AMPICILLIN TRIHYDRATE

BELCHER PHARMS

EQ 250MG BASE

A061602 001

EQ 500MG BASE

A061602 002

CHARTWELL RX

EQ 250MG BASE

A061502 001

EQ 500MG BASE

A061502 002

IVAX SUB TEVA PHARMS

EQ 250MG BASE

A060765 001

EQ 500MG BASE

A060765 002

LEDERLE

EQ 250MG BASE

A062208 001

EQ 500MG BASE

A062208 002

MYLAN

EQ 250MG BASE

A061755 001

EQ 500MG BASE

A061755 002

PUREPAC PHARM

EQ 250MG BASE

A061853 001

EQ 500MG BASE

A061853 002

SANDOZ

EQ 250MG BASE

A064082 001 Aug 29, 1995

STRIDES PHARMA

EQ 250MG BASE

A062883 001 Feb 25, 1988

EQ 500MG BASE

A062882 001 Feb 25, 1988

VITARINE

EQ 250MG BASE

A061387 001

EQ 500MG BASE

A061387 003

OMNIPEN (AMPICILLIN)

WYETH AYERST

250MG

A060624 001

500MG

A060624 002

PENBRITIN

WYETH AYERST

EQ 250MG BASE

A060908 001

EQ 500MG BASE

A060908 002

PFIZERPEN-A

PFIZER

EQ 250MG BASE

A062050 001

EQ 500MG BASE

A062050 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE;ORAL

POLYCILLIN

BRISTOL	EQ 250MG BASE	N050310 001	
	EQ 500MG BASE	N050310 002	

PRINCIPEN

APOTHECON	EQ 250MG BASE	A062888 001	Mar 04, 1988
	EQ 500MG BASE	A062888 002	Mar 04, 1988
BRISTOL MYERS SQUIBB	EQ 250MG BASE	A061392 001	
	EQ 500MG BASE	A061392 002	

PRINCIPEN '250'

APOTHECON	EQ 250MG BASE	A062157 002	
+	EQ 250MG BASE	N050056 001	

PRINCIPEN '500'

APOTHECON	EQ 500MG BASE	A062157 001	
+	EQ 500MG BASE	N050056 002	

TOTACILLIN

GLAXOSMITHKLINE	EQ 250MG BASE	A060060 001	
	EQ 250MG BASE	A062212 001	
	EQ 500MG BASE	A060060 002	
	EQ 500MG BASE	A062212 002	

FOR SUSPENSION;ORAL

AMCILL

PARKE DAVIS	EQ 125MG BASE/5ML	A062030 001	
	EQ 250MG BASE/5ML	A062030 002	

AMPICILLIN TRIHYDRATE

BELCHER PHARMS	EQ 125MG BASE/5ML	A061601 001	
	EQ 250MG BASE/5ML	A061601 002	
DAVA PHARMS INC	EQ 125MG BASE/5ML	A062982 001	Feb 10, 1989
	EQ 250MG BASE/5ML	A062982 002	Feb 10, 1989
MYLAN	EQ 125MG BASE/5ML	A061829 002	
	EQ 250MG BASE/5ML	A061829 001	
PUREPAC PHARM	EQ 125MG BASE/5ML	A061980 001	
	EQ 250MG BASE/5ML	A061980 002	
TEVA	EQ 125MG BASE/5ML	A061370 001	
	EQ 250MG BASE/5ML	A061370 002	

OMNIPEN (AMPICILLIN)

WYETH AYERST	100MG/ML	A060625 001	
	125MG/5ML	A060625 002	
	250MG/5ML	A060625 003	
	500MG/5ML	A060625 004	

PENBRITIN

WYETH AYERST	EQ 100MG BASE/ML	N050019 001	
	EQ 125MG BASE/5ML	N050019 002	
	EQ 250MG BASE/5ML	N050019 003	

PFIZERPEN-A

PFIZER	EQ 125MG BASE/5ML	A062049 001	
	EQ 250MG BASE/5ML	A062049 002	

POLYCILLIN

APOTHECON	EQ 125MG BASE/5ML	A062297 001	
	EQ 250MG BASE/5ML	A062297 002	
BRISTOL	EQ 100MG BASE/ML	N050308 004	
	EQ 125MG BASE/5ML	N050308 001	
	EQ 250MG BASE/5ML	N050308 002	
	EQ 500MG BASE/5ML	N050308 003	

PRINCIPEN

APOTHECON	EQ 100MG BASE/ML	A061394 001	
	EQ 125MG BASE/5ML	A061394 002	
	EQ 250MG BASE/5ML	A061394 003	

PRINCIPEN '125'

APOTHECON	EQ 125MG BASE/5ML	A060127 002	
	EQ 125MG BASE/5ML	A062151 001	

PRINCIPEN '250'

APOTHECON	EQ 250MG BASE/5ML	A060127 001	
	EQ 250MG BASE/5ML	A062151 002	

TOTACILLIN

GLAXOSMITHKLINE	EQ 125MG BASE/5ML	A060666 001	
	EQ 125MG BASE/5ML	A062223 001	
	EQ 250MG BASE/5ML	A060666 002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATEFOR SUSPENSION;ORAL
TOTACILLIN

EQ 250MG BASE/5ML A062223 002

TABLET, CHEWABLE;ORAL
POLYCILLIN

BRISTOL EQ 125MG BASE N050093 001

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

CAPSULE;ORAL

PRINCIPEN W/ PROBENECID

APOTHECON EQ 389MG BASE;111MG A062150 001

EQ 389MG BASE;111MG N050488 001

FOR SUSPENSION;ORAL

POLYCILLIN-PRB

APOTHECON EQ 3.5GM BASE/BOT;1GM/BOT A061898 001

BRISTOL EQ 3.5GM BASE/BOT;1GM/BOT N050457 001

PROBAMPACIN

COSETTE EQ 3.5GM BASE/BOT;1GM/BOT A061741 001

AMPRENAVIR

CAPSULE;ORAL

AGENERASE

GLAXOSMITHKLINE 50MG N021007 001 Apr 15, 1999

150MG N021007 002 Apr 15, 1999

SOLUTION;ORAL

AGENERASE

+ GLAXOSMITHKLINE 15MG/ML ** N021039 001 Apr 15, 1999

ANAGRELIDE HYDROCHLORIDE

CAPSULE;ORAL

AGRYLIN

+ TAKEDA PHARMS USA EQ 1MG BASE ** N020333 002 Mar 14, 1997

ANAGRELIDE HYDROCHLORIDE

BARR EQ 0.5MG BASE A076530 001 Apr 18, 2005

EQ 1MG BASE A076530 002 Apr 18, 2005

CHARTWELL RX EQ 0.5MG BASE A076683 001 Apr 18, 2005

EQ 1MG BASE A076683 002 Apr 18, 2005

NATCO PHARMA USA EQ 0.5MG BASE A076811 001 Apr 18, 2005

EQ 1MG BASE A076811 002 Apr 18, 2005

RISING EQ 0.5MG BASE A077613 001 Jun 27, 2006

EQ 1MG BASE A077613 002 Jun 27, 2006

ROXANE EQ 0.5MG BASE A076489 001 Apr 18, 2005

EQ 1MG BASE A076489 002 Apr 18, 2005

WATSON LABS EQ 0.5MG BASE A076417 001 Apr 18, 2005

EQ 1MG BASE A076417 002 Apr 18, 2005

ANASTROZOLE

TABLET;ORAL

ANASTROZOLE

APOTEX INC 1MG A200654 001 May 11, 2012

CHARTWELL MOLECULAR 1MG A091331 001 Jan 05, 2011

CHARTWELL RX 1MG A090732 001 Jun 28, 2010

FRESENIUS KABI USA 1MG A090088 001 Jun 28, 2010

HIKMA 1MG A078485 001 Jun 28, 2010

IMPAX LABS INC 1MG A091242 001 May 31, 2012

MYLAN 1MG A091051 001 Jun 28, 2010

SANDOZ 1MG A079007 001 Jun 28, 2010

SUN PHARM INDS LTD 1MG A091177 001 Jul 15, 2011

SYNTHON PHARMS 1MG A078322 001 Jun 28, 2010

WATSON LABS TEVA 1MG A078984 001 Jun 28, 2010

ANGIOTENSIN II ACETATE

SOLUTION;INTRAVENOUS

GIAPREZA

+ LA JOLLA PHARMA EQ 5MG BASE/2ML (EQ 2.5MG BASE/ML) N209360 002 Dec 21, 2017

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ANILERIDINE HYDROCHLORIDE

TABLET; ORAL

LERITINE

MERCK

EQ 25MG BASE

N010585 002

ANILERIDINE PHOSPHATE

INJECTABLE; INJECTION

LERITINE

MERCK

25MG/ML

N010520 003

ANISINDIONE

TABLET; ORAL

MIRADON

SCHERING

50MG

N010909 003

ANISOTROPINE METHYLBROMIDE

TABLET; ORAL

ANISOTROPINE METHYLBROMIDE

WATSON LABS

50MG

A086046 001

VALPIN 50

ENDO PHARMS

50MG

N013428 001

ANTAZOLINE PHOSPHATE; NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

VASOCON-A

NOVARTIS

0.5%; 0.05%

N018746 002 Jul 11, 1994

APIXABAN

TABLET; ORAL

APIXABAN

AUROBINDO PHARMA LTD 2.5MG

A210026 001 May 26, 2023

5MG

A210026 002 May 26, 2023

BIONPHARMA 2.5MG

A210152 001 Apr 08, 2020

5MG

A210152 002 Apr 08, 2020

BRECKENRIDGE 2.5MG

A209845 001 Jul 26, 2021

5MG

A209845 002 Jul 26, 2021

MICRO LABS 2.5MG

A210013 001 Dec 23, 2019

5MG

A210013 002 Dec 23, 2019

MYLAN 2.5MG

A210128 001 Dec 23, 2019

5MG

A210128 002 Dec 23, 2019

ZYDUS PHARMS 2.5MG

A210185 001 Feb 27, 2023

5MG

A210185 002 Feb 27, 2023

APOMORPHINE HYDROCHLORIDE

FILM; SUBLINGUAL

KYNMOBI

+ SUMITOMO PHARMA AM 10MG

N210875 001 May 21, 2020

+ 15MG

N210875 002 May 21, 2020

+ 20MG

N210875 003 May 21, 2020

+ 25MG

N210875 004 May 21, 2020

+ 30MG

N210875 005 May 21, 2020

INJECTABLE; SUBCUTANEOUS

APOKYN

MDD US

20MG/2ML (10MG/ML)

N021264 001 Apr 20, 2004

APREMILAST

TABLET; ORAL

APREMILAST

AMNEAL 10MG

A211782 001 Jun 30, 2021

20MG

A211782 002 Jun 30, 2021

30MG

A211782 003 Jun 30, 2021

AUROBINDO PHARMA LTD 10MG

A211716 001 Jul 19, 2023

20MG

A211716 002 Jul 19, 2023

30MG

A211716 003 Jul 19, 2023

DR REDDYS 10MG

A211756 001 Jul 14, 2023

20MG

A211756 002 Jul 14, 2023

30MG

A211756 003 Jul 14, 2023

TEVA PHARMS USA INC 10MG

A211897 001 Aug 18, 2022

20MG

A211897 002 Aug 18, 2022

30MG

A211897 003 Aug 18, 2022

UNICHEM 10MG

A211819 001 Feb 17, 2021

20MG

A211819 002 Feb 17, 2021

30MG

A211819 003 Feb 17, 2021

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

APREPITANT

CAPSULE; ORAL

EMEND

+ MERCK

40MG **

N021549 003 Jun 30, 2006

ARBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

GENESA

GENSIA AUTOMEDICS

0.05MG/ML

N020420 001 Sep 12, 1997

ARDEPARIN SODIUM

INJECTABLE; INJECTION

NORMIFLO

+ PHARMACIA AND UPJOHN

5,000 UNITS/0.5ML **

N020227 002 May 23, 1997

+

10,000 UNITS/0.5ML **

N020227 001 May 23, 1997

ARFORMOTEROL TARTRATE

SOLUTION; INHALATION

ARFORMOTEROL TARTRATE

GLENMARK PHARMS LTD

EQ 0.015MG BASE/2ML

A213132 001 Jun 22, 2021

ARGATROBAN

INJECTABLE; INJECTION

ARGATROBAN

+ SANDOZ

250MG/2.5ML (100MG/ML)

N020883 001 Jun 30, 2000

INJECTABLE; INTRAVENOUS

ARGATROBAN IN 0.9% SODIUM CHLORIDE

TEVA PHARMS USA

250MG/250ML (1MG/ML)

N206769 001 Dec 15, 2014

ARGATROBAN IN SODIUM CHLORIDE

+ CIPLA

50MG/50ML (1MG/ML) **

N022434 001 Jun 29, 2011

SOLUTION; INTRAVENOUS

ARGATROBAN IN DEXTROSE

SANDOZ

125MG/125ML (1MG/ML)

N201743 001 May 09, 2011

ARIPIPRAZOLE

INJECTABLE; INTRAMUSCULAR

ABILIFY

OTSUKA

9.75MG/1.3ML (7.5MG/ML)

N021866 001 Sep 20, 2006

SOLUTION; ORAL

ABILIFY

+ OTSUKA

1MG/ML **

N021713 001 Dec 10, 2004

TABLET; ORAL

ARIPIPRAZOLE

MYLAN

2MG

A206240 001 Sep 19, 2018

5MG

A206240 002 Sep 19, 2018

10MG

A206240 003 Sep 19, 2018

15MG

A206240 004 Sep 19, 2018

20MG

A206240 005 Sep 19, 2018

30MG

A206240 006 Sep 19, 2018

TEVA PHARMS USA

2MG

A078607 001 Apr 28, 2015

5MG

A078607 002 Apr 28, 2015

10MG

A078608 001 Apr 28, 2015

15MG

A078708 001 Apr 28, 2015

20MG

A078708 002 Apr 28, 2015

30MG

A078708 003 Apr 28, 2015

ZYDUS PHARMS

2MG

A090472 001 Jan 07, 2019

5MG

A090472 002 Jan 07, 2019

10MG

A090472 003 Jan 07, 2019

15MG

A090472 004 Jan 07, 2019

20MG

A090472 005 Jan 07, 2019

30MG

A090472 006 Jan 07, 2019

TABLET, ORALLY DISINTEGRATING; ORAL

ABILIFY

+ OTSUKA

10MG **

N021729 002 Jun 07, 2006

+

15MG **

N021729 003 Jun 07, 2006

+

20MG **

N021729 004 Jun 07, 2006

+

30MG **

N021729 005 Jun 07, 2006

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ARMODAFINIL

TABLET; ORAL

ARMODAFINIL

COREPHARMA	50MG	A201514 001	Mar 25, 2019
	150MG	A201514 002	Mar 25, 2019
	250MG	A201514 003	Mar 25, 2019
WATSON LABS INC	100MG	A200156 002	Aug 29, 2012
	200MG	A200156 004	Aug 29, 2012
NUVIGIL			
+ CEPHALON	100MG **	N021875 002	Mar 26, 2009

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

ARSENIC TRIOXIDE

INGENUS PHARMS LLC	2MG/ML	A209315 002	Jan 14, 2021
TRISENOX			
+ CEPHALON	1MG/ML **	N021248 001	Sep 25, 2000

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

HOSPIRA	4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)	A079138 001	Jun 18, 2010
ULTACAN			
HANSAMED INC	4%;EQ 0.0085MG BASE/1.7ML (4%; EQ 0.005MG BASE/ML)	A201751 001	Jul 11, 2017
ULTACAN FORTE			
HANSAMED INC	4%;EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)	A201750 001	Jul 11, 2017

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PHYTONADIONE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

FOR SOLUTION; INTRAVENOUS

M.V.I. PEDIATRIC

+ HOSPIRA	80MG/VIAL;0.02MG/VIAL;0.001MG/VIAL;5MG/VIAL;0.01MG/VIAL;0.14MG/VIAL;17MG/VIAL;0.2MG/VIAL;1MG/VIAL;1.4MG/VIAL;EQ 1.2MG BASE/VIAL;0.7MG/VIAL;7MG/VIAL	N018920 001	Sep 21, 2000
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ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION

BEROCCA PN

ROCHE	50MG/ML;0.03MG/ML;0.0025MG/ML;7.5MG/ML;100IU/ML;0.2MG/ML;20MG/ML;2MG/ML;1.8MG/ML;	N006071 003	Oct 10, 1985
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ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.C. 9+3

ABRAXIS PHARM	10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;20MG/ML;0.006MG/ML;0.05MCG/ML;1.5MG/ML;2IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/	N018440 002	Aug 08, 1985
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M.V.I.-12 ADULT

HOSPIRA	10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;20MG/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/	N008809 004	Aug 08, 1985
+ HOSPIRA	10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;20MG/ML;0.006MG/ML;0.05MCG/ML;1.5MG/ML;0.0005MG/ML;0.06MG/ML;4MG/ML;0.6MG/ML;0.36MG/ML;0.6MG/ML;0.1MG/ML;1MG/ML	N008809 006	Sep 09, 2004

MVC PLUS

WATSON LABS	10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;20MG/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/	N018439 002	Aug 08, 1985
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E; VITAMIN K

INJECTABLE; INTRAVENOUS

M.V.I. ADULT

+	HOSPIRA	200MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15MG/VIAL;0.005MG/VIAL;0.6MG/VIAL;40MG/VIAL;6MG/VIAL;3.6MG/VIAL;6MG/VIAL;1MG/VIAL;10MG/VIAL;0.15MG/VIAL	N021625 001 Jan 30, 2004
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M.V.I. ADULT (PHARMACY BULK PACKAGE)

+	HOSPIRA	200MG/5ML;0.06MG/5ML;0.005MG/5ML;15MG/5ML;0.005MG/5ML;0.6MG/5ML;40MG/5ML;6MG/5ML;3.6MG/5ML;6MG/5ML;1MG/5ML;10MG/5ML;0.15MG/5ML	N021643 001 Feb 18, 2004
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ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 ADULT

	HOSPIRA	20MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;20IU/ML;0.6MG/ML;4MG/ML;0.4MG/ML;0.36MG/ML	N008809 005 Apr 22, 2004
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ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 LYOPHILIZED

	TELIGENT	100MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15MG/VIAL;5MCG/VIAL;0.4MG/VIAL;40MG/VIAL;4MG/VIAL;3.6MG/VIAL;3MG/VIAL;1MG/VIAL;10MG/VIAL	N018933 002 Aug 08, 1985
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ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION

VITAPED

	HOSPIRA	N/A,80MG/VIAL;N/A,0.02MG/VIAL;N/A,0.001MG/VIAL;400IU/10ML,N/A;N/A,0.14MG/VIAL;N/A,17MG/VIAL;N/A,5MG/VIAL;0.2MG/10ML,N/A;N/A,1MG/VIAL;N/A,1.4MG/VIAL;N/A,1.2MG/VIAL;EQ 2,300 UNITS BASE/10ML,N/A;7 IU/10ML,N/A	N020176 001 Dec 29, 1993
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ASPIRIN

CAPSULE, EXTENDED RELEASE; ORAL

DURLAZA

+	HESP	162.5MG	N200671 001 Sep 04, 2015
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TABLET; ORAL

BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN

	BAYER	500MG	N021317 001 Oct 18, 2001
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TABLET, EXTENDED RELEASE; ORAL

8-HOUR BAYER

	BAYER	650MG	N016030 001
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MEASURIN

	BAYER	650MG	N016030 002
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ASPIRIN; BUTALBITAL

TABLET; ORAL

AXOTAL

	SAVAGE LABS	650MG;50MG	A088305 001 Oct 13, 1983
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ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

	LGM PHARMA	500MG;50MG;40MG	A205230 001 Oct 18, 2021
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	WATSON LABS	325MG;50MG;40MG	A086231 002 Feb 12, 1985
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FIORINAL

+	ALLERGAN	325MG;50MG;40MG **	N017534 005 Apr 16, 1986
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TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

	ACTAVIS ELIZABETH	325MG;50MG;40MG	A086710 002 Aug 23, 1983
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	FOSUN PHARMA	325MG;50MG;40MG	A086398 002 Apr 06, 1984
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	HALSEY	325MG;50MG;40MG	A089448 001 Dec 01, 1986
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+	HIKMA INTL PHARMS	325MG;50MG;40MG	A086162 002 Feb 16, 1984
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	IVAX PHARMS	325MG;50MG;40MG	A085441 002 Oct 31, 1984
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

PURACAP PHARM	325MG;50MG;40MG	A087048	002	Dec 09, 1983
QUANTUM PHARMICS	325MG;50MG;40MG	A088972	001	Jun 18, 1985
WATSON LABS	325MG;50MG;40MG	A086237	002	Mar 23, 1984

FIORINAL

+ ALLERGAN	325MG;50MG;40MG **	N017534	003	Apr 16, 1986
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LANORINAL

LANNETT	325MG;50MG;40MG	A086986	002	Oct 18, 1985
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ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

DR REDDYS LABS SA	325MG;50MG;40MG;30MG	A203335	001	Oct 30, 2015
STRIDES PHARMA	325MG;50MG;40MG;30MG	A075351	001	Mar 05, 1999
WATSON LABS	325MG;50MG;40MG;30MG	A074359	001	Aug 31, 1995

FIORINAL W/CODEINE

+ ALLERGAN	325MG;50MG;40MG;30MG	N019429	003	Oct 26, 1990
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ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

SYNALGOS-DC

+ SUN PHARM INDUSTRIES	356.4MG;30MG;16MG	N011483	004	Sep 06, 1983
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ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

INVAGESIC

SANDOZ	385MG;30MG;25MG	A074817	001	Nov 27, 1996
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INVAGESIC FORTE

SANDOZ	770MG;60MG;50MG	A074817	002	Nov 27, 1996
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NORGESIC

+ BAUSCH	385MG;30MG;25MG **	N013416	003	Oct 27, 1982
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NORGESIC FORTE

+ BAUSCH	770MG;60MG;50MG **	N013416	004	Oct 27, 1982
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ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

SANDOZ	385MG;30MG;25MG	A074654	001	Dec 31, 1996
	770MG;60MG;50MG	A074654	002	Dec 31, 1996
STEVENS J	385MG;30MG;25MG	A074988	001	Apr 30, 1999
	770MG;60MG;50MG	A074988	002	Apr 30, 1999

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

COMPOUND 65

ALRA	389MG;32.4MG;65MG	A084553	002	Aug 17, 1983
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DARVON COMPOUND

XANODYNE PHARM	389MG;32.4MG;32MG	N010996	006	Mar 08, 1983
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DARVON COMPOUND-65

XANODYNE PHARM	389MG;32.4MG;65MG	N010996	007	Mar 08, 1983
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PROPOXYPHENE COMPOUND 65

IVAX SUB TEVA PHARMS	389MG;32.4MG;65MG	A083077	002	Dec 07, 1984
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+ SANDOZ	389MG;32.4MG;65MG	A080044	002	Sep 16, 1983
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TEVA	389MG;32.4MG;65MG	A089025	001	Mar 29, 1985
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PROPOXYPHENE COMPOUND-65

SANDOZ	389MG;32.4MG;65MG	A083101	002	Jun 24, 1985
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PROPOXYPHENE HYDROCHLORIDE W/ ASPIRIN AND CAFFEINE

WATSON LABS	389MG;32.4MG;65MG	A085732	002	Sep 03, 1984
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ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL AND ASPIRIN

CHARTWELL RX	325MG;200MG	A089594	001	Mar 31, 1989
NOVAST LABS	325MG;200MG	A040832	001	Jan 07, 2010
OXFORD PHARMS	325MG;200MG	A040252	001	Dec 10, 1997
SANDOZ	325MG;200MG	A040116	001	Apr 25, 1996

CARISOPRODOL COMPOUND

WATSON LABS	325MG;200MG	A088809	001	Oct 03, 1985
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SOMA COMPOUND

MEDA PHARMS	325MG;200MG **	N012365	005	Jul 11, 1983
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET;ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

INGENUS PHARMS NJ	325MG;200MG;16MG	A040860	001	Jan 07, 2010
OXFORD PHARMS	325MG;200MG;16MG	A040283	001	Dec 29, 1998
SANDOZ	325MG;200MG;16MG	A040118	001	Apr 16, 1996
SOMA COMPOUND W/ CODEINE				
MEDA PHARMS	325MG;200MG;16MG **	N012366	002	Jul 11, 1983

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE;ORAL

AGGRENOX

+ BOEHRINGER INGELHEIM	25MG;200MG **	N020884	001	Nov 22, 1999
ASPIRIN AND DIPYRIDAMOLE				
ANI PHARMS	25MG;200MG	A206964	001	Jan 18, 2017
CHARTWELL MOLECULAR	25MG;200MG	A204552	001	Mar 20, 2019
SUN PHARM	25MG;200MG	A208572	001	Aug 21, 2018

ASPIRIN; HYDROCODONE BITARTRATE

TABLET;ORAL

AZDONE

SCHWARZ PHARMA	500MG;5MG **	A089420	001	Jan 25, 1988
VICOPRIN				
ABBOTT	500MG;5MG	A086333	001	Sep 14, 1983

ASPIRIN; MEPROBAMATE

TABLET;ORAL

EQUAGESIC

SUN PHARM INDUSTRIES	325MG;200MG	N011702	003	Dec 29, 1983
MEPRO-ASPIRIN				
SANDOZ	325MG;200MG	A089127	001	Mar 02, 1987
MEPROBAMATE AND ASPIRIN				
PAR PHARM	325MG;200MG	A089126	001	Aug 19, 1986
MICRAININ				
MEDPOINTE PHARM HLC	325MG;200MG	A084978	001	
Q-GESIC				
QUANTUM PHARMICS	325MG;200MG	A088740	001	Jun 01, 1984

ASPIRIN; METHOCARBAMOL

TABLET;ORAL

METHOCARBAMOL AND ASPIRIN

IVAX SUB TEVA PHARMS	325MG;400MG	A087211	001	Dec 22, 1982
MCNEIL	325MG;400MG	A089193	001	Feb 12, 1986
PAR PHARM	325MG;400MG	A089657	001	Nov 04, 1988
STEVENS J	325MG;400MG	A081145	001	Jan 31, 1995
ROBAXISAL				
+ ROBINS AH	325MG;400MG	N012281	001	

ASPIRIN; OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

YOSPRALA

+ GENUS LIFESCIENCES	81MG;40MG	N205103	001	Sep 14, 2016
+	325MG;40MG	N205103	002	Sep 14, 2016

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

OXYCODONE AND ASPIRIN

ACTAVIS LABS FL INC	325MG;4.8355MG	A090084	001	Mar 22, 2011
DR REDDYS LABS SA	325MG;4.8355MG	A091670	001	Mar 16, 2011

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET;ORAL

CODOXY

HALSEY	325MG;4.5MG;0.38MG	A087464	001	Jul 01, 1982
OXYCODONE AND ASPIRIN				
ANI PHARMS	325MG;4.5MG;0.38MG	A040255	001	Feb 27, 1998
SUN PHARM INDUSTRIES	325MG;4.5MG;0.38MG	A040260	001	Jul 17, 1998
	325MG;4.5MG;0.38MG	A087794	001	May 26, 1982
OXYCODONE AND ASPIRIN (HALF-STRENGTH)				
ROXANE	325MG;2.25MG;0.19MG	A087742	001	Jun 04, 1982
PERCODAN				
ENDO PHARMS	325MG;4.5MG;0.38MG **	N007337	006	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

PERCODAN-DEMI

ENDO PHARMS

325MG; 2.25MG; 0.19MG **

N007337 005

ROXIPRIN

ROXANE

325MG; 4.5MG; 0.38MG

A087743 001 Jun 04, 1982

ASPIRIN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN COMPOUND

+ SANOFI AVENTIS US

325MG; EQ 12.5MG BASE **

N016891 001

ASPIRIN; PRAVASTATIN SODIUM

TABLET; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB

81MG; 20MG

N021387 001 Jun 24, 2003

81MG; 40MG

N021387 002 Jun 24, 2003

81MG; 80MG

N021387 003 Jun 24, 2003

325MG; 20MG

N021387 004 Jun 24, 2003

325MG; 40MG

N021387 005 Jun 24, 2003

325MG; 80MG

N021387 006 Jun 24, 2003

ASPIRIN; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON W/ ASA

XANODYNE PHARM

325MG; 65MG

N010996 005

ASPIRIN; PROPOXYPHENE NAPSYLATE

CAPSULE; ORAL

DARVON-N W/ ASA

AAIPHARMA LLC

325MG; 100MG

N016829 001

TABLET; ORAL

DARVON-N W/ ASA

AAIPHARMA LLC

325MG; 100MG

N016863 001

ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

AMNEAL

EQ 100MG BASE

A209717 001 Jun 01, 2020

CIPLA

EQ 100MG BASE

A200626 001 Aug 09, 2018

EQ 150MG BASE

A200626 002 Aug 09, 2018

EQ 200MG BASE

A200626 003 Aug 09, 2018

EQ 300MG BASE

A200626 004 Aug 09, 2018

MYLAN

EQ 150MG BASE

A208177 001 Sep 24, 2018

EQ 200MG BASE

A208177 002 Sep 24, 2018

EQ 300MG BASE

A208177 003 Sep 24, 2018

ZYDUS PHARMS

EQ 150MG BASE

A210575 001 Jun 04, 2020

EQ 200MG BASE

A210575 002 Jun 04, 2020

EQ 300MG BASE

A210575 003 Jun 04, 2020

REYATAZ

+ BRISTOL MYERS SQUIBB

EQ 100MG BASE **

N021567 001 Jun 20, 2003

+

EQ 150MG BASE **

N021567 002 Jun 20, 2003

ATENOLOL

INJECTABLE; INJECTION

TENORMIN

+ ASTRAZENECA

0.5MG/ML **

N019058 001 Sep 13, 1989

TABLET; ORAL

ATENOLOL

ABLE

25MG

A076907 001 Jul 30, 2004

50MG

A076907 002 Jul 30, 2004

100MG

A076907 003 Jul 30, 2004

APOTHECON

50MG

A073317 001 Mar 20, 1992

100MG

A073318 001 Mar 20, 1992

CHARTWELL RX

25MG

A074265 001 Feb 28, 1994

50MG

A074265 002 Feb 28, 1994

100MG

A074265 003 Feb 28, 1994

MYLAN

25MG

A074126 003 Aug 26, 1998

50MG

A074126 001 Mar 23, 1994

100MG

A074126 002 Mar 23, 1994

NORTHSTAR HLTHCARE

25MG

A078254 001 Sep 25, 2009

50MG

A078254 002 Sep 25, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATENOLOL

TABLET;ORAL

ATENOLOL

	100MG	A078254	003	Sep 25, 2009
NOSTRUM LABS	50MG	A074127	001	Feb 21, 1995
	100MG	A074127	002	Feb 21, 1995
PLIVA	25MG	A074101	001	Jul 17, 1997
	50MG	A074101	002	Jul 17, 1997
	100MG	A074101	003	Jul 17, 1997
SCS	50MG	A073676	001	Oct 30, 1992
	100MG	A073676	002	Oct 30, 1992
STRIDES PHARMA	25MG	A074099	001	Apr 28, 1992
	50MG	A073542	001	Dec 19, 1991
	100MG	A073543	001	Dec 19, 1991
SUN PHARM INDS INC	25MG	A078210	001	Jul 10, 2007
	50MG	A078210	002	Jul 10, 2007
	100MG	A078210	003	Jul 10, 2007
SUN PHARM INDUSTRIES	25MG	A074499	001	Jul 30, 1997
	50MG	A073475	001	Mar 30, 1993
	100MG	A073476	001	Mar 30, 1993
TEVA	50MG	A073315	001	May 28, 1993
	100MG	A073316	001	May 28, 1993
TEVA PHARMS	50MG	A074120	001	Feb 24, 1995
	100MG	A074120	002	Feb 24, 1995
WATSON LABS	50MG	A073352	001	Dec 27, 1991
WATSON LABS TEVA	100MG	A073353	001	Dec 27, 1991

ATENOLOL; CHLORTHALIDONE

TABLET;ORAL

ATENOLOL AND CHLORTHALIDONE

AUROBINDO PHARMA USA	50MG;25MG	A074203	001	Oct 31, 1993
	100MG;25MG	A074203	002	Oct 31, 1993
NOSTRUM LABS	50MG;25MG	A074404	001	May 14, 1998
	100MG;25MG	A074404	002	May 14, 1998
PLIVA	50MG;25MG	A074107	001	Sep 24, 1997
	100MG;25MG	A074107	002	Sep 24, 1997
SUN PHARM INDUSTRIES	50MG;25MG	A073582	002	Apr 29, 1993
	100MG;25MG	A073582	001	Apr 29, 1993

ATOMOXETINE HYDROCHLORIDE

CAPSULE;ORAL

ATOMOXETINE HYDROCHLORIDE

STRIDES PHARMA	10MG	A079021	001	Feb 18, 2021
	18MG	A079021	002	Feb 18, 2021
	25MG	A079021	003	Feb 18, 2021
	40MG	A079021	004	Feb 18, 2021
	60MG	A079021	005	Feb 18, 2021
	80MG	A079021	006	Feb 18, 2021
	100MG	A079021	007	Feb 18, 2021
STRATTERA				
LILLY	5MG	N021411	001	Nov 26, 2002

ATORVASTATIN CALCIUM

TABLET;ORAL

ATORVASTATIN CALCIUM

LEPU PHARM	EQ 10MG BASE	A216848	001	Nov 03, 2022
	EQ 20MG BASE	A216848	002	Nov 03, 2022
	EQ 40MG BASE	A216848	003	Nov 03, 2022
	EQ 80MG BASE	A216848	004	Nov 03, 2022
PERRIGO R AND D	EQ 10MG BASE	A208478	001	Jun 23, 2020
	EQ 20MG BASE	A208478	002	Jun 23, 2020
TEVA PHARMS	EQ 10MG BASE	A078773	001	May 29, 2012
	EQ 20MG BASE	A078773	002	May 29, 2012
	EQ 40MG BASE	A078773	003	May 29, 2012
	EQ 80MG BASE	A078773	004	May 29, 2012

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET; ORAL

LIPTRUZET

+	ORGANON	EQ 10MG BASE;10MG **	N200153 001	May 03, 2013
+		EQ 20MG BASE;10MG **	N200153 002	May 03, 2013
+		EQ 40MG BASE;10MG **	N200153 003	May 03, 2013
+		EQ 80MG BASE;10MG **	N200153 004	May 03, 2013

ATOVAQUONE

TABLET; ORAL

MEPRON

+	GLAXOSMITHKLINE LLC	250MG **	N020259 001	Nov 25, 1992
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ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

	BAXTER HLTHCARE	10MG/ML	A074824 001	Sep 30, 1997
	BAXTER HLTHCARE CORP	10MG/ML	A074753 001	Jan 23, 1997
	HOSPIRA	10MG/ML	A074632 001	Dec 23, 1996
		10MG/ML	A074740 001	Mar 28, 1997
	MYLAN LABS LTD	10MG/ML	A206096 001	Jun 22, 2017
	TEVA PARENTERAL	10MG/ML	A074784 001	Jun 11, 1997
	WATSON PHARMS TEVA	10MG/ML	A074945 001	Jul 28, 1998
	ATRACURIUM BESYLATE PRESERVATIVE FREE			
	BAXTER HLTHCARE	10MG/ML	A074825 001	Sep 30, 1997
	BAXTER HLTHCARE CORP	10MG/ML	A074768 001	Jan 23, 1997
	HOSPIRA	10MG/ML	A074633 001	Dec 23, 1996
		10MG/ML	A074639 001	Mar 25, 1997
		10MG/ML	A074741 001	Mar 28, 1997
	MYLAN LABS LTD	10MG/ML	A206001 001	Apr 07, 2017
	WATSON LABS INC	10MG/ML	A074944 001	Jul 28, 1998
	TRACRIUM			
+	HOSPIRA	10MG/ML **	N018831 002	Jun 20, 1985
	TRACRIUM PRESERVATIVE FREE			
+	HOSPIRA	10MG/ML **	N018831 001	Nov 23, 1983

ATROPINE

SOLUTION; INTRAMUSCULAR

ATROPEN

+	MMT	EQ 0.25MG SULFATE/0.3ML	N017106 004	Sep 17, 2004
+		EQ 0.5MG SULFATE/0.7ML	N017106 003	Jun 19, 2003
+		EQ 1MG SULFATE/0.7ML	N017106 002	Jun 19, 2003
+		EQ 2MG SULFATE/0.7ML	N017106 001	

ATROPINE

	ABBVIE	EQ 2MG SULFATE/0.7ML	A071295 001	Jan 30, 1987
	ATROPINE (AUTOINJECTOR)			
	RAFA LABS LTD	EQ 2MG SULFATE/0.7ML (EQ 2MG SULFATE/0.7ML)	N212319 001	Jul 09, 2018

ATROPINE SULFATE

AEROSOL, METERED; INHALATION

ATROPINE SULFATE

	US ARMY	EQ 0.36MG BASE/INH	N020056 001	Sep 19, 1990
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ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN HALF-STRENGTH

	SEBELA IRELAND LTD	0.025MG;0.5MG	N017744 001	
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ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE

	SCHERER RP	0.025MG;2.5MG	A086440 001	
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SOLUTION; ORAL

COLONAIID

	MEDPOINTE PHARM HLC	0.025MG/5ML;2.5MG/5ML	A085735 001	
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LOMANATE

	ALPHARMA US PHARMS	0.025MG/5ML;2.5MG/5ML	A085746 001	
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LOMOTIL

	GD SEARLE LLC	0.025MG/5ML;2.5MG/5ML	N012699 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

COLONOID

MEDPOINTE PHARM HLC 0.025MG;2.5MG A085737 001

DI-ATRO

MD PHARM 0.025MG;2.5MG A085266 001

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

ABLE 0.025MG;2.5MG A040395 001 Nov 27, 2000

ASCOT 0.025MG;2.5MG A087934 001 Jul 19, 1983

DR REDDYS LABS SA 0.025MG;2.5MG A210789 001 Jun 03, 2020

FOSUN PHARMA 0.025MG;2.5MG A086173 001

HEATHER 0.025MG;2.5MG A086798 001

HIKMA 0.025MG;2.5MG A087765 001 Mar 15, 1982

INWOOD LABS 0.025MG;2.5MG A085509 001

KV PHARM 0.025MG;2.5MG A085659 001

LEDERLE 0.025MG;2.5MG A086950 001

PARKE DAVIS 0.025MG;2.5MG A087131 001

PVT FORM 0.025MG;2.5MG A085766 001

R AND S PHARMA 0.025MG;2.5MG A085035 001

ROXANE 0.025MG;2.5MG A086057 001

STRIDES PHARMA 0.025MG;2.5MG A040357 001 May 02, 2000

SUN PHARM INDUSTRIES 0.025MG;2.5MG A085506 001

UPSHER SMITH LABS 0.025MG;2.5MG A210571 001 Aug 31, 2018

USL PHARMA 0.025MG;2.5MG A087842 001 Mar 29, 1982

VALEANT PHARM INTL 0.025MG;2.5MG A087195 001 Feb 16, 1982

WATSON LABS 0.025MG;2.5MG A085876 001

LO-TROL

VANGARD 0.025MG;2.5MG A088009 001 Mar 25, 1983

LOGEN

SUPERPHARM 0.025MG;2.5MG A088962 001 May 10, 1985

LONOX

FOSUN PHARMA 0.025MG;2.5MG A085311 002

LOW-QUEL

HALSEY 0.025MG;2.5MG A085211 001

ATROPINE SULFATE; EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON-PLUS

+ MYLAN 0.14MG/ML;10MG/ML N019678 001 Nov 06, 1991

MYLAN INSTITUTIONAL 0.14MG/ML;10MG/ML N019677 001 Nov 06, 1991

ATROPINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

ATROPINE AND DEMEROL

ABBVIE 0.4MG/ML;50MG/ML A087853 001 Nov 26, 1982

0.4MG/ML;75MG/ML A087847 001 Nov 26, 1982

0.4MG/ML;100MG/ML A087848 001 Nov 26, 1982

MEPERIDINE AND ATROPINE SULFATE

WYETH AYERST 0.4MG/ML;50MG/ML A085121 001

0.4MG/ML;75MG/ML A085121 002

0.4MG/ML;100MG/ML A085121 003

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

ATNAA

US ARMY 2.1MG/0.7ML;600MG/2ML N021175 001 Jan 17, 2002

AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION; TOPICAL

SHADE UVAGUARD

+ BAYER HEALTHCARE LLC 3%;7.5%;3% N020045 001 Dec 07, 1992

AZACITIDINE

POWDER; INTRAVENOUS, SUBCUTANEOUS

AZACITIDINE

LUPIN LTD 100MG/VIAL A210748 001 Feb 27, 2019

MYLAN INSTITUTIONAL 100MG/VIAL A204949 001 Apr 28, 2016

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AZATADINE MALEATETABLET; ORAL
OPTIMINE

SCHERING 1MG N017601 001

AZATADINE MALEATE; PSEUDOEPHEDRINE SULFATETABLET, EXTENDED RELEASE; ORAL
TRINALIN

SCHERING 1MG; 120MG N018506 001 Mar 23, 1982

AZATHIOPRINE

TABLET; ORAL

IMURAN

+ SEBELA IRELAND LTD 25MG ** N016324 002

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

IMURAN

+ CASPER PHARMA LLC EQ 100MG BASE/VIAL ** N017391 001

AZELAIC ACID

AEROSOL, FOAM; TOPICAL

AZELAIC ACID

TEVA PHARMS USA 15% A210928 001 Oct 07, 2020

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AZELASTINE HYDROCHLORIDE

EPIC PHARMA LLC 0.05% A203660 001 Nov 08, 2016

OPTIVAR

+ MYLAN SPECIALITY LP 0.05% ** N021127 001 May 22, 2000

SPRAY, METERED; NASAL

ASTELIN

+ MYLAN SPCLT VIATRIS 0.137MG/SPRAY ** N020114 001 Nov 01, 1996

ASTEPRO

+ MYLAN SPECIALITY LP 0.137MG/SPRAY ** N022203 001 Oct 15, 2008

0.2055MG/SPRAY ** N022203 002 Aug 31, 2009

AZELASTINE HYDROCHLORIDE

AKORN 0.2055MG/SPRAY A210032 001 Aug 23, 2019

AMNEAL 0.2055MG/SPRAY A208199 001 Dec 15, 2017

APOTEX INC 0.2055MG/SPRAY A201846 001 Aug 31, 2012

AUROBINDO PHARMA LTD 0.2055MG/SPRAY A212775 001 Nov 12, 2020

HIKMA 0.2055MG/SPRAY A207243 001 Sep 22, 2017

PADAGIS ISRAEL 0.2055MG/SPRAY A202743 001 May 08, 2014

AZILSARTAN KAMEDOXOMIL

TABLET; ORAL

AZILSARTAN MEDOXOMIL

LUPIN LTD EQ 40MG MEDOXOMIL A214489 001 Jul 20, 2022

EQ 80MG MEDOXOMIL A214489 002 Jul 20, 2022

AZITHROMYCIN

CAPSULE; ORAL

ZITHROMAX

+ PFIZER EQ 250MG BASE ** N050670 001 Nov 01, 1991

FOR SUSPENSION; ORAL

AZITHROMYCIN

CHARTWELL RX EQ 100MG BASE/5ML A065488 001 May 15, 2015

EQ 200MG BASE/5ML A065488 002 May 15, 2015

SANDOZ EQ 100MG BASE/5ML A065297 001 Sep 18, 2006

EQ 200MG BASE/5ML A065297 002 Sep 18, 2006

TARO EQ 200MG BASE/5ML A211521 001 Dec 11, 2019

TEVA PHARMS EQ 100MG BASE/5ML A065419 001 Jun 24, 2008

EQ 200MG BASE/5ML A065419 002 Jun 24, 2008

FOR SUSPENSION, EXTENDED RELEASE; ORAL

ZMAX

+ PF PRISM CV EQ 2GM BASE/BOT N050797 001 Jun 10, 2005

INJECTABLE; INJECTION

AZITHROMYCIN

HIKMA EQ 500MG BASE/VIAL A065265 001 Jan 18, 2007

MYLAN ASI EQ 500MG BASE/VIAL A065506 001 Mar 24, 2009

RISING EQ 500MG BASE/VIAL A204732 001 Jan 26, 2017

TEVA PARENTERAL EQ 500MG BASE/VIAL N050809 001 Dec 19, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AZITHROMYCIN

INJECTABLE; INJECTION

AZITHROMYCIN

EQ 2.5GM BASE/VIAL

N050809 002 Dec 19, 2006

TABLET; ORAL

AZITHROMYCIN

APOTEX CORP

EQ 250MG BASE

A065507 001 Jul 13, 2011

EQ 500MG BASE

A065509 001 Jul 13, 2011

EQ 600MG BASE

A065508 001 Jul 13, 2011

AUROBINDO PHARMA USA

EQ 600MG BASE

A065360 001 Jan 08, 2007

MYLAN

EQ 250MG BASE

A065365 001 May 30, 2007

EQ 500MG BASE

A065366 001 May 30, 2007

TEVA

EQ 250MG BASE

A065153 001 Nov 14, 2005

EQ 600MG BASE

A065150 001 Nov 14, 2005

YUNG SHIN PHARM

EQ 500MG BASE

A211318 001 Mar 17, 2021

ZYDUS PHARMS

EQ 250MG BASE

A213275 001 Apr 11, 2023

EQ 500MG BASE

A213056 001 Apr 07, 2023

ZITHROMAX

+ PFIZER

EQ 600MG BASE **

N050730 001 Jun 12, 1996

AZITHROMYCIN DIHYDRATE; TROVAFLOXACIN MESYLATE

FOR SUSPENSION, TABLET; ORAL

TROVAN/ZITHROMAX COMPLIANCE PAK

PFIZER

EQ 1GM BASE, N/A; N/A, EQ 100MG BASE

N050762 001 Dec 18, 1998

AZLOCILLIN SODIUM

INJECTABLE; INJECTION

AZLIN

BAYER PHARMS

EQ 2GM BASE/VIAL

A062388 001 Sep 08, 1982

EQ 2GM BASE/VIAL

A062417 001 Oct 12, 1982

EQ 2GM BASE/VIAL

N050562 001 Sep 03, 1982

EQ 3GM BASE/VIAL

A062388 002 Sep 08, 1982

EQ 3GM BASE/VIAL

A062417 002 Oct 12, 1982

EQ 3GM BASE/VIAL

N050562 002 Sep 03, 1982

EQ 4GM BASE/VIAL

A062388 003 Sep 08, 1982

EQ 4GM BASE/VIAL

A062417 003 Oct 12, 1982

EQ 4GM BASE/VIAL

N050562 003 Sep 03, 1982

AZTREONAM

INJECTABLE; INJECTION

AZACTAM

BRISTOL MYERS SQUIBB 500MG/VIAL **

N050580 001 Dec 31, 1986

AZACTAM IN PLASTIC CONTAINER

BRISTOL MYERS SQUIBB 10MG/ML

N050632 003 May 24, 1989

+ 20MG/ML

N050632 002 May 24, 1989

+ 40MG/ML

N050632 001 May 24, 1989

AZTREONAM

HIKMA

1GM/VIAL

A065286 001 Mar 23, 2011

2GM/VIAL

A065286 002 Mar 23, 2011

BACAMPICILLIN HYDROCHLORIDE

FOR SUSPENSION; ORAL

SPECTROBID

PFIZER

125MG/5ML

N050556 001 Mar 23, 1982

TABLET; ORAL

SPECTROBID

PFIZER

400MG

N050520 001

800MG

N050520 002 Sep 12, 1983

BACITRACIN

OINTMENT; OPHTHALMIC

BACIGUENT

PHARMACIA AND UPJOHN 500 UNITS/GM

A060734 001

BACITRACIN

LILLY

500 UNITS/GM

A060687 001

PHARMADERM

500 UNITS/GM

A062158 001

PHARMAFAIR

500 UNITS/GM

A062453 001 Mar 28, 1984

OINTMENT; TOPICAL

BACITRACIN

COMBE

500 UNITS/GM

A062799 001 May 14, 1987

NASKA

500 UNITS/GM

A062857 001 Nov 13, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BACITRACIN

POWDER;FOR RX COMPOUNDING

BACI-RX

X GEN PHARMS 5,000,000 UNITS/BOT A061580 001

BACITRACIN

APOTHEKERNES 5,000,000 UNITS/BOT A061699 001

PADDOCK LLC 5,000,000 UNITS/BOT A062456 001 Jul 27, 1983

BACITRACIN ZINC

POWDER;FOR RX COMPOUNDING

ZIBA-RX

X GEN PHARMS 500,000 UNITS/BOT A061737 001

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

CORTISPORIN

+ CASPER PHARMA LLC 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM ** N050416 002

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

AKORN 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A065213 001 Jul 25, 2012

ZINC BACITRACIN,NEOMYCIN SULFATE,POLYMYXIN B SULFATE & HYDROCORTISONE

PHARMAFAIR 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062389 001 Jul 02, 1982

OINTMENT;TOPICAL

CORTISPORIN

+ MONARCH PHARMS 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM N050168 002 May 04, 1984

NEOMYCIN & POLYMYXIN B SULFATES & BACITRACIN ZINC & HYDROCORTISONE

PHARMAFAIR 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM A062381 001 Sep 06, 1985

BACITRACIN ZINC; LIDOCAINE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;TOPICAL

LANABIOTIC

COMBE 400 UNITS/GM;40MG/GM;EQ 5MG BASE/GM;5,000 UNITS/GM A062499 001 Jun 03, 1985

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE

PHARMAFAIR 400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062386 001 Sep 09, 1982

BACITRACIN-NEOMYCIN-POLYMYXIN

PHARMADERM 400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM A062167 001

NEO-POLYICIN

DOW PHARM 500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM A060647 001

NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC

AKORN 400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM A065088 001 Feb 06, 2004

OINTMENT;TOPICAL

BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE

NASKA 400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM A062833 001 Nov 09, 1987

BACITRACIN ZINC; POLYMYXIN B SULFATE

AEROSOL;TOPICAL

POLYSPORIN

GLAXOSMITHKLINE 10,000 UNITS/GM;2,000,000 UNITS/GM N050167 002 Mar 01, 1985

OINTMENT;OPHTHALMIC

OCUMYCIN

PHARMAFAIR 500 UNITS/GM;10,000 UNITS/GM A062430 001 Apr 08, 1983

POLYSPORIN

MONARCH PHARMS 500 UNITS/GM;10,000 UNITS/GM ** A061229 001

OINTMENT;TOPICAL

BACITRACIN ZINC-POLYMYXIN B SULFATE

NASKA 500 UNITS/GM;10,000 UNITS/GM A062849 001 Nov 13, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

ALTANA	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A060731	002	
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BACITRACIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

MYCITRACIN

PHARMACIA AND UPJOHN	500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A061048	001	
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BACITRACIN; POLYMYXIN B SULFATE

DISC;TOPICAL

LANABIOTIC

COMBE	500 UNITS/GM;5,000 UNITS/GM	N050598	001	Sep 22, 1986
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BACLOFEN

SOLUTION;ORAL

OZOBAX

+ METACEL PHARMS LLC

5MG/5ML

N208193	001	Sep 18, 2019
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TABLET;ORAL

BACLOFEN

APPCO

10MG

A090334	001	Feb 18, 2010
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20MG

A090334	002	Feb 18, 2010
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MYLAN

10MG

A077181	001	Jul 29, 2005
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20MG

A077121	002	Jul 29, 2005
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SUN PHARM INDS INC

10MG

A077862	001	Aug 14, 2006
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20MG

A077862	002	Aug 14, 2006
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TEVA

10MG

A073043	001	Feb 27, 1992
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20MG

A073044	001	Feb 27, 1992
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USL PHARMA

10MG

A071260	001	May 06, 1988
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20MG

A071261	001	May 06, 1988
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WATSON LABS

10MG

A072824	001	Sep 18, 1991
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10MG

A073092	001	Jan 28, 1994
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10MG

A074698	001	Aug 20, 1996
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20MG

A072825	001	Sep 18, 1991
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20MG

A073093	001	Jan 28, 1994
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20MG

A074698	002	Aug 20, 1996
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Lioresal

+ NOVARTIS

10MG **

N017851	001	
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+

20MG **

N017851	003	Jan 20, 1982
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TABLET, ORALLY DISINTEGRATING;ORAL

KEMSTRO

+ UCB INC

10MG

N021589	001	Oct 30, 2003
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+

20MG

N021589	002	Oct 30, 2003
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BALOXAVIR MARBOXIL

TABLET;ORAL

XOFLUZA

+ GENENTECH INC

20MG

N210854	001	Oct 24, 2018
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BALSALAZIDE DISODIUM

CAPSULE;ORAL

BALSALAZIDE DISODIUM

DASH PHARMS NATCO

750MG

A077807	001	Dec 28, 2007
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TABLET;ORAL

BALSALAZIDE DISODIUM

STRIDES PHARMA

1.1GM

A206336	001	Sep 08, 2015
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GIAZO

+ VALEANT PHARMS INTL

1.1GM **

N022205	001	Feb 03, 2012
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BARIUM SULFATE

FOR SUSPENSION;ORAL

E-Z-CAT DRY

+ BRACCO

40% (9GM/POUCH)

N208036	003	Jan 03, 2017
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

BECLOVENT

GLAXOSMITHKLINE 0.042MG/INH

N018153 001

QVAR 40

+ TEVA BRANDED PHARM 0.04MG/INH **

N020911 002 Sep 15, 2000

QVAR 80

+ TEVA BRANDED PHARM 0.08MG/INH **

N020911 001 Sep 15, 2000

VANCERIL

SCHERING 0.042MG/INH

N017573 001

VANCERIL DOUBLE STRENGTH

SCHERING 0.084MG/INH

N020486 001 Dec 24, 1996

AEROSOL, METERED; NASAL

BECONASE

GLAXOSMITHKLINE 0.042MG/INH

N018584 001

VANCENASE

SCHERING 0.042MG/INH

N018521 001

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED; NASAL

BECONASE AQ

+ GLAXOSMITHKLINE EQ 0.042MG DIPROP/SPRAY

N019389 001 Jul 27, 1987

VANCENASE AQ

SCHERING EQ 0.042MG DIPROP/SPRAY

N019589 001 Dec 23, 1987

EQ 0.084MG DIPROP/SPRAY

N020469 001 Jun 26, 1996

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

GENPHARM 5MG

A076476 001 Feb 11, 2004

10MG

A076476 002 Feb 11, 2004

20MG

A076476 003 Feb 11, 2004

40MG

A076476 004 Feb 11, 2004

RISING 5MG

A076430 001 Feb 11, 2004

10MG

A076430 002 Feb 11, 2004

20MG

A076430 003 Feb 11, 2004

40MG

A076430 004 Feb 11, 2004

LOTENSIN

+ VALIDUS PHARMS 5MG **

N019851 001 Jun 25, 1991

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ANI PHARMS 5MG; 6.25MG

A076348 001 Feb 11, 2004

10MG; 12.5MG

A076348 002 Feb 11, 2004

20MG; 12.5MG

A076348 003 Feb 11, 2004

20MG; 25MG

A076348 004 Feb 11, 2004

AUROBINDO PHARMA USA 5MG; 6.25MG

A076688 001 Feb 11, 2004

10MG; 12.5MG

A076688 002 Feb 11, 2004

20MG; 12.5MG

A076688 003 Feb 11, 2004

20MG; 25MG

A076688 004 Feb 11, 2004

MYLAN PHARMS INC 5MG; 6.25MG

A076612 001 Feb 11, 2004

10MG; 12.5MG

A076612 002 Feb 11, 2004

20MG; 12.5MG

A076612 003 Feb 11, 2004

20MG; 25MG

A076612 004 Feb 11, 2004

SUN PHARM INDS LTD 5MG; 6.25MG

A077483 001 Sep 08, 2005

10MG; 12.5MG

A077483 002 Sep 08, 2005

20MG; 12.5MG

A077483 003 Sep 08, 2005

20MG; 25MG

A077483 004 Sep 08, 2005

LOTENSIN HCT

+ VALIDUS PHARMS 5MG; 6.25MG **

N020033 001 May 19, 1992

BENDAMUSTINE HYDROCHLORIDE

POWDER; IV (INFUSION)

BENDAMUSTINE HYDROCHLORIDE

MYLAN LABS LTD 25MG/VIAL

A204104 001 Apr 28, 2023

100MG/VIAL

A204104 002 Apr 28, 2023

NANG KUANG PHARM CO 25MG/VIAL

A206554 001 Jun 07, 2023

100MG/VIAL

A206554 002 Jun 07, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENDAMUSTINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

BENDAMUSTINE HYDROCHLORIDE

+	HOSPIRA	25MG/ML (25MG/ML)	N211530 001	Dec 15, 2022
+		100MG/4ML (25MG/ML)	N211530 002	Dec 15, 2022
+		200MG/8ML (25MG/ML)	N211530 003	Dec 15, 2022

SOLUTION; IV (INFUSION)

TREANDA

+	CEPHALON	45MG/0.5ML (90MG/ML)	N022249 003	Sep 13, 2013
+		180MG/2ML (90MG/ML)	N022249 004	Sep 13, 2013

BENDROFLUMETHIAZIDE

TABLET; ORAL

NATURETIN-10

	APOTHECON	10MG	N012164 003	
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NATURETIN-2.5

	APOTHECON	2.5MG	N012164 001	
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NATURETIN-5

	APOTHECON	5MG	N012164 002	
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BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL

CORZIDE

+	KING PHARMS LLC	5MG; 40MG **	N018647 001	May 25, 1983
+		5MG; 80MG **	N018647 002	May 25, 1983

NADOLOL AND BENDROFLUMETHIAZIDE

	IMPAX LABS	5MG; 40MG	A077833 001	Mar 30, 2007
		5MG; 80MG	A077833 002	Mar 30, 2007
	MYLAN	5MG; 40MG	A078688 001	Feb 15, 2008
		5MG; 80MG	A078688 002	Feb 15, 2008

BENOXINATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BENOXINATE HYDROCHLORIDE

	SOLA BARNES HIND	0.4%	A084149 001	
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BENTIROMIDE

SOLUTION; ORAL

CHYMEX

	SAVAGE LABS	500MG/7.5ML	N018366 001	Dec 29, 1983
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BENZONATATE

CAPSULE; ORAL

BENZONATATE

	CHARTWELL RX	100MG	A210562 001	Nov 09, 2018
		150MG	A210562 002	Nov 09, 2018
		200MG	A210562 003	Nov 09, 2018
	MIKART	100MG	A040851 001	Nov 09, 2009
		150MG	A040851 002	Nov 09, 2009
		200MG	A040851 003	Nov 09, 2009
	NESHER PHARMS	100MG	A040795 001	Oct 31, 2007
		200MG	A040795 002	Oct 31, 2007
	SUN PHARM INDS INC	100MG	A040587 001	Mar 19, 2008
		200MG	A040587 002	Mar 19, 2008
	THEPHARMANETWORK LLC	100MG	A040627 001	Mar 30, 2007
		150MG	A040627 003	Sep 24, 2014
		200MG	A040627 002	Jul 25, 2007

TESSALON

+	PFIZER	200MG **	N011210 003	Jun 25, 1999
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BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

BENZACLIN

+	BAUSCH	5%;EQ 1% BASE	N050756 001	Dec 21, 2000
		5%;EQ 1% BASE	N050756 002	Apr 20, 2007

CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

	CHARTWELL RX	5%;1.2%	A203688 001	Aug 25, 2016
	ENCUBE	2.5%;EQ 1.2% BASE	A207194 001	Aug 19, 2019
		5%;EQ 1% BASE	A204087 001	Jun 27, 2017

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

AKTIPAK

+ BIOFRONTERA 5%;3%

N050769 001 Nov 27, 2000

ERYTHROMYCIN AND BENZOYL PEROXIDE

ENCUBE 5%;3%

A065112 001 Mar 29, 2004

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

AVET LIFESCIENCES 50MG

A202061 001 Jan 27, 2012

EPIC PHARMA LLC 50MG

A040714 001 Oct 29, 2007

IMPAX LABS 50MG

A040845 001 Nov 18, 2008

SCINOPHARM TAIWAN 50MG

A040578 001 Apr 17, 2006

SPECGX LLC 50MG

A040773 001 Apr 25, 2007

TEDOR PHARM 25MG

A040747 002 Nov 20, 2015

50MG

A040747 001 Mar 30, 2007

DIDREX

+ PFIZER 25MG **

N012427 003

+ 50MG **

N012427 002

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

EMETE-CON

PFIZER EQ 50MG BASE/VIAL

N016820 001

SUPPOSITORY; RECTAL

EMETE-CON

ROERIG EQ 100MG BASE

N016818 006

BENZTHIAZIDE

TABLET; ORAL

AQUATAG

SOLVAY 25MG

N016001 001

50MG

N016001 002

BENZTHIAZIDE

PVT FORM 50MG

A083206 001

EXNA

AH ROBINS INC 50MG

N012489 001

FOVANE

PFIZER 50MG

N012128 002

URESE

PFIZER 25MG

N012128 003

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

LUITPOLD 1MG/ML

A091152 001 Mar 29, 2010

COGENTIN

+ EPIC PHARMA LLC 1MG/ML **

N012015 001

TABLET; ORAL

BENZTROPINE MESYLATE

AIPING PHARM INC 0.5MG

A040103 001 Dec 12, 1996

1MG

A040103 002 Dec 12, 1996

2MG

A040103 003 Dec 12, 1996

LANNETT CO INC 0.5MG **

A088877 001 Apr 11, 1985

1MG **

A088894 001 Apr 11, 1985

2MG **

A088895 001 Apr 11, 1985

OXFORD PHARMS 0.5MG

A040706 002 Feb 14, 2008

1MG

A040706 003 Feb 14, 2008

2MG

A040706 001 Feb 14, 2008

QUANTUM PHARMICS 0.5MG

A088514 001 Jan 31, 1984

1MG

A088510 001 Jan 31, 1984

2MG

A088511 001 Jan 31, 1984

USL PHARMA 0.5MG

A089211 001 Jun 14, 1988

1MG

A089212 001 Jun 14, 1988

2MG

A089213 001 Jun 14, 1988

COGENTIN

+ MERCK 0.5MG **

N009193 004

+ 1MG **

N009193 003

+ 2MG **

N009193 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENZYL ALCOHOL

LOTION; TOPICAL

ULESFIA

+ SHIONOGI INC

5% **

N022129 001 Apr 09, 2009

BENZYL BENZOATE

EMULSION; TOPICAL

BENZYL BENZOATE

LANNETT

50%

A084535 001

BEPRIDIL HYDROCHLORIDE

TABLET; ORAL

BEPADIN

MEDPOINTE PHARM HLC

200MG

N019001 001 Dec 28, 1990

300MG

N019001 002 Dec 28, 1990

400MG

N019001 003 Dec 28, 1990

VASCOR

JOHNSON AND JOHNSON

200MG

N019002 001 Dec 28, 1990

300MG

N019002 002 Dec 28, 1990

400MG

N019002 003 Dec 28, 1990

BETA CAROTENE

CAPSULE; ORAL

SOLATENE

ROCHE

30MG

N017589 001

BETAMETHASONE

CREAM; TOPICAL

CELESTONE

SCHERING

0.2%

N014762 001

SYRUP; ORAL

CELESTONE

MERCK SHARP DOHME

0.6MG/5ML

N014215 002

TABLET; ORAL

CELESTONE

SCHERING

0.6MG

N012657 003

BETAMETHASONE BENZOATE

CREAM; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N016998 002

GEL; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N017244 001

LOTION; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N017528 001

OINTMENT; TOPICAL

UTICORT

PARKE DAVIS

0.025%

N018089 001

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

ALPHATREX

SAVAGE LABS

EQ 0.05% BASE

N019138 001 Jun 26, 1984

BETAMETHASONE DIPROPIONATE

PERRIGO NEW YORK

EQ 0.05% BASE

A072536 001 Jan 31, 1990

EQ 0.05% BASE

A074579 001 Nov 26, 1997

PHARMADERM

EQ 0.05% BASE

N019136 001 Jun 26, 1984

TARO

EQ 0.05% BASE

A071143 001 Jun 17, 1987

TEVA

EQ 0.05% BASE

A071476 001 Aug 10, 1987

DIPROSONE

SCHERING

EQ 0.05% BASE

N017536 001

CREAM, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

ANDA REPOSITORY

EQ 0.05% BASE

A076603 001 Jan 23, 2004

DIPROLENE

SCHERING

EQ 0.05% BASE

N019408 001 Jan 31, 1986

DIPROLENE AF

+ ORGANON

EQ 0.05% BASE **

N019555 001 Apr 27, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAMETHASONE DIPROPIONATE

DISC;TOPICAL

DIPROSONE

SCHERING

EQ 0.1% BASE

N017829 001

GEL, AUGMENTED;TOPICAL

DIPROLENE

SCHERING

EQ 0.05% BASE

N019408 002 Nov 22, 1991

LOTION;TOPICAL

ALPHATREX

SAVAGE LABS

EQ 0.05% BASE

A070273 001 Aug 12, 1985

BETAMETHASONE DIPROPIONATE

ACTAVIS MID ATLANTIC

EQ 0.05% BASE

A070281 001 Jul 31, 1985

ALPHARMA US PHARMS

EQ 0.05% BASE

A071085 001 Feb 03, 1987

COSETTE

EQ 0.05% BASE

A071882 001 Jun 06, 1988

HIKMA

EQ 0.05% BASE

A209896 001 Feb 06, 2018

SHREE HARI INTL

EQ 0.05% BASE

A070274 001 Aug 12, 1985

TARO

EQ 0.05% BASE

A072276 001 Aug 24, 1988

EQ 0.05% BASE

A074272 001 Sep 30, 1994

DIPROSONE

+ SCHERING

EQ 0.05% BASE **

N017781 001

LOTION, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

VERTICE

EQ 0.05% BASE

A206389 001 Feb 13, 2018

DIPROLENE

+ ORGANON

EQ 0.05% BASE **

N019716 001 Aug 01, 1988

OINTMENT;TOPICAL

ALPHATREX

SAVAGE LABS

EQ 0.05% BASE

N019143 001 Sep 04, 1984

BETAMETHASONE DIPROPIONATE

PERRIGO NEW YORK

EQ 0.05% BASE

A072526 001 Jan 31, 1990

PHARMADERM

EQ 0.05% BASE

N019140 001 Sep 04, 1984

TEVA

EQ 0.05% BASE

A071477 001 Aug 10, 1987

DIPROSONE

SCHERING

EQ 0.05% BASE

N017691 001

OINTMENT, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

PAI HOLDINGS PHARM

EQ 0.05% BASE

A206118 001 Nov 09, 2017

SPRAY;TOPICAL

BETAMETHASONE DIPROPIONATE

TARO

EQ 0.05% BASE/SPRAY

A211722 001 Jun 17, 2020

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE

OINTMENT;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE

CHARTWELL RX

0.064%;0.005%

A201615 001 Jan 14, 2013

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM;TOPICAL

LOTRISONE

+ ORGANON

EQ 0.05% BASE;1%

N018827 001 Jul 10, 1984

LOTION;TOPICAL

LOTRISONE

+ MERCK SHARP DOHME

EQ 0.05% BASE;1% **

N020010 001 Dec 08, 2000

BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE SODIUM PHOSPHATE

WATSON LABS

EQ 3MG BASE/ML

A085738 001

CELESTONE

+ SCHERING

EQ 3MG BASE/ML **

N017561 001

BETAMETHASONE VALERATE

AEROSOL, FOAM;TOPICAL

BETAMETHASONE VALERATE

NOVAST LABS

0.12%

A207144 001 May 24, 2017

LUXIQ

+ MYLAN

0.12%

N020934 001 Feb 28, 1999

CREAM;TOPICAL

BETADERM

ROACO

EQ 0.1% BASE

N018839 001 Jun 30, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAMETHASONE VALERATE

CREAM; TOPICAL

BETAMETHASONE VALERATE

PERRIGO NEW YORK	EQ 0.1% BASE	A070053 001	Jun 10, 1986
PHARMADERM	EQ 0.1% BASE	N018860 002	Aug 31, 1983
PHARMAFAIR	EQ 0.1% BASE	A070485 001	May 29, 1987
TARO	EQ 0.1% BASE	A070062 001	May 14, 1985

BETATREX

SAVAGE LABS	EQ 0.1% BASE	N018862 001	Aug 31, 1983
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VALISONE

SCHERING	EQ 0.01% BASE	N016322 002	
	EQ 0.1% BASE	N016322 001	

LOTION; TOPICAL

BETA-VAL

COSETTE	EQ 0.1% BASE	A070072 001	Jun 27, 1985
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BETAMETHASONE VALERATE

PHARMADERM	EQ 0.1% BASE	N018870 001	Aug 31, 1983
PHARMAFAIR	EQ 0.1% BASE	A070484 001	May 29, 1987
TEVA PHARMS	EQ 0.1% BASE	A071883 001	Apr 22, 1988

BETATREX

SAVAGE LABS	EQ 0.1% BASE	N018867 001	Aug 31, 1983
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VALISONE

SCHERING	EQ 0.1% BASE	N016932 001	
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OINTMENT; TOPICAL

BETAMETHASONE VALERATE

PERRIGO NEW YORK	EQ 0.1% BASE	A071478 001	Dec 23, 1987
PHARMADERM	EQ 0.1% BASE	N018864 001	Aug 31, 1983
PHARMAFAIR	EQ 0.1% BASE	A070486 001	May 29, 1987

BETATREX

SAVAGE LABS	EQ 0.1% BASE	N018863 001	Aug 31, 1983
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VALISONE

SCHERING	EQ 0.1% BASE	N016740 001	
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BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

AKORN	EQ 0.5% BASE	A075386 001	Jun 30, 2000
APOTEX INC	EQ 0.5% BASE	A075446 001	Sep 28, 2000

TABLET; ORAL

KERLONE

SANOFI AVENTIS US	10MG **	N019507 001	Oct 27, 1989
	20MG **	N019507 002	Oct 27, 1989

BETAXOLOL HYDROCHLORIDE; CHLORTHALIDONE

TABLET; ORAL

KERLEDEX

SANOFI AVENTIS US	5MG;12.5MG	N019807 001	Oct 30, 1992
	10MG;12.5MG	N019807 002	Oct 30, 1992

BETAXOLOL HYDROCHLORIDE; PILOCARPINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC PILO

ALCON	EQ 0.25% BASE;1.75%	N020619 001	Apr 17, 1997
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BETAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

HISTALOG

LILLY	50MG/ML	N009344 001	
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BETHANECHOL CHLORIDE

INJECTABLE; INJECTION

URECHOLINE

+ ODYSSEY PHARMS	5MG/ML **	N006536 001	
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TABLET; ORAL

BETHANECHOL CHLORIDE

ABLE	5MG	A040492 001	Jul 27, 2004
	10MG	A040483 001	Jul 27, 2004
	25MG	A040485 001	Jul 27, 2004
	50MG	A040509 001	Jul 27, 2004
ACTAVIS ELIZABETH	5MG	A040552 001	Oct 28, 2004
	10MG	A040553 001	Oct 28, 2004
	25MG	A040554 001	Oct 28, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

	50MG	A040551 001	Oct 28, 2004
ASCOT	10MG	A088288 001	Jun 08, 1983
	25MG	A088289 001	Jun 08, 1983
HERITAGE PHARMA	5MG	A091256 001	May 04, 2010
	10MG	A091256 002	May 04, 2010
	25MG	A091256 003	May 04, 2010
	50MG	A091256 004	May 04, 2010
IMPAX LABS	5MG	A040721 001	Nov 01, 2006
	10MG	A040721 002	Nov 01, 2006
	25MG	A040721 003	Nov 01, 2006
	50MG	A040721 004	Nov 01, 2006
IVAX SUB TEVA PHARMS	25MG	A084689 001	
LANNETT	5MG	A084702 001	
	10MG	A084712 001	
	25MG	A084074 001	
LANNETT CO INC	5MG	A040677 002	Mar 27, 2008
	10MG	A040677 003	Mar 27, 2008
	25MG	A040677 004	Mar 27, 2008
	50MG	A040677 001	Mar 27, 2008
SANDOZ	5MG	A084353 001	
	10MG	A084378 001	
	10MG	A084379 001	
	25MG	A084383 001	
	25MG	A084384 001	
SUN PHARM INDS INC	5MG	A040897 001	Apr 22, 2009
	10MG	A040897 002	Apr 22, 2009
	25MG	A040897 003	Apr 22, 2009
	50MG	A040897 004	Apr 22, 2009
WATSON LABS	5MG	A084402 001	
	5MG	A085230 002	
	5MG	A085841 001	
	10MG	A084408 001	
	10MG	A085228 001	
	10MG	A085842 001	
	25MG	A084441 001	
	25MG	A085229 001	
	25MG	A085839 001	
	50MG	A087397 001	
	50MG	A087444 001	
WOCKHARDT	5MG	A040532 001	Sep 29, 2003
	10MG	A040533 001	Sep 29, 2003
	25MG	A040534 001	Sep 29, 2003
	50MG	A040518 001	Sep 29, 2003
MYOTONACHOL			
GLENWOOD	5MG	A084188 001	
	10MG	A084188 003	
	25MG	A084188 004	
URECHOLINE			
ODYSSEY PHARMS	5MG	A089095 001	Dec 19, 1985
+	5MG **	N006536 003	
+	10MG	A088440 001	May 29, 1984
+	10MG **	N006536 002	
+	25MG	A088441 001	May 29, 1984
+	25MG **	N006536 004	
	50MG	A089096 001	Dec 19, 1985
+	50MG **	N006536 005	

BETHANIDINE SULFATE

TABLET; ORAL

TENATHAN

ROBINS AH	10MG	N017675 001	
	25MG	N017675 002	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETRIXABAN

CAPSULE; ORAL

BEVYXXA

+	PORTOLA PHARMS INC	40MG	N208383	001	Jun 23, 2017
+		80MG	N208383	002	Jun 23, 2017

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

	FRESENIUS KABI USA	50MG	A079045	001	May 13, 2010
	KUDCO IRELAND	50MG	A077995	001	Jul 06, 2009
	MYLAN	50MG	A079185	001	Jul 06, 2009
	ROXANE	50MG	A078285	001	Mar 24, 2011
	SYNTHON PHARMS	50MG	A077973	001	Jul 06, 2009
	TEVA	50MG	A076932	001	Jul 06, 2009

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC

BIMATOPROST

	HIKMA	0.03%	A203299	001	Nov 08, 2018
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LUMIGAN

+	ABBVIE	0.03% **	N021275	001	Mar 16, 2001
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BIPERIDEN HYDROCHLORIDE

TABLET; ORAL

AKINETON

	ABBVIE	2MG	N012003	001	
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BIPERIDEN LACTATE

INJECTABLE; INJECTION

AKINETON

	ABBVIE	5MG/ML	N012418	002	
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BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION, TABLET, DELAYED RELEASE; ORAL

HALFLYTELY

+	BRAINTREE	5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM ; N/A, 5.6GM **	N021551	003	Jul 16, 2010
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PEG-3350, SODIUM CHLORIDE, SODIUM BICARBONATE, POTASSIUM CHLORIDE AND BISACODYL

	NOVEL LABS INC	5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM ; N/A, 5.6GM	A202217	001	Aug 20, 2014
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BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE; ORAL

HELIDAC

+	CASPER PHARMA LLC	262.4MG, N/A, N/A; N/A, 250MG, N/A; N/A, N/A, 5 00MG **	N050719	001	Aug 15, 1996
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BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

	DASH PHARMS NATCO	5MG	A075831	001	Dec 14, 2005
		10MG	A075831	002	Dec 14, 2005
	TEVA PHARMS	5MG	A075644	001	Jun 26, 2001
		10MG	A075644	002	Jun 26, 2001

ZEBETA

+	TEVA WOMENS	5MG **	N019982	002	Jul 31, 1992
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+		10MG **	N019982	001	Jul 31, 1992
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BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

	ACTAVIS ELIZABETH	2.5MG; 6.25MG	A075672	001	Sep 25, 2000
		5MG; 6.25MG	A075672	002	Sep 25, 2000
		10MG; 6.25MG	A075672	003	Sep 25, 2000
	APOTHECON	2.5MG; 6.25MG	A075642	002	Dec 27, 2000
		5MG; 6.25MG	A075642	001	Dec 27, 2000
		10MG; 6.25MG	A075642	003	Dec 27, 2000
	CHARTWELL RX	2.5MG; 6.25MG	A075527	001	Sep 25, 2000
		5MG; 6.25MG	A075527	003	Sep 25, 2000
		10MG; 6.25MG	A075527	002	Sep 25, 2000
	IVAX SUB TEVA PHARMS	2.5MG; 6.25MG	A075632	001	Sep 27, 2000
		5MG; 6.25MG	A075632	002	Sep 27, 2000
		10MG; 6.25MG	A075632	003	Sep 27, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

TEVA	2.5MG; 6.25MG	A075686 001	Jan 19, 2001
	5MG; 6.25MG	A075686 002	Jan 19, 2001
	10MG; 6.25MG	A075686 003	Jan 19, 2001
WATSON LABS TEVA	2.5MG; 6.25MG	A075469 001	Sep 25, 2000
	5MG; 6.25MG	A075469 002	Sep 25, 2000
	10MG; 6.25MG	A075469 003	Sep 25, 2000

BITOLTEROL MESYLATE

AEROSOL, METERED; INHALATION

TORNALATE

SANOFI AVENTIS US	0.37MG/INH	N018770 001	Dec 28, 1984
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SOLUTION; INHALATION

TORNALATE

SANOFI AVENTIS US	0.2%	N019548 001	Feb 19, 1992
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BIVALIRUDIN

INJECTABLE; INTRAVENOUS

BIVALIRUDIN

APOTEX	250MG/VIAL	A204876 001	Jul 06, 2017
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SOLUTION; INTRAVENOUS

BIVALIRUDIN IN 0.9% SODIUM CHLORIDE

+ BAXTER HLTHCARE CORP	250MG/50ML (5MG/ML)	N208374 001	Dec 21, 2017
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+	500MG/100ML (5MG/ML)	N208374 002	Dec 21, 2017
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BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLENOXANE

+ BRISTOL MYERS SQUIBB	EQ 15 UNITS BASE/VIAL **	N050443 001	
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+	EQ 30 UNITS BASE/VIAL **	N050443 002	Sep 07, 1995
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BLEOMYCIN SULFATE

CIPLA	EQ 15 UNITS BASE/VIAL	A209439 001	Mar 11, 2019
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PHARMACHEMIE BV	EQ 15 UNITS BASE/VIAL	A065201 001	Dec 13, 2007
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TEVA PARENTERAL	EQ 15 UNITS BASE/VIAL	A064084 001	Jun 01, 1996
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	EQ 30 UNITS BASE/VIAL	A064084 002	Jun 01, 1996
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BOCEPREVIR

CAPSULE; ORAL

VICTRELIS

MERCK SHARP DOHME	200MG	N202258 001	May 13, 2011
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BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

ACCORD HLTHCARE	3.5MG/VIAL	A204405 001	Jul 26, 2022
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MYLAN LABS LTD	3.5MG/VIAL	A205160 001	Oct 31, 2022
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SCINOPHARM TAIWAN	3.5MG/VIAL	A216912 001	Sep 26, 2023
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TEVA PHARMS USA	3.5MG/VIAL	A205857 001	May 02, 2022
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POWDER; INTRAVENOUS

BORTEZOMIB

DR REDDYS	3.5MG/VIAL	N206927 001	Oct 04, 2019
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FRESENIUS KABI USA	3.5MG/VIAL	N205004 001	Nov 06, 2017
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SOLUTION; INTRAVENOUS

BORTEZOMIB

+ ACCORD HLTHCARE	2.5MG/ML (2.5MG/ML) **	N215441 001	Jul 26, 2022
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+	3.5MG/1.4ML (2.5MG/ML) **	N215441 002	Jul 26, 2022
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+ MAIA PHARMS INC	3.5MG/3.5ML (1MG/ML)	N215331 001	Jul 27, 2022
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+	3.5MG/1.4ML (2.5MG/ML)	N215331 002	Jul 27, 2022
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BOSENTAN

TABLET; ORAL

BOSENTAN

ALEMBIC	62.5MG	A211461 001	Jan 23, 2020
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	125MG	A211461 002	Jan 23, 2020
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ALVOGEN PINE BROOK	62.5MG	A206002 001	Apr 26, 2019
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	125MG	A206002 002	Apr 26, 2019
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AMNEAL PHARMS CO	62.5MG	A209742 001	Apr 26, 2019
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	125MG	A209742 002	Apr 26, 2019
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CHARTWELL MOLECULAR	62.5MG	A210342 001	Jan 03, 2020
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	125MG	A210342 002	Jan 03, 2020
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BOSENTAN

TABLET; ORAL

BOSENTAN

HIKMA	62.5MG	A208695 001	Apr 26, 2019
	125MG	A208695 002	Apr 26, 2019
MYLAN	62.5MG	A205173 001	Jan 15, 2020
	125MG	A205173 002	Jan 15, 2020
NATCO PHARMA LTD	62.5MG	A206987 001	Apr 26, 2019
	125MG	A206987 002	Apr 26, 2019
PAR PHARM INC	62.5MG	A205699 001	Apr 26, 2019
	125MG	A205699 002	Apr 26, 2019

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

ABRAXIS PHARM	50MG/ML	A070134 001	Apr 29, 1986
	100MG/ML	A071298 001	Feb 13, 1987
ASTRAZENECA	50MG/ML	A071151 001	Aug 10, 1987
	50MG/ML	A071152 001	Aug 10, 1987
	50MG/ML	A071153 001	Aug 10, 1987
BRECKENRIDGE	50MG/ML	A204386 001	Dec 21, 2018
HIKMA	50MG/ML	A070545 001	May 14, 1986
	50MG/ML	A070546 001	May 14, 1986
+ HOSPIRA	50MG/ML **	N019030 001	Apr 29, 1986
	50MG/ML	N019033 001	Apr 29, 1986
INTL MEDICATION	50MG/ML	A070119 001	Apr 29, 1986
LUITPOLD	50MG/ML	A070891 001	Jul 26, 1988
BRETYLIUM TOSYLATE IN DEXTROSE 5%			
ABBOTT	200MG/100ML	N019005 002	Apr 29, 1986
	400MG/100ML	N019005 003	Apr 29, 1986
	800MG/100ML	N019005 001	Apr 29, 1986
BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	100MG/100ML	N019121 001	Apr 29, 1986
	200MG/100ML	N019121 002	Apr 29, 1986
	400MG/100ML	N019121 003	Apr 29, 1986
BAXTER HLTHCARE	200MG/100ML	N019837 002	Apr 12, 1989
	400MG/100ML	N019837 001	Apr 12, 1989
HOSPIRA INC	200MG/100ML	N019008 002	Apr 29, 1986
	400MG/100ML	N019008 003	Apr 29, 1986
	800MG/100ML	N019008 001	Apr 29, 1986
BRETYLOL			
HOSPIRA	50MG/ML	N017954 001	

BREXPIPIRAZOLE

TABLET; ORAL

BREXPIPIRAZOLE

AJANTA PHARMA LTD	0.25MG	A213718 001	Feb 03, 2023
	0.5MG	A213718 002	Feb 03, 2023
	1MG	A213718 003	Feb 03, 2023
	2MG	A213718 004	Feb 03, 2023
	3MG	A213718 005	Feb 03, 2023
	4MG	A213718 006	Feb 03, 2023
AMNEAL	0.25MG	A213562 001	Jan 31, 2023
	0.5MG	A213562 002	Jan 31, 2023
	1MG	A213562 003	Jan 31, 2023
	2MG	A213562 004	Jan 31, 2023
	3MG	A213562 005	Jan 31, 2023
	4MG	A213562 006	Jan 31, 2023
AUROBINDO PHARMA LTD	0.25MG	A213659 001	Nov 02, 2023
	0.5MG	A213659 002	Nov 02, 2023
	1MG	A213659 003	Nov 02, 2023
	2MG	A213659 004	Nov 02, 2023
	3MG	A213659 005	Nov 02, 2023
	4MG	A213659 006	Nov 02, 2023
LUPIN LTD	0.25MG	A213512 001	Mar 17, 2023
	0.5MG	A213512 002	Mar 17, 2023
	1MG	A213512 003	Mar 17, 2023
	2MG	A213512 004	Mar 17, 2023
	3MG	A213512 005	Mar 17, 2023
	4MG	A213512 006	Mar 17, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BREXPIRAZOLE

TABLET; ORAL

BREXPIRAZOLE

SANDOZ	0.25MG	A213570 001	Sep 26, 2022
	0.5MG	A213570 002	Sep 26, 2022
	1MG	A213570 003	Sep 26, 2022
	2MG	A213570 004	Sep 26, 2022
	3MG	A213570 005	Sep 26, 2022
	4MG	A213570 006	Sep 26, 2022
TEVA PHARMS USA INC	0.25MG	A213692 001	Aug 11, 2022
	0.5MG	A213692 002	Aug 11, 2022
	1MG	A213692 003	Aug 11, 2022
	2MG	A213692 004	Aug 11, 2022
	3MG	A213692 005	Aug 11, 2022
	4MG	A213692 006	Aug 11, 2022
ZYDUS PHARMS	0.25MG	A213660 001	Jan 10, 2023
	0.5MG	A213660 002	Jan 10, 2023
	1MG	A213660 003	Jan 10, 2023
	2MG	A213660 004	Jan 10, 2023
	3MG	A213660 005	Jan 10, 2023
	4MG	A213660 006	Jan 10, 2023

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN

+ ALLERGAN	0.2% **	N020613 001	Sep 06, 1996
	0.5%	N020490 001	Mar 13, 1997
BRIMONIDINE TARTRATE			
TEVA PARENTERAL	0.2%	A076372 001	Sep 10, 2004

BRIVARACETAM

TABLET; ORAL

BRIVARACETAM

AUROBINDO PHARMA LTD	50MG	A214848 001	Jan 06, 2023
	100MG	A214848 002	Jan 06, 2023
LUPIN LTD	10MG	A214918 001	Dec 20, 2022
	25MG	A214918 002	Dec 20, 2022
	50MG	A214918 003	Dec 20, 2022
	75MG	A214918 004	Dec 20, 2022
	100MG	A214918 005	Dec 20, 2022
SUNSHINE	10MG	A214748 001	Jun 09, 2022
	25MG	A214748 002	Jun 09, 2022
	50MG	A214748 003	Jun 09, 2022
	75MG	A214748 004	Jun 09, 2022
	100MG	A214748 005	Jun 09, 2022
ZYDUS PHARMS	10MG	A214501 001	Oct 03, 2022
	25MG	A214501 002	Oct 03, 2022
	50MG	A214501 003	Oct 03, 2022
	75MG	A214501 004	Oct 03, 2022
	100MG	A214501 005	Oct 03, 2022

BROMFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC

BROMDAY

+ BAUSCH AND LOMB INC	EQ 0.09% ACID **	N021664 002	Oct 16, 2010
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BROMFENAC SODIUM

AMRING PHARMS	EQ 0.09% ACID	A202030 001	Jan 09, 2013
APOTEX	EQ 0.09% ACID	A202435 001	Jun 19, 2014
	EQ 0.09% ACID	A202620 001	Jun 23, 2014
CHARTWELL RX	EQ 0.09% ACID	A201941 001	Feb 10, 2015
COASTAL PHARMS	EQ 0.09% ACID	A201211 001	May 11, 2011
RISING	EQ 0.09% ACID	A203368 001	Jun 03, 2019
XIBROM			
+ BAUSCH AND LOMB INC	EQ 0.09% ACID **	N021664 001	Mar 24, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BROMOCRIPTINE MESYLATE

CAPSULE; ORAL

BROMOCRIPTINE MESYLATE

LEK PHARM

EQ 5MG BASE

A075100 001 Dec 10, 1998

TABLET; ORAL

BROMOCRIPTINE MESYLATE

AUROBINDO PHARMA USA EQ 2.5MG BASE

A076962 001 Sep 24, 2004

BROMODIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

AMBODRYL

PARKE DAVIS

25MG

N007984 001

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL

AMBENYL

FOREST LABS

12.5MG/5ML; 10MG/5ML

N009319 006 Jan 10, 1984

BROMANYL

ALPHARMA US PHARMS

12.5MG/5ML; 10MG/5ML

A088343 001 Aug 15, 1984

BROMODIPHENHYDRAMINE HYDROCHLORIDE AND CODEINE PHOSPHATE

WOCKHARDT

12.5MG/5ML; 10MG/5ML

A088626 001 Oct 12, 1984

BROMPHENIRAMINE MALEATE

ELIXIR; ORAL

BROMPHENIRAMINE MALEATE

ALPHARMA US PHARMS

2MG/5ML

A086936 001

KV PHARM

2MG/5ML

A085466 001

PHARM ASSOC

2MG/5ML

A087517 001

USL PHARMA

2MG/5ML

A087964 001 Jan 25, 1983

INJECTABLE; INJECTION

BROMPHENIRAMINE MALEATE

WATSON LABS

10MG/ML

A083821 001

100MG/ML

A083820 001

DIMETANE-TEN

WYETH AYERST

10MG/ML

N011418 002

TABLET; ORAL

BROMPHENIRAMINE MALEATE

BARR

4MG

A084468 001

IVAX SUB TEVA PHARMS

4MG

A084351 001

NEWTRON PHARMS

4MG

A086987 001

NEXGEN PHARMA INC

4MG

A086187 001

PAR PHARM

4MG

A087009 001

PIONEER PHARMS

4MG

A088604 001 Jul 13, 1984

UPSHER SMITH LABS

4MG

A083215 001

VITARINE

4MG

A085850 001

WATSON LABS

4MG

A083123 001

4MG

A085769 001

DIMETANE

WYETH CONS

4MG

N010799 003

TABLET, EXTENDED RELEASE; ORAL

DIMETANE

WYETH CONS

8MG

N010799 010 Jun 10, 1983

12MG

N010799 011 Jun 10, 1983

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMANATE DM

ALPHARMA US PHARMS

2MG/5ML; 10MG/5ML; 30MG/5ML

A088722 001 Mar 07, 1985

BROMFED-DM

WOCKHARDT

2MG/5ML; 10MG/5ML; 30MG/5ML

A089681 001 Dec 22, 1988

BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

BIONPHARMA

2MG/5ML; 10MG/5ML; 30MG/5ML

A203997 001 Sep 30, 2020

RHODES PHARMS

2MG/5ML; 10MG/5ML; 30MG/5ML

A202955 001 Nov 05, 2020

DIMETANE-DX

+

ROBINS AH

2MG/5ML; 10MG/5ML; 30MG/5ML **

N019279 001 Aug 24, 1984

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE

ALZA 16MG;240MG

N019672 001 Mar 29, 1996

BUCLIZINE HYDROCHLORIDE

TABLET;ORAL

BUCLADIN-S

STUART PHARMS 50MG

N010911 006

BUDESONIDE

AEROSOL, METERED;NASAL

RHINOCORT

ASTRAZENECA 0.032MG/INH

N020233 001 Feb 14, 1994

CAPSULE, DELAYED RELEASE;ORAL

BUDESONIDE

BARR LABS DIV TEVA 3MG

A090379 001 Apr 02, 2014

NATCO 3MG

A206724 001 Nov 23, 2016

SCIECURE PHARMA INC 3MG

A209041 001 Sep 28, 2017

ORTIKOS

+ SUN PHARM INDS INC 6MG

N211929 001 Jun 13, 2019

+ 9MG

N211929 002 Jun 13, 2019

POWDER, METERED;INHALATION

PULMICORT

ASTRAZENECA 0.16MG/INH

N020441 002 Jun 24, 1997

0.32MG/INH

N020441 003 Jun 24, 1997

BUMETANIDE

INJECTABLE;INJECTION

BUMETANIDE

HOSPIRA 0.25MG/ML

A074160 001 Oct 30, 1997

TEVA PARENTERAL 0.25MG/ML

A074613 001 Nov 18, 1997

BUMEX

+ VALIDUS PHARMS 0.25MG/ML **

N018226 001 Feb 28, 1983

BUPIVACAINE HYDROCHLORIDE

INJECTABLE;INJECTION

BUPIVACAINE HYDROCHLORIDE

CIVICA 0.5%

A211096 001 Feb 19, 2019

HOSPIRA 0.25%

A070586 001 Mar 03, 1987

0.25%

N018053 002

0.5%

N018053 001

0.75%

A070587 001 Mar 03, 1987

0.75%

N018053 003

BUPIVACAINE HYDROCHLORIDE KIT

HOSPIRA 0.075%

N019978 001 Sep 03, 1992

0.114%

N019978 002 Sep 03, 1992

0.23%

N019978 003 Sep 03, 1992

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

INTL MEDICATED 0.25%

A076012 001 Jan 09, 2002

0.5%

A076012 002 Jan 09, 2002

0.75%

A076012 003 Jan 09, 2002

INJECTABLE;SPINAL

BUPIVACAINE HYDROCHLORIDE

BAXTER HLTHCARE CORP 0.75%

A207266 001 Jul 25, 2016

SENSORCAINE

FRESENIUS KABI USA 0.75%

A071202 001 Apr 15, 1987

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE;INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

HOSPIRA 0.25%;0.005MG/ML

A071166 001 Jun 16, 1988

0.5%;0.005MG/ML

A071169 001 Jun 16, 1988

0.75%;0.005MG/ML

A071171 001 Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE;INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

+ HOSPIRA 0.5%;0.0091MG/ML

N022046 001 Jul 13, 1983

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPIVACAINE HYDROCHLORIDE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DUOCAINE

AMPHASTAR PHARMS INC	EQ 0.375% (37.5MG/10ML); EQ 1% (100MG/10ML)	N021496 001	May 23, 2003
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BUPIVACAINE; MELOXICAM

SOLUTION, EXTENDED RELEASE; PERIARTICULAR

ZYNRELEF KIT

+	HERON THERAPS INC	60MG/2.3ML (29.25MG/ML); 1.8MG/2.3ML (0.88MG/ML)	N211988 001	May 12, 2021
+		300MG/10.5ML (29.25MG/ML); 9MG/10.5ML (0.88MG/ML)	N211988 003	May 12, 2021

BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

ALVOGEN	7.5MCG/HR	A207490 005	Mar 21, 2023
MYLAN TECH VIATRIS	5MCG/HR	A210162 001	May 03, 2021
	7.5MCG/HR	A210162 002	May 03, 2021
	10MCG/HR	A210162 003	May 03, 2021
	15MCG/HR	A210162 004	May 03, 2021
	20MCG/HR	A210162 005	May 03, 2021

BUPRENORPHINE HYDROCHLORIDE

FILM; BUCCAL

BUPRENORPHINE HYDROCHLORIDE

ALVOGEN	EQ 0.075MG BASE	A211594 001	Aug 03, 2021
	EQ 0.15MG BASE	A211594 002	Aug 03, 2021
	EQ 0.3MG BASE	A211594 003	Aug 03, 2021
	EQ 0.45MG BASE	A211594 004	Aug 03, 2021
	EQ 0.6MG BASE	A211594 005	Aug 03, 2021
	EQ 0.75MG BASE	A211594 006	Aug 03, 2021
	EQ 0.9MG BASE	A211594 007	Aug 03, 2021

IMPLANT; IMPLANTATION

PROBUPHINE

+	TITAN PHARMS	EQ 80MG BASE/IMPLANT	N204442 001	May 26, 2016
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INJECTABLE; INJECTION

BUPRENEX

+	INDIVIOR	EQ 0.3MG BASE/ML **	N018401 001	
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TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

BARR	EQ 2MG BASE	A090360 001	May 07, 2010
	EQ 8MG BASE	A090360 002	May 07, 2010
MYLAN	EQ 2MG BASE	A201066 001	Mar 06, 2015
	EQ 8MG BASE	A201066 002	Mar 06, 2015

SUBUTEX

+	INDIVIOR	EQ 2MG BASE **	N020732 002	Oct 08, 2002
+		EQ 8MG BASE **	N020732 003	Oct 08, 2002

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM; BUCCAL

BUNAVAIL

+	BDSI	EQ 2.1MG BASE; EQ 0.3MG BASE	N205637 001	Jun 06, 2014
+		EQ 4.2MG BASE; EQ 0.7MG BASE	N205637 002	Jun 06, 2014
+		EQ 6.3MG BASE; EQ 1MG BASE	N205637 003	Jun 06, 2014

FILM; SUBLINGUAL

CASSIPA

+	TEVA PHARMS USA	EQ 16MG BASE; EQ 4MG BASE	N208042 001	Sep 07, 2018
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TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

DR REDDYS LABS SA	EQ 2MG BASE; EQ 0.5MG BASE	A206953 001	Jul 17, 2020
	EQ 8MG BASE; EQ 2MG BASE	A206953 002	Jul 17, 2020
TEVA PHARMS USA	EQ 2MG BASE; EQ 0.5MG BASE	A091149 001	Sep 08, 2014
	EQ 8MG BASE; EQ 2MG BASE	A091149 002	Sep 08, 2014

SUBOXONE

+	INDIVIOR	EQ 2MG BASE; EQ 0.5MG BASE **	N020733 001	Oct 08, 2002
+		EQ 8MG BASE; EQ 2MG BASE **	N020733 002	Oct 08, 2002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPROPION HYDROCHLORIDE

TABLET; ORAL

BUPROPION HYDROCHLORIDE

AUROBINDO PHARMA USA	75MG	A075491 001	Apr 17, 2000
	100MG	A075491 002	Apr 17, 2000
HERITAGE PHARMA	75MG	A075310 001	Nov 29, 1999
	100MG	A075310 002	Nov 29, 1999
INVATECH	75MG	A075613 002	Oct 10, 2000
	100MG	A075613 001	Oct 10, 2000

WELLBUTRIN

+	GLAXOSMITHKLINE	50MG **	N018644 001	Dec 30, 1985
+		75MG **	N018644 002	Dec 30, 1985
+		100MG **	N018644 003	Dec 30, 1985

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

ACTAVIS LABS FL INC	300MG	A077715 002	Jun 13, 2007
ANCHEN PHARMS	150MG	A077284 001	Dec 14, 2006
	300MG	A077284 002	Dec 14, 2006
AUROBINDO PHARMA USA	100MG	A090325 001	Apr 08, 2010
	150MG	A090325 002	Apr 08, 2010
	150MG	A090942 001	Jul 14, 2010
	200MG	A090325 003	Apr 08, 2010
	300MG	A090942 002	Jul 14, 2010
IMPAX LABS	150MG	A077415 001	Nov 26, 2008
	200MG	A076711 001	Dec 03, 2004
	300MG	A077415 002	Dec 15, 2006
JUBILANT GENERICS	100MG	A202774 001	Oct 11, 2013
	150MG	A202774 002	Oct 11, 2013
	150MG	A202775 001	Oct 11, 2013
	150MG	A207459 001	Jun 30, 2017
	200MG	A202774 003	Oct 11, 2013
	300MG	A207459 002	Jun 30, 2017
MYLAN	150MG	A090941 001	May 03, 2010
SANDOZ	100MG	A076845 001	Jul 14, 2005
	150MG	A076834 001	Jul 14, 2005
	150MG	A076845 002	Jul 14, 2005
SUN PHARM	100MG	A078866 001	Apr 06, 2010
	150MG	A200695 001	Dec 18, 2014
TORRENT	100MG	A203969 001	Oct 31, 2014
	150MG	A203969 002	Oct 31, 2014
	200MG	A203969 003	Oct 31, 2014
WATSON LABS INC	100MG	A077455 001	Jul 19, 2010
	150MG	A077455 002	Mar 12, 2008
	200MG	A077455 003	Jul 19, 2010
WOCKHARDT LTD	100MG	A201331 001	Aug 30, 2012
	150MG	A201331 002	Aug 30, 2012
	200MG	A201331 003	Aug 30, 2012

WELLBUTRIN SR

GLAXOSMITHKLINE	50MG	N020358 001	Oct 04, 1996
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ZYBAN

GLAXOSMITHKLINE	100MG	N020711 002	May 14, 1997
+	150MG **	N020711 003	May 14, 1997

BUSPIRONE HYDROCHLORIDE

CAPSULE; ORAL

BUSPAR

+	BRISTOL MYERS SQUIBB	5MG **	N021190 001	Dec 20, 2000
+		7.5MG **	N021190 002	Dec 20, 2000
+		10MG **	N021190 003	Dec 20, 2000
+		15MG **	N021190 004	Dec 20, 2000

TABLET; ORAL

BUSPAR

+	BRISTOL MYERS SQUIBB	5MG **	N018731 001	Sep 29, 1986
+		10MG **	N018731 002	Sep 29, 1986
+		15MG **	N018731 003	Apr 22, 1996
+		30MG **	N018731 004	Apr 22, 1996

BUSPIRONE HYDROCHLORIDE

EGIS	5MG	A075119 001	Mar 14, 2002
	10MG	A075119 002	Mar 14, 2002
	15MG	A075119 003	Jan 23, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

IVAX SUB TEVA PHARMS	5MG **	A075385 001	Mar 01, 2002
	10MG **	A075385 002	Mar 01, 2002
	15MG **	A075385 003	Mar 01, 2002
MYLAN	5MG	A075467 001	Feb 28, 2002
	7.5MG	A075467 002	Mar 28, 2001
	10MG	A075467 003	Feb 28, 2002
	15MG	A075467 004	Feb 28, 2002
NESHER PHARMS	5MG	A075572 001	Feb 27, 2002
	10MG	A075572 002	Feb 27, 2002
	15MG	A075572 003	Feb 27, 2002
RISING	5MG	A075413 001	Mar 19, 2002
	10MG	A075413 002	Mar 19, 2002
	15MG	A075413 003	Mar 19, 2002

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

ACTAVIS	6MG/ML	A205139 001	Dec 08, 2017
AM REGENT	6MG/ML	A202259 001	Dec 22, 2015
ARTHUR GRP	6MG/ML	A205106 001	Sep 21, 2018
MYLAN LABS LTD	6MG/ML	A205184 001	Jul 13, 2018

BUTABARBITAL SODIUM

CAPSULE; ORAL

BUTICAPS

MEDPOINTE PHARM HLC	15MG	A085381 001	
	30MG	A085381 002	
	50MG	A085381 003	
	100MG	A085381 004	

ELIXIR; ORAL

BUTABARB

ALPHARMA US PHARMS	30MG/5ML	A085873 001	
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BUTABARBITAL SODIUM

WOCKHARDT	30MG/5ML	A085383 001	
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BUTALAN

LANNETT	33.3MG/5ML	A085880 001	
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BUTISOL SODIUM

MEDA PHARMS	30MG/5ML	A085380 001	
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SARISOL

HALSEY	30MG/5ML	A084723 001	
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TABLET; ORAL

BUTABARBITAL

BUNDY	30MG	A085550 001	
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BUTABARBITAL SODIUM

SANDOZ	15MG	A084292 003	Feb 09, 1982
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+

	15MG	A085938 001	
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	30MG	A084272 002	
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	30MG	A085934 001	
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SOLVAY	16.2MG	A083606 001	
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	32.4MG	A083898 001	
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	48.6MG	A083897 001	
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	97.2MG	A083896 001	
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TEVA	15MG	A088632 001	May 18, 1985
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	30MG	A088631 001	May 01, 1985
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WATSON LABS	15MG	A085764 001	
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	30MG	A085772 001	
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WHITEWORTH TOWN PLSN	15MG	A083325 002	
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	30MG	A083337 001	
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BUTISOL SODIUM

MYLAN SPECIALITY LP	15MG **	N000793 002	
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+

	30MG	N000793 004	
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	50MG **	N000793 003	
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	100MG **	N000793 005	
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SARISOL NO. 1

HALSEY	15MG	A084719 001	
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SARISOL NO. 2

HALSEY	30MG	A084719 002	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUTABARBITAL SODIUM

TABLET; ORAL

SODIUM BUTABARBITAL

HIKMA	15MG	A085418	001
	30MG	A085432	001
IVAX SUB TEVA PHARMS	15MG	A083484	001
	30MG	A084040	001
LANNETT	15MG	A085849	001
	30MG	A085866	001
	100MG	A085881	001
MARSHALL PHARMA	16.2MG	A083524	001
	32.4MG	A083858	001

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

MENTAX

+ MYLAN	1%	N020524	001	Oct 18, 1996
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MENTAX-TC

MYLAN	1%	N021408	001	Oct 17, 2002
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BUTOCONAZOLE NITRATE

CREAM; VAGINAL

BUTOCONAZOLE NITRATE

PADAGIS US	2%	N019881	001	Feb 07, 1997
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FEMSTAT

ROCHE PALO	2%	N019215	001	Nov 25, 1985
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FEMSTAT 3

+ BAYER	2%	N020421	001	Dec 21, 1995
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SUPPOSITORY; VAGINAL

FEMSTAT

ROCHE PALO	100MG	N019359	001	Nov 25, 1985
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BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

BAXTER HLTHCARE CORP	2MG/ML	A075697	001	Oct 23, 2001
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HIKMA FARMACEUTICA	2MG/ML	A078247	001	Apr 29, 2009
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HOSPIRA	1MG/ML	A075342	001	Nov 04, 1999
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	1MG/ML	A075559	001	Mar 20, 2000
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	2MG/ML	A075342	002	Nov 04, 1999
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	2MG/ML	A075559	002	Mar 20, 2000
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BUTORPHANOL TARTRATE PRESERVATIVE FREE

BAXTER HLTHCARE CORP	1MG/ML	A075695	001	Oct 23, 2001
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	2MG/ML	A075695	002	Oct 23, 2001
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HOSPIRA	1MG/ML	A074620	001	Jan 22, 1997
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	1MG/ML	A075170	001	Sep 28, 1998
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	2MG/ML	A074620	002	Jan 22, 1997
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	2MG/ML	A075170	002	Sep 28, 1998
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STADOL

+ APOTHECON	2MG/ML **	N017857	004
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STADOL PRESERVATIVE FREE

+ APOTHECON	1MG/ML **	N017857	001
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+	2MG/ML **	N017857	002
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SPRAY, METERED; NASAL

STADOL

BRISTOL MYERS SQUIBB	1MG/SPRAY **	N019890	001	Dec 12, 1991
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CABAZITAXEL

SOLUTION; INTRAVENOUS

CABAZITAXEL

APOTEX	60MG/1.5ML (40MG/ML)	A207736	001	Feb 10, 2023
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BRECKENRIDGE	60MG/1.5ML (40MG/ML)	A207619	001	Jun 23, 2022
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MYLAN LABS LTD	60MG/1.5ML (40MG/ML)	A207381	001	Jul 05, 2023
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CABERGOLINE

TABLET; ORAL

CABERGOLINE

ACTAVIS LABS FL INC	0.5MG	A078035	001	Apr 21, 2008
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APOTEX CORP	0.5MG	A201503	001	Mar 08, 2013
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IMPAX LABS INC	0.5MG	A077843	001	Jul 03, 2007
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MYLAN	0.5MG	A202947	001	Dec 02, 2013
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CABERGOLINE

TABLET; ORAL

DOSTINEX

+ PFIZER

0.5MG **

N020664 001 Dec 23, 1996

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAFFEINE CITRATE

SUN PHARM

EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A090077 001 Sep 30, 2009

SOLUTION; ORAL

CAFECIT

+ HIKMA

EQ 30MG BASE/3ML (EQ 10MG BASE/ML) **

N020793 002 Apr 12, 2000

CAFFEINE CITRATE

AM REGENT

EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A090064 001 Nov 20, 2009

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

CAFERGOT

+ NOVARTIS

100MG; 2MG **

N009000 002

TABLET; ORAL

CAFERGOT

NOVARTIS

100MG; 1MG

N006620 001

+ SANDOZ

100MG; 1MG

A084294 001

ERGOTAMINE TARTRATE AND CAFFEINE

HIKMA INTL PHARMS

100MG; 1MG

A040510 001 Sep 17, 2004

WIGRAINE

ORGANON USA INC

100MG; 1MG

A086562 001

CALCIFEDIOL

CAPSULE; ORAL

CALDEROL

ORGANON USA INC

0.02MG

N018312 001

0.05MG

N018312 002

CALCIPOTRIENE

CREAM; TOPICAL

CALCIPOTRIENE

CHARTWELL RX

0.005%

A200935 001 May 30, 2012

SOLUTION; TOPICAL

CALCIPOTRIENE

CHARTWELL RX

0.005%

A077029 001 Nov 20, 2009

DOVONEX

+ LEO PHARM

0.005% **

N020611 001 Mar 03, 1997

CALCITONIN HUMAN

INJECTABLE; INJECTION

CIBACALCIN

NOVARTIS

0.5MG/VIAL

N018470 001 Oct 31, 1986

CALCITONIN SALMON

INJECTABLE; INJECTION

CALCIMAR

+ SANOFI AVENTIS US

200 IU/ML **

N017769 001

400 IU/VIAL

N017497 001

CALCITONIN-SALMON

IGI LABS INC

200 IU/ML

A073690 001 Apr 14, 1995

MIACALCIN

+ MYLAN IRELAND LTD

100 IU/ML **

N017808 001 Jul 03, 1986

SPRAY, METERED; NASAL

CALCITONIN-SALMON

PAR PHARM

200 IU/SPRAY

A076979 001 Jun 08, 2009

MIACALCIN

+ MYLAN IRELAND LTD

200 IU/SPRAY **

N020313 002 Aug 17, 1995

CALCITONIN SALMON RECOMBINANT

SPRAY, METERED; NASAL

FORTICAL

UPSHER SMITH LABS

200 IU/SPRAY **

N021406 001 Aug 12, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCITRIOL

INJECTABLE; INJECTION

CALCIJEX

+	ABBVIE	0.001MG/ML **	N018874 001	Sep 25, 1986
+		0.002MG/ML **	N018874 002	Sep 25, 1986

CALCITRIOL

AM REGENT	0.001MG/ML	A075746 001	Sep 26, 2003
	0.002MG/ML	A075746 002	Sep 26, 2003
FRESENIUS KABI USA	0.001MG/ML	A075836 001	Dec 31, 2002
	0.002MG/ML	A075836 002	Dec 31, 2002
FRESENIUS MEDCL	0.001MG/ML	A075766 001	Feb 20, 2003
	0.002MG/ML	A075766 002	Feb 20, 2003
HOSPIRA	0.001MG/ML	A075816 001	Jan 16, 2004
	0.002MG/ML	A075816 002	Jan 16, 2004
LONG GROVE PHARMS	0.002MG/ML	A078066 002	Jan 29, 2008
ROCKWELL MEDCL	0.001MG/ML	A076206 001	Sep 17, 2003
SAGENT PHARMS	0.001MG/ML	A077102 001	Feb 08, 2006
TEVA PARENTERAL	0.001MG/ML	A075823 001	Mar 31, 2003
	0.002MG/ML	A075823 002	Mar 31, 2003

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

AMNEAL PHARMS	667MG	A201658 001	Oct 06, 2014
LOTUS PHARM CO LTD	667MG	A203298 001	Jul 26, 2016

PHOSLO

FRESENIUS MEDCL	333.5MG	N021160 001	Apr 02, 2001
	667MG	N021160 002	Apr 02, 2001

SOLUTION; ORAL

PHOSLYRA

+	FRESENIUS MEDCL	667MG/5ML	N022581 001	Apr 18, 2011
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TABLET; ORAL

CALCIUM ACETATE

HIKMA	667MG	A077693 001	Jan 30, 2008
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ELIPHOS

CYPRESS PHARM	667MG	A078502 001	Nov 25, 2008
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PHOSLO

+	FRESENIUS MEDCL	667MG **	N019976 001	Dec 10, 1990
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CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL

FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE

PERRIGO R AND D	800MG;10MG;165MG	A204782 001	Aug 29, 2016
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CALCIUM CARBONATE; RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL WITH CALCIUM (COPACKAGED)

+	WARNER CHILCOTT	EQ 500MG BASE; 35MG **	N021823 001	Aug 12, 2005
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CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

METHOTREXATE SODIUM

EPIC PHARMA LLC	0.154MG/ML; 0.92MG/ML; 0.184MG/ML; 0.2MG/ML; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/ML	N020079 001	Feb 26, 1999
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CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL B22GK 2/0 IN PLASTIC CONTAINER

+	BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.157GM/1000ML; 2.21GM/1000ML; 7.07GM/1000ML (5000ML)	N021703 010	Oct 10, 2008
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PRISMASOL B22GK 2/2.5 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.157GM/1000ML; 2.21GM/1000ML; 7.07GM/1000ML (5000ML)	N021703 012	Oct 10, 2008
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PRISMASOL B22GK 4/2.5 IN PLASTIC CONTAINER

+	BAXTER HLTHCARE CORP	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML; 7.07GM/1000ML (5000ML)	N021703 013	Oct 10, 2008
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL BGK 4/0 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)	N021703 005	Oct 25, 2006
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PRISMASOL BGK 4/3.5 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	5.15GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)	N021703 008	Oct 25, 2006
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PRISMASOL BK 0/0 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	N/A/1000ML; N/A/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)	N021703 007	Oct 25, 2006
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PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER

+ BAXTER HLTHCARE CORP	5.15GM/1000ML; N/A/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)	N021703 001	Oct 25, 2006
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PRISMASOL BK 4/2.5 IN PLASTIC CONTAINER

BAXTER HLTHCARE CORP	3.68GM/1000ML; N/A/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML)	N021703 009	Oct 25, 2006
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATIONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

NAVSTEL

ALCON PHARMS LTD	0.154MG/ML; 0.92MG/ML; 0.2MG/ML; 0.184MG/ML; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/ML	N022193 001	Jul 24, 2008
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	37MG/100ML; 5GM/100ML; 31MG/100ML; 120MG/100ML; 330MG/100ML; 88MG/100ML	N019864 001	Jun 10, 1993
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ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	37MG/100ML; 5GM/100ML; 31MG/100ML; 120MG/100ML; 330MG/100ML; 88MG/100ML	N018271 001	
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	35MG/100ML; 5GM/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML	N019867 001	Dec 20, 1993
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ISOLYTE E W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	35MG/100ML; 5GM/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML	N018269 002	Jan 17, 1983
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE	37MG/100ML; 5GM/100ML; 30MG/100ML; 119MG/100ML; 161MG/100ML; 94MG/100ML; 138MG/100ML	N017390 001	
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CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION; INTRAPERITONEAL

DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER

B BRAUN	510MG/100ML; 30GM/100ML; 200MG/100ML; 9.2GM/100ML; 9.6GM/100ML	N018807 001	Aug 26, 1983
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	510MG/100ML; 30GM/100ML; 200MG/100ML; 9.4GM/100ML; 11GM/100ML	N018807 003	Aug 26, 1983
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DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER

B BRAUN	510MG/100ML; 50GM/100ML; 200MG/100ML; 9.2GM/100ML; 9.6GM/100ML	N018807 002	Aug 26, 1983
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	510MG/100ML; 50GM/100ML; 200MG/100ML; 9.4GM/100ML; 11GM/100ML	N018807 004	Aug 26, 1983
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DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN	29MG/100ML; 2.5GM/100ML; 15MG/100ML; 610MG/100ML; 560MG/100ML	N018460 006	Jan 29, 1986
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION;INTRAPERITONEAL

DIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER			
B BRAUN	29MG/100ML;1.5GM/100ML;15MG/100ML;610MG	N018460	001
	/100ML;560MG/100ML		
DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER			
B BRAUN	29MG/100ML;4.25GM/100ML;15MG/100ML;610M	N018460	003
	G/100ML;560MG/100ML		

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION;INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5	N018379	002
	67MG/100ML;392MG/100ML		
DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N018379	003
	67MG/100ML;392MG/100ML		
DELFLX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;5	N018379	007 Jun 24, 1988
	67MG/100ML;392MG/100ML		
DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;	N018379	001
	567MG/100ML;392MG/100ML		
DELFLX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5	N018379	004 Jul 07, 1982
	38MG/100ML;448MG/100ML		
DELFLX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5	N018379	005 Jul 07, 1982
	38MG/100ML;448MG/100ML		
DELFLX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;5	N018379	008 Jun 24, 1988
	38MG/100ML;448MG/100ML		
DELFLX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER			
FRESENIUS MEDCL	25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;	N018379	006 Jul 07, 1982
	538MG/100ML;448MG/100ML		
DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER			
B BRAUN	26MG/100ML;1.5GM/100ML;5MG/100ML;530MG/	N018460	007 Jan 29, 1986
	100ML;450MG/100ML		
	26MG/100ML;1.5GM/100ML;15MG/100ML;560MG	N018460	002
	/100ML;390MG/100ML		
DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER			
B BRAUN	26MG/100ML;2.5GM/100ML;5MG/100ML;530MG/	N018460	005 Nov 02, 1983
	100ML;450MG/100ML		
	26MG/100ML;5GM/100ML;5MG/100ML;530MG/10	N018460	008 Jan 29, 1986
	0ML;450MG/100ML		
DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER			
B BRAUN	26MG/100ML;4.25GM/100ML;5MG/100ML;530MG	N018460	009 Jan 29, 1986
	/100ML;450MG/100ML		
	26MG/100ML;4.25GM/100ML;15MG/100ML;560M	N018460	004
	G/100ML;390MG/100ML		
DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5	N017512	001
	67MG/100ML;392MG/100ML		
DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N017512	003
	67MG/100ML;392MG/100ML		
DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;	N017512	002
	567MG/100ML;392MG/100ML		
DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER			
+ BAXTER HLTHCARE	18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;5	N020183	003 Dec 04, 1992
	38MG/100ML;448MG/100ML		
DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5	N017512	007 Jul 09, 1984
	67MG/100ML;392MG/100ML		
DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5	N017512	008 Jul 09, 1984
	67MG/100ML;392MG/100ML		
DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;5	N017512	010 Nov 18, 1985
	67MG/100ML;392MG/100ML		
DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;	N017512	009 Jul 09, 1984
	567MG/100ML;392MG/100ML		

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION;INTRAPERITONEAL

DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER					
+ BAXTER HLTHCARE	25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;5	N017512	011	Nov 18, 1985	
	38MG/100ML;448MG/100ML				
INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5	A020374	001	Jun 13, 1994	
	38MG/100ML;448MG/100ML				
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML;2.5GM/100ML;5.08MG/100ML;5	A020374	002	Jun 13, 1994	
	38MG/100ML;448MG/100ML				
INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML;3.5GM/100ML;5.08MG/100ML;5	A020374	003	Jun 13, 1994	
	38MG/100ML;448MG/100ML				
INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER					
FRESENIUS	18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;	A020374	004	Jun 13, 1994	
	538MG/100ML;448MG/100ML				

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE;INJECTION

DEXTROSE 5% IN ACETATED RINGER'S IN PLASTIC CONTAINER					
B BRAUN	20MG/100ML;5GM/100ML;30MG/100ML;380MG/1	N018258	001		
	00ML;600MG/100ML				

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE;INJECTION

DEXTROSE 5% AND RINGER'S IN PLASTIC CONTAINER					
HOSPIRA	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1	N018254	001		
	00ML				
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER					
B BRAUN	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1	N018256	001		
	00ML				
	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1	N020000	001	Apr 17, 1992	
	00ML				
BAXTER HLTHCARE	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1	N016695	001		
	00ML				

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE;INJECTION

DEXTROSE 4% IN MODIFIED LACTATED RINGER'S IN PLASTIC CONTAINER					
B BRAUN	4MG/100ML;4GM/100ML;6MG/100ML;120MG/100	N019634	002	Feb 24, 1988	
	ML;62MG/100ML				
DEXTROSE 5% AND LACTATED RINGER'S					
FRESENIUS KABI USA	20MG/100ML;5GM/100ML;30MG/100ML;600MG/1	A210332	001	Mar 28, 2022	
	00ML;310MG/100ML				
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER					
B BRAUN	20MG/100ML;5GM/100ML;30MG/100ML;600MG/1	N017510	001		
	00ML;310MG/100ML				
MILES	20MG/100ML;5GM/100ML;30MG/100ML;600MG/1	N018499	001		
	00ML;310MG/100ML				
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;104MG/100ML;600MG/	N019685	005	Oct 17, 1988	
	100ML;310MG/100ML				
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;179MG/100ML;600MG/	N019685	006	Oct 17, 1988	
	100ML;310MG/100ML				
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;254MG/100ML;600MG/	N019685	007	Oct 17, 1988	
	100ML;310MG/100ML				
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
FRESENIUS KABI USA	20MG/100ML;5GM/100ML;179MG/100ML;600MG/	A211428	001	Nov 09, 2021	
	100ML;310MG/100ML				
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;328MG/100ML;600MG/	N019685	008	Oct 17, 1988	
	100ML;310MG/100ML				
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;254MG/100ML;600MG/	N019685	003	Oct 17, 1988	
	100ML;310MG/100ML				
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;328MG/100ML;600MG/	N019685	004	Oct 17, 1988	
	100ML;310MG/100ML				
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER					
+ ICU MEDICAL INC	20MG/100ML;5GM/100ML;104MG/100ML;600MG/	N019685	001	Oct 17, 1988	
	100ML;310MG/100ML				

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

INPERSOL-ZM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL 25.7MG/100ML; 1.5GM/100ML; 538MG/100ML; 44 N019395 001 Mar 26, 1986
8MG/100ML

INPERSOL-ZM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL 25.7MG/100ML; 2.5GM/100ML; 538MG/100ML; 44 N019395 002 Mar 26, 1986
8MG/100ML

INPERSOL-ZM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

FRESENIUS MEDCL 25.7MG/100ML; 4.25GM/100ML; 538MG/100ML; 4 N019395 003 Mar 26, 1986
48MG/100MLCALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

ABBOTT 16.5MG/ML; 25.4MG/ML; 74.6MG/ML; 121MG/ML; N019399 001 Jun 16, 1986
16.1MG/MLCALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN PLASTIC CONTAINER

B BRAUN 35MG/100ML; 30MG/100ML; 74MG/100ML; 640MG/ N018899 001 Oct 31, 1983
100ML; 500MG/100ML; 74MG/100ML35MG/100ML; 30MG/100ML; 74MG/100ML; 640MG/ N019718 001 Sep 29, 1989
100ML; 500MG/100ML; 74MG/100ML

SOLUTION; IRRIGATION

BALANCED SALT

EPIC PHARMA LLC 0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6 A075503 001 Sep 27, 2006
.4MG/ML; 1.7MG/MLCALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE R IN PLASTIC CONTAINER

BAXTER HLTHCARE 36.8MG/100ML; 30.5MG/100ML; 74.6MG/100ML; N017438 001
640MG/100ML; 496MG/100ML; 89.6MG/100MLCALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ACETATED RINGER'S IN PLASTIC CONTAINER

B BRAUN 20MG/100ML; 30MG/100ML; 380MG/100ML; 600MG N018725 001 Nov 29, 1982
/100MLCALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

B BRAUN 33MG/100ML; 30MG/100ML; 860MG/100ML N018721 001 Nov 09, 1982

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

ABBOTT 33MG/100ML; 30MG/100ML; 860MG/100ML N018462 001

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

ABBOTT 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG N019485 001 Oct 24, 1985
/100MLB BRAUN 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG N018023 001
/100MLMILES 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG N018417 001
/100ML

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG N019933 001 Aug 29, 1989
/100MLCALCIUM GLUCEPTATE

INJECTABLE; INJECTION

CALCIUM GLUCEPTATE

+ ABBOTT EQ 90MG CALCIUM/5ML A080001 001

EQ 90MG CALCIUM/5ML A083159 001

ABRAXIS PHARM EQ 90MG CALCIUM/5ML A089373 001 Apr 30, 1987

LILLY EQ 90MG CALCIUM/5ML N006470 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM METRIZOATE; MEGLUMINE METRIZOATE; METRIZOATE MAGNESIUM; METRIZOATE SODIUM

INJECTABLE; INJECTION

ISOPAQUE 440

GE HEALTHCARE 0.78MG/ML; 75.9MG/ML; 0.15MG/ML; 16.6MG/ML N016847 001

CALCIUM; MEGLUMINE; METRIZOIC ACID

INJECTABLE; INJECTION

ISOPAQUE 280

GE HEALTHCARE 0.35MG/ML; 140.1MG/ML; 461.8MG/ML N017506 001

CANDESARTAN CILEXETIL

TABLET; ORAL

CANDESARTAN CILEXETIL

APOTEX	4MG	A202079 001	Jan 10, 2014
	8MG	A202079 002	Jan 10, 2014
	16MG	A202079 003	Jan 10, 2014
	32MG	A202079 004	Jan 10, 2014

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

APOTEX INC	16MG; 12.5MG	A202884 001	Dec 04, 2012
	32MG; 12.5MG	A202884 002	Dec 04, 2012
	32MG; 25MG	A202884 003	Jun 03, 2013

CANDICIDIN

OINTMENT; VAGINAL

VANOVID

SANOFI AVENTIS US 0.6MG/GM A061596 001

TABLET; VAGINAL

VANOVID

SANOFI AVENTIS US 3MG A061613 001

CAPECITABINE

TABLET; ORAL

CAPECITABINE

AMNEAL PHARMS	150MG	A204741 001	Feb 28, 2017
	500MG	A204741 002	Feb 28, 2017

CAPREOMYCIN SULFATE

INJECTABLE; INJECTION

CAPASTAT SULFATE

+ EPIC PHARMA LLC EQ 1GM BASE/VIAL N050095 001

CAPREOMYCIN SULFATE

HISUN PHARM HANGZHOU	EQ 1GM BASE/VIAL	A204796 001	Oct 18, 2018
MYLAN LABS LTD	EQ 1GM BASE/VIAL	A202634 001	Nov 27, 2017

CAPTOPRIL

TABLET; ORAL

CAPOTEN

+ STRIDES PHARMA	12.5MG **	N018343 005	Jan 17, 1985
+	25MG **	N018343 002	
+	37.5MG **	N018343 006	Sep 17, 1986
+	50MG **	N018343 001	
+	75MG **	N018343 007	Jun 13, 1995
+	100MG **	N018343 003	
+	150MG **	N018343 004	Jun 13, 1995

CAPTOPRIL

APOTHECON	12.5MG	A074472 001	Mar 31, 1995
	25MG	A074472 002	Mar 31, 1995
	50MG	A074472 003	Mar 31, 1995
	100MG	A074472 004	Mar 31, 1995
AUROBINDO PHARMA USA	12.5MG	A074434 001	Feb 13, 1996
	25MG	A074434 002	Feb 13, 1996
	50MG	A074434 003	Feb 13, 1996
	100MG	A074434 004	Feb 13, 1996
CHARTWELL RX	12.5MG	A074363 001	Nov 09, 1995
	12.5MG	A074519 001	Feb 13, 1996
	25MG	A074363 002	Nov 09, 1995
	25MG	A074519 002	Feb 13, 1996
	50MG	A074363 003	Nov 09, 1995
	50MG	A074519 003	Feb 13, 1996
	100MG	A074363 004	Nov 09, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRILTABLET; ORAL
CAPTOPRIL

	100MG	A074519	004	Feb 13, 1996
COSETTE	12.5MG	A074433	001	Feb 13, 1996
	12.5MG	A074462	001	Feb 13, 1996
	12.5MG	A074483	001	Feb 13, 1996
	25MG	A074433	002	Feb 13, 1996
	25MG	A074462	002	Feb 13, 1996
	25MG	A074483	002	Feb 13, 1996
	50MG	A074433	003	Feb 13, 1996
	50MG	A074462	003	Feb 13, 1996
	50MG	A074483	003	Feb 13, 1996
	100MG	A074433	004	Feb 13, 1996
	100MG	A074462	004	Feb 13, 1996
	100MG	A074483	004	Feb 13, 1996
DAVA PHARMS INC	12.5MG	A074423	001	Feb 13, 1996
	25MG	A074423	002	Feb 13, 1996
	50MG	A074423	003	Feb 13, 1996
	100MG	A074423	004	Feb 13, 1996
EGIS PHARMS	12.5MG	A074748	004	May 29, 1997
	25MG	A074748	002	May 29, 1997
	50MG	A074748	001	May 29, 1997
	100MG	A074748	003	May 29, 1997
G AND W LABS INC	12.5MG	A074590	004	Aug 30, 1996
	25MG	A074590	002	Aug 30, 1996
	50MG	A074590	001	Aug 30, 1996
	100MG	A074590	003	Aug 30, 1996
OXFORD PHARMS	12.5MG	A074418	001	Feb 13, 1996
	25MG	A074418	002	Feb 13, 1996
	50MG	A074418	003	Feb 13, 1996
	100MG	A074418	004	Feb 13, 1996
PUREPAC PHARM	12.5MG	A074640	001	Mar 31, 1997
	25MG	A074640	002	Mar 31, 1997
	50MG	A074640	003	Mar 31, 1997
	100MG	A074640	004	Mar 31, 1997
SANDOZ	12.5MG	A074481	001	Feb 13, 1996
	25MG	A074481	002	Feb 13, 1996
	50MG	A074481	003	Feb 13, 1996
	100MG	A074481	004	Feb 13, 1996
SETON PHARMS	12.5MG	A212223	001	Oct 30, 2019
	25MG	A212223	002	Oct 30, 2019
	50MG	A212223	003	Oct 30, 2019
	100MG	A212223	004	Oct 30, 2019
STRIDES PHARMA	12.5MG	A074493	001	Feb 13, 1996
	25MG	A074493	002	Feb 13, 1996
	50MG	A074493	003	Feb 13, 1996
	100MG	A074493	004	Feb 13, 1996
TEVA	12.5MG	A074322	001	Feb 13, 1996
	25MG	A074322	002	Feb 13, 1996
	50MG	A074322	003	Feb 13, 1996
	100MG	A074322	004	Feb 13, 1996
WATSON LABS	12.5MG	A074386	001	May 23, 1996
	12.5MG	A074451	001	Feb 13, 1996
	12.5MG	A074576	001	Apr 23, 1996
	25MG	A074386	002	May 23, 1996
	25MG	A074451	002	Feb 13, 1996
	25MG	A074576	002	Apr 23, 1996
	50MG	A074386	003	May 23, 1996
	50MG	A074451	003	Feb 13, 1996
	50MG	A074576	003	Apr 23, 1996
	100MG	A074386	004	May 23, 1996
	100MG	A074451	004	Feb 13, 1996
	100MG	A074576	004	Apr 23, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPOZIDE 25/15					
+ APOTHECON	25MG;15MG **		N018709 001	Oct 12, 1984	
CAPOZIDE 25/25					
+ APOTHECON	25MG;25MG **		N018709 002	Oct 12, 1984	
CAPOZIDE 50/15					
+ APOTHECON	50MG;15MG **		N018709 004	Oct 12, 1984	
CAPOZIDE 50/25					
+ APOTHECON	50MG;25MG **		N018709 003	Oct 12, 1984	
CAPTOPRIL AND HYDROCHLOROTHIAZIDE					
COSETTE	25MG;15MG		A074827 001	Dec 29, 1997	
	25MG;25MG		A074827 002	Dec 29, 1997	
	50MG;15MG		A074827 004	Dec 29, 1997	
	50MG;25MG		A074827 003	Dec 29, 1997	
IVAX SUB TEVA PHARMS	25MG;15MG		A075055 001	Jun 18, 1998	
	25MG;25MG		A075055 002	Jun 18, 1998	
	50MG;15MG		A075055 004	Jun 18, 1998	
	50MG;25MG		A075055 003	Jun 18, 1998	
STRIDES PHARMA	25MG;15MG		A074788 001	Dec 29, 1997	
	25MG;25MG		A074788 002	Dec 29, 1997	
	50MG;15MG		A074788 004	Dec 29, 1997	
	50MG;25MG		A074788 003	Dec 29, 1997	
WATSON LABS	50MG;25MG		A074832 001	Dec 29, 1997	

CARBACHOL

SOLUTION; INTRAOCULAR

CARBACHOL					
PHARMAFAIR	0.01%		A070292 001	May 21, 1986	
CARBASTAT					
NOVARTIS	0.01%		A073677 001	Apr 28, 1995	

CARBAMAZEPINE

SOLUTION; INTRAVENOUS

CARNEXIV					
+ LUNDBECK PHARMS LLC	200MG/20ML (10MG/ML)		N206030 001	Oct 07, 2016	

SUSPENSION; ORAL

CARBAMAZEPINE					
TARO	100MG/5ML		A075875 001	Dec 21, 2000	

TABLET; ORAL

CARBAMAZEPINE					
ACTAVIS ELIZABETH	200MG		A071696 001	Nov 09, 1987	
INWOOD LABS	200MG		A070231 001	Aug 14, 1986	
PLIVA	200MG		A071479 001	Jul 24, 1987	
RK PHARMA	200MG		A214328 001	Aug 16, 2021	
USL PHARMA	200MG		A070300 001	May 15, 1986	
WARNER CHILCOTT	200MG		A070429 001	Jan 02, 1987	

TERIL

TARO	200MG		A076525 001	Sep 26, 2003	
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TABLET, CHEWABLE; ORAL

CARBAMAZEPINE					
JUBILANT CADISTA	100MG		A071940 001	Feb 01, 1988	
TEGRETOL					
+ NOVARTIS	100MG **		N018281 001		

TABLET, EXTENDED RELEASE; ORAL

CARBAMAZEPINE					
AJANTA PHARMA LTD	100MG		A216193 001	Mar 24, 2023	
	200MG		A216193 002	Mar 24, 2023	
	400MG		A216193 003	Mar 24, 2023	

CARBENICILLIN DISODIUM

INJECTABLE; INJECTION

GEOPEN

ROERIG	EQ 1GM BASE/VIAL		N050306 001		
	EQ 2GM BASE/VIAL		N050306 004		
	EQ 5GM BASE/VIAL		N050306 002		
	EQ 10GM BASE/VIAL		N050306 006		
	EQ 30GM BASE/VIAL		N050306 007		

PYOPEN

GLAXOSMITHKLINE	EQ 1GM BASE/VIAL		N050298 001		
	EQ 2GM BASE/VIAL		N050298 002		

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBENICILLIN DISODIUMINJECTABLE; INJECTION
PYOPEN

EQ 5GM BASE/VIAL	N050298	003
EQ 10GM BASE/VIAL	N050298	006
EQ 20GM BASE/VIAL	N050298	007

CARBENICILLIN INDANYL SODIUM

TABLET; ORAL

GEOCILLIN

PFIZER

EQ 382MG BASE

N050435 001

CARBIDOPA

TABLET; ORAL

CARBIDOPA

ANI PHARMS

25MG

A203261 001 Mar 10, 2014

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

MYLAN

12.5MG; 200MG; 50MG	A203424	001	Aug 13, 2020
18.75MG; 200MG; 75MG	A203424	002	Aug 13, 2020
25MG; 200MG; 100MG	A203424	003	Aug 13, 2020
31.25MG; 200MG; 125MG	A203424	004	Aug 13, 2020
37.5MG; 200MG; 150MG	A203424	005	Aug 13, 2020
50MG; 200MG; 200MG	A203424	006	Aug 13, 2020
12.5MG; 200MG; 50MG	A090786	001	Nov 20, 2012
18.75MG; 200MG; 75MG	A090833	001	Nov 20, 2012
25MG; 200MG; 100MG	A090833	002	Nov 20, 2012
31.25MG; 200MG; 125MG	A090833	003	Nov 20, 2012
37.5MG; 200MG; 150MG	A090833	004	Nov 20, 2012
50MG; 200MG; 200MG	A090833	005	Nov 20, 2012

WOCKHARDT LTD

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

ANI PHARMS

10MG; 100MG	A073587	002	Jun 29, 1995
25MG; 100MG	A073587	001	Jun 29, 1995
25MG; 250MG	A073587	003	Jun 29, 1995
10MG; 100MG	A074080	001	Mar 25, 1994
25MG; 100MG	A074080	002	Mar 25, 1994
25MG; 250MG	A074080	003	Mar 25, 1994
10MG; 100MG	A073381	001	Sep 28, 1993
25MG; 100MG	A073382	001	Sep 28, 1993
25MG; 250MG	A073383	001	Sep 28, 1993
10MG; 100MG	A215999	001	Apr 04, 2023
25MG; 100MG	A215999	002	Apr 04, 2023
25MG; 250MG	A215999	003	Apr 04, 2023

SCS

WATSON LABS

ZYDUS PHARMS

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

KV PHARM

50MG; 200MG

A076663 001 Jun 24, 2004

SINEMET CR

+ ORGANON LLC

25MG; 100MG **

N019856 002 Dec 24, 1992

+

50MG; 200MG **

N019856 001 May 30, 1991

TABLET, FOR SUSPENSION; ORAL

CARBILEV

RANBAXY

10MG; 100MG	A076643	001	Jun 10, 2005
25MG; 100MG	A076643	002	Jun 10, 2005
25MG; 250MG	A076643	003	Jun 10, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPA

IMPAX LABS

10MG; 100MG	A090631	001	Jun 08, 2010
25MG; 100MG	A090631	002	Jun 08, 2010
25MG; 250MG	A090631	003	Jun 08, 2010
10MG; 100MG	A078893	001	Sep 18, 2008
25MG; 100MG	A078893	002	Sep 18, 2008
25MG; 250MG	A078893	003	Sep 18, 2008

RISING

PARCOPA

UCB INC

10MG; 100MG **	A076699	001	Aug 27, 2004
25MG; 100MG **	A076699	002	Aug 27, 2004
25MG; 250MG **	A076699	003	Aug 27, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBINOXAMINE MALEATE

ELIXIR; ORAL

CLISTIN

+ MCNEIL

4MG/5ML **

N008955 001

SOLUTION; ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM

4MG/5ML

A090418 001 May 04, 2010

VINTAGE PHARMS

4MG/5ML

A040814 001 Feb 26, 2008

TABLET; ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM

4MG

A090417 001 Aug 23, 2010

STRIDES PHARMA

4MG

A040639 002 May 30, 2008

CLISTIN

+ ORTHO MCNEIL PHARM

4MG **

N008915 001

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

CIPLA LTD

50MG/VIAL

A077383 001 Jan 27, 2006

150MG/VIAL

A077383 002 Jan 27, 2006

450MG/VIAL

A077383 003 Jan 27, 2006

FRESENIUS KABI USA

50MG/VIAL

A076235 001 Oct 14, 2004

150MG/VIAL

A076235 002 Oct 14, 2004

450MG/VIAL

A076235 003 Oct 14, 2004

HIKMA

50MG/VIAL

A076099 001 Oct 14, 2004

150MG/VIAL

A076099 002 Oct 14, 2004

450MG/VIAL

A076099 003 Oct 14, 2004

HOSPIRA

50MG/VIAL

A076473 001 Oct 27, 2004

150MG/VIAL

A076473 002 Oct 27, 2004

450MG/VIAL

A076473 003 Oct 27, 2004

NATCO PHARMA USA

50MG/VIAL

A091510 001 May 29, 2012

150MG/VIAL

A091510 002 May 29, 2012

450MG/VIAL

A091510 003 May 29, 2012

PLIVA

50MG/VIAL

A076602 001 Nov 16, 2004

150MG/VIAL

A076602 002 Nov 16, 2004

450MG/VIAL

A076602 003 Nov 16, 2004

SANDOZ

50MG/VIAL

A076959 001 Mar 18, 2005

150MG/VIAL

A076959 002 Mar 18, 2005

450MG/VIAL

A076959 003 Mar 18, 2005

WATSON LABS TEVA

50MG/VIAL

A076162 001 Oct 14, 2004

150MG/VIAL

A076162 002 Oct 14, 2004

450MG/VIAL

A076162 003 Oct 14, 2004

PARAPLATIN

+ CORDEN PHARMA

50MG/VIAL **

N019880 001 Mar 03, 1989

+

150MG/VIAL **

N019880 002 Mar 03, 1989

+

450MG/VIAL **

N019880 003 Mar 03, 1989

INJECTABLE; INTRAVENOUS

CARBOPLATIN

ACTAVIS TOTOWA

50MG/5ML (10MG/ML)

A078732 001 Feb 06, 2012

150MG/15ML (10MG/ML)

A078732 002 Feb 06, 2012

450MG/45ML (10MG/ML)

A078732 003 Feb 06, 2012

600MG/60ML (10MG/ML)

A078732 004 Feb 06, 2012

FRESENIUS KABI USA

50MG/5ML (10MG/ML)

A077432 001 Sep 29, 2006

150MG/15ML (10MG/ML)

A077432 002 Sep 29, 2006

450MG/45ML (10MG/ML)

A077432 003 Sep 29, 2006

50MG/5ML (10MG/ML)

A077247 001 Oct 21, 2004

50MG/5ML (10MG/ML)

A077266 001 Feb 15, 2006

150MG/15ML (10MG/ML)

A077247 002 Oct 21, 2004

150MG/15ML (10MG/ML)

A077266 002 Feb 15, 2006

MEITHEAL

50MG/5ML (10MG/ML)

A077096 001 Jun 14, 2005

150MG/15ML (10MG/ML)

A077096 002 Jun 14, 2005

450MG/45ML (10MG/ML)

A077096 003 Jun 14, 2005

600MG/60ML (10MG/ML)

A077096 004 Jun 03, 2013

MYLAN INSTITUTIONAL

50MG/5ML (10MG/ML)

A077998 001 Apr 24, 2007

150MG/15ML (10MG/ML)

A077998 002 Apr 24, 2007

450MG/45ML (10MG/ML)

A077998 003 Apr 24, 2007

MYLAN LABS LTD

50MG/5ML (10MG/ML)

A091063 001 Nov 09, 2011

150MG/15ML (10MG/ML)

A091063 002 Nov 09, 2011

450MG/45ML (10MG/ML)

A091063 003 Nov 09, 2011

600MG/60ML (10MG/ML)

A091063 004 Nov 09, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBOPLATININJECTABLE; INTRAVENOUS
CARBOPLATIN

	1GM/100ML (10MG/ML)	A091478 001	Nov 23, 2011
PHARMACHEMIE BV	50MG/5ML (10MG/ML)	A077679 001	Feb 25, 2009
	150MG/15ML (10MG/ML)	A077679 002	Feb 25, 2009
	450MG/45ML (10MG/ML)	A077679 003	Feb 25, 2009
PLIVA LACHEMA	50MG/5ML (10MG/ML)	A078631 001	Dec 02, 2008
	150MG/15ML (10MG/ML)	A078631 002	Dec 02, 2008
	450MG/45ML (10MG/ML)	A078631 003	Dec 02, 2008
	600MG/60ML (10MG/ML)	A078631 004	Dec 02, 2008
TEVA PARENTERAL	50MG/5ML (10MG/ML)	A077389 001	Mar 30, 2007
	150MG/15ML (10MG/ML)	A077389 002	Mar 30, 2007
	450MG/45ML (10MG/ML)	A077389 003	Mar 30, 2007
TEVA PHARMS USA	50MG/5ML (10MG/ML)	A077139 001	Sep 21, 2005
	150MG/15ML (10MG/ML)	A077139 002	Sep 21, 2005
	450MG/45ML (10MG/ML)	A077139 003	Sep 21, 2005
	600MG/60ML (10MG/ML)	A077139 004	Sep 21, 2005
PARAPLATIN			
+ CORDENPHARMA	50MG/5ML (10MG/ML) **	N020452 001	Jul 14, 2003
+	150MG/15ML (10MG/ML) **	N020452 002	Jul 14, 2003
+	450MG/45ML (10MG/ML) **	N020452 003	Jul 14, 2003
+	600MG/60ML (10MG/ML) **	N020452 004	Jan 15, 2004

CARFILZOMIBPOWDER; INTRAVENOUS
CARFILZOMIB

APOTEX	30MG/VIAL	A211185 001	Mar 20, 2020
	60MG/VIAL	A209425 001	Mar 16, 2020
BRECKENRIDGE	10MG/VIAL	A209330 001	Jun 11, 2021
	60MG/VIAL	A209330 002	Jun 11, 2021

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL

CARIPRAZINE HYDROCHLORIDE

SUN PHARM	EQ 1.5MG BASE	A213932 001	Sep 30, 2022
	EQ 3MG BASE	A213932 002	Sep 30, 2022
	EQ 4.5MG BASE	A213932 003	Sep 30, 2022
	EQ 6MG BASE	A213932 004	Sep 30, 2022
ZYDUS	EQ 1.5MG BASE	A213984 001	Sep 09, 2022
	EQ 3MG BASE	A213984 002	Sep 09, 2022
	EQ 4.5MG BASE	A213984 003	Sep 09, 2022
	EQ 6MG BASE	A213984 004	Sep 09, 2022

CARISOPRODOL

CAPSULE; ORAL

SOMA

+ MYLAN SPECIALITY LP 250MG N011792 003

TABLET; ORAL

CARISOPRODOL

ABLE	350MG	A040421 001	Jun 21, 2001
EPIC PHARMA LLC	350MG	A040397 001	Sep 21, 2000
HIKMA INTL PHARMS	350MG	A040124 001	Jan 24, 1996
PIONEER PHARMS	350MG	A089390 001	Oct 13, 1988
SANDOZ	350MG	A089566 001	Aug 30, 1988
STRIDES PHARMA	250MG	A205513 001	Nov 12, 2015
	350MG	A205513 002	Nov 12, 2015
SUN PHARM INDS LTD	350MG	A040755 001	Feb 27, 2007
SUN PHARM INDUSTRIES	350MG	A089346 001	Oct 17, 1991
WATSON LABS	350MG	A040152 001	Dec 03, 1996
	350MG	A085433 001	
WATSON LABS TEVA	350MG	A086179 001	
RELA			
SCHERING	350MG	N012155 001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARMUSTINE

INJECTABLE; INJECTION

CARMUSTINE

ACCORD HLTHCARE	100MG/VIAL	A214117 001	Dec 27, 2022
MYLAN LABS LTD	100MG/VIAL	A215368 001	Mar 03, 2023

CARPHENAZINE MALEATE

CONCENTRATE; ORAL

PROKETAZINE

WYETH AYERST	50MG/ML	N014173 001	
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TABLET; ORAL

PROKETAZINE

WYETH AYERST	12.5MG	N012768 001	
	25MG	N012768 002	
	50MG	N012768 004	

CARPROFEN

TABLET; ORAL

RIMADYL

ROCHE	100MG	N018550 002	Dec 31, 1987
	150MG	N018550 003	Dec 31, 1987

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

APOTEX INC	1%	A076097 001	Feb 06, 2002
BAUSCH AND LOMB	1%	A075546 001	Jan 20, 2000

OCUPRESS

+ NOVARTIS

1% **

N019972 001 May 23, 1990

TABLET; ORAL

CARTROL

ABBVIE	2.5MG	N019204 001	Dec 28, 1988
	5MG	N019204 002	Dec 28, 1988
	10MG	N019204 003	Dec 28, 1988

CARVEDILOL

TABLET; ORAL

CARVEDILOL

ACI	3.125MG	A078786 001	Dec 22, 2009
	6.25MG	A078786 002	Dec 22, 2009
	12.5MG	A078786 003	Dec 22, 2009
	25MG	A078786 004	Dec 22, 2009
HIKMA	3.125MG	A077887 001	Sep 07, 2007
	6.25MG	A077887 002	Sep 07, 2007
	12.5MG	A077887 003	Sep 07, 2007
	25MG	A077887 004	Sep 07, 2007
PLIVA HRVATSKA DOO	3.125MG	A078240 001	Oct 30, 2007
	6.25MG	A078240 002	Oct 30, 2007
	12.5MG	A078240 003	Oct 30, 2007
	25MG	A078240 004	Oct 30, 2007
SUN PHARM INDS INC	3.125MG	A077346 004	Sep 05, 2007
	6.25MG	A077346 001	Sep 05, 2007
	12.5MG	A077346 002	Sep 05, 2007
	25MG	A077346 003	Sep 05, 2007

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CASPOFUNGIN ACETATE

CIPLA	50MG/VIAL	A209489 001	Jul 12, 2018
	70MG/VIAL	A209489 002	Jul 12, 2018
MYLAN LABS LTD	50MG/VIAL	A207650 001	Sep 29, 2017
	70MG/VIAL	A207650 002	Sep 29, 2017
XELLIA PHARMS APS	50MG/VIAL	A205923 001	Jul 02, 2018
	70MG/VIAL	A205923 002	Jul 02, 2018

CEFACLOR

CAPSULE; ORAL

CECLOR

+ LILLY

EQ 250MG BASE **

N050521 001

+

EQ 500MG BASE **

N050521 002

CEFACLOR

CEPH INTL	EQ 250MG BASE	A062205 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFACTOR

CAPSULE; ORAL

CEFACTOR

	EQ 500MG BASE	A062205 002	
CHARTWELL RX	EQ 250MG BASE	A064148 001	May 23, 1996
	EQ 500MG BASE	A064148 002	May 23, 1996
DAVA PHARMS INC	EQ 250MG BASE	A064107 001	Apr 27, 1995
	EQ 500MG BASE	A064107 002	Apr 27, 1995
HIKMA	EQ 250MG BASE	A065350 001	Apr 03, 2007
	EQ 500MG BASE	A065350 002	Apr 03, 2007
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A064061 001	Apr 27, 1995
	EQ 500MG BASE	A064061 002	Apr 27, 1995
RANBAXY	EQ 250MG BASE	A064156 001	Aug 28, 1997
	EQ 500MG BASE	A064156 002	Aug 28, 1997
TEVA	EQ 250MG BASE	A064081 001	Sep 16, 1996
	EQ 250MG BASE	A064145 001	Jun 24, 1996
	EQ 500MG BASE	A064081 002	Sep 16, 1996
	EQ 500MG BASE	A064145 002	Jun 24, 1996

FOR SUSPENSION; ORAL

CECLOR

+ LILLY

EQ 125MG BASE/5ML **

N050522 001

+

EQ 250MG BASE/5ML **

N050522 002

CEFACTOR

DAVA PHARMS INC

EQ 125MG BASE/5ML

A064114 001 Apr 28, 1995

EQ 187MG BASE/5ML

A064115 001 Apr 28, 1995

EQ 250MG BASE/5ML

A064116 001 Apr 28, 1995

EQ 375MG BASE/5ML

A064110 001 Apr 28, 1995

FACTA FARMA

EQ 125MG BASE/5ML

A062206 001

EQ 187MG BASE/5ML

A062206 003 Apr 20, 1988

EQ 250MG BASE/5ML

A062206 002

EQ 375MG BASE/5ML

A062206 004 Apr 20, 1988

IVAX SUB TEVA PHARMS

EQ 125MG BASE/5ML

A064087 001 Apr 28, 1995

EQ 187MG BASE/5ML

A064086 001 Apr 28, 1995

EQ 250MG BASE/5ML

A064085 001 Apr 28, 1995

EQ 375MG BASE/5ML

A064070 001 Apr 28, 1995

RANBAXY

EQ 125MG BASE/5ML

A064166 001 Oct 02, 1997

EQ 187MG BASE/5ML

A064165 001 Oct 02, 1997

EQ 250MG BASE/5ML

A064164 001 Oct 02, 1997

EQ 375MG BASE/5ML

A064155 001 Oct 02, 1997

WATSON LABS INC

EQ 125MG BASE/5ML

A064204 001 Feb 18, 1998

EQ 187MG BASE/5ML

A064205 001 Feb 18, 1998

EQ 250MG BASE/5ML

A064206 001 Feb 18, 1998

EQ 375MG BASE/5ML

A064207 001 Feb 18, 1998

TABLET, CHEWABLE; ORAL

RANICLOR

RANBAXY LABS LTD

EQ 125MG BASE

A065092 001 Dec 22, 2003

EQ 187MG BASE

A065092 002 Dec 22, 2003

EQ 250MG BASE

A065092 003 Dec 22, 2003

EQ 375MG BASE

A065092 004 Dec 22, 2003

TABLET, EXTENDED RELEASE; ORAL

CECLOR CD

LILLY

EQ 375MG BASE **

N050673 001 Jun 28, 1996

EQ 500MG BASE **

N050673 002 Jun 28, 1996

CEFACTOR

WORLD GEN

EQ 500MG BASE

A065057 001 Jan 05, 2001

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

CHARTWELL RX

EQ 500MG BASE

A065309 001 Sep 18, 2006

CSPC OUYI

EQ 500MG BASE

A205072 001 Jul 28, 2017

HIKMA

EQ 500MG BASE

A065311 001 Feb 07, 2006

IVAX SUB TEVA PHARMS

EQ 500MG BASE

A062766 001 Mar 03, 1987

PUREPAC PHARM

EQ 500MG BASE

A063017 001 Jan 05, 1989

RANBAXY LABS LTD

EQ 500MG BASE

A065015 001 Jun 22, 1999

SANDOZ

EQ 500MG BASE

A062291 001

TEVA

EQ 500MG BASE

A062695 001 Feb 10, 1989

DURICEF

WARNER CHILCOTT

EQ 250MG BASE

N050512 002

+

EQ 500MG BASE **

N050512 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

ULTRACEF

BRISTOL EQ 500MG BASE A062378 001 Mar 16, 1982

FOR SUSPENSION; ORAL

CEFADROXIL

ANI PHARMS EQ 125MG BASE/5ML A062698 001 Mar 01, 1989

EQ 250MG BASE/5ML A062698 002 Mar 01, 1989

EQ 250MG BASE/5ML A065278 001 Jan 20, 2006

EQ 500MG BASE/5ML A062698 003 Mar 01, 1989

EQ 500MG BASE/5ML A065278 002 Jan 20, 2006

APOTHECON EQ 125MG BASE/5ML A062334 001

EQ 250MG BASE/5ML A062334 002

EQ 500MG BASE/5ML A062334 003

CHARTWELL RX EQ 250MG BASE/5ML A065307 002 Oct 16, 2006

EQ 500MG BASE/5ML A065307 003 Oct 16, 2006

HIKMA PHARMS EQ 250MG BASE/5ML A091036 001 Nov 28, 2012

EQ 500MG BASE/5ML A091036 002 Nov 28, 2012

SUN PHARM INDS LTD EQ 125MG BASE/5ML A065115 001 Mar 26, 2003

EQ 250MG BASE/5ML A065115 002 Mar 26, 2003

EQ 500MG BASE/5ML A065115 003 Mar 26, 2003

DURICEF

+ WARNER CHILCOTT EQ 125MG BASE/5ML ** N050527 002

+ EQ 250MG BASE/5ML ** N050527 003

+ EQ 500MG BASE/5ML ** N050527 001

ULTRACEF

BRISTOL EQ 125MG BASE/5ML A062376 001 Mar 16, 1982

EQ 250MG BASE/5ML A062376 002 Mar 16, 1982

EQ 500MG BASE/5ML A062376 003 Mar 16, 1982

TABLET; ORAL

CEFADROXIL

CHARTWELL RX EQ 1GM BASE A065301 001 Sep 18, 2006

HIKMA EQ 1GM BASE A065260 001 Mar 30, 2006

RANBAXY EQ 1GM BASE A065018 001 Apr 23, 1999

DURICEF

+ WARNER CHILCOTT EQ 1GM BASE ** N050528 001

ULTRACEF

APOTHECON EQ 1GM BASE A062390 001 Jun 10, 1982

BRISTOL EQ 1GM BASE A062408 001 Aug 31, 1982

CEFAMANDOLE NAFATE

INJECTABLE; INJECTION

MANDOL

LILLY EQ 500MG BASE/VIAL N050504 001

EQ 1GM BASE/VIAL A062560 001 Sep 10, 1985

EQ 1GM BASE/VIAL N050504 002

EQ 2GM BASE/VIAL A062560 002 Sep 10, 1985

EQ 2GM BASE/VIAL N050504 003

EQ 10GM BASE/VIAL N050504 004

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

ANCEF

+ GLAXOSMITHKLINE EQ 250MG BASE/VIAL ** N050461 001

+ EQ 500MG BASE/VIAL N050461 002

+ EQ 1GM BASE/VIAL ** N050461 003

+ EQ 5GM BASE/VIAL ** N050461 004

+ EQ 10GM BASE/VIAL ** N050461 005

ANCEF IN DEXTROSE 5% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE EQ 10MG BASE/ML N050566 003 Jun 08, 1983

+ EQ 20MG BASE/ML N050566 004 Jun 08, 1983

ANCEF IN PLASTIC CONTAINER

BAXTER HLTHCARE EQ 10MG BASE/ML A063002 001 Mar 28, 1991

EQ 20MG BASE/ML A063002 002 Mar 28, 1991

ANCEF IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE EQ 10MG BASE/ML N050566 001 Jun 08, 1983

+ EQ 20MG BASE/ML N050566 002 Jun 08, 1983

CEFAZOLIN AND DEXTROSE

B BRAUN EQ 500MG BASE/VIAL N050779 001 Jul 27, 2000

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

ABRAXIS PHARM	EQ 500MG BASE/VIAL	A062688 002	Nov 17, 1986
	EQ 1GM BASE/VIAL	A062688 003	Nov 17, 1986
	EQ 10GM BASE/VIAL	A062688 004	Nov 17, 1986
	EQ 20GM BASE/VIAL	A062688 005	Aug 03, 1987
AUROBINDO PHARMA	EQ 500MG BASE/VIAL	A065395 001	Aug 08, 2008
	EQ 1GM BASE/VIAL	A065395 002	Aug 08, 2008
BEDFORD	EQ 250MG BASE/VIAL	A062894 001	Jul 21, 1988
	EQ 500MG BASE/VIAL	A062894 002	Jul 21, 1988
	EQ 1GM BASE/VIAL	A062894 003	Jul 21, 1988
	EQ 5GM BASE/VIAL	A062894 004	Jul 21, 1988
	EQ 10GM BASE/VIAL	A062894 005	Jul 21, 1988
CEPHAZONE PHARMA	EQ 500MG BASE/VIAL	A065280 001	Mar 18, 2009
	EQ 1GM BASE/VIAL	A065280 002	Mar 18, 2009
	EQ 10GM BASE/VIAL	A065295 001	Mar 18, 2009
	EQ 20GM BASE/VIAL	A065296 001	Mar 18, 2009
DR REDDYS	EQ 250MG BASE/VIAL	A062988 001	Dec 29, 1989
	EQ 500MG BASE/VIAL	A062988 002	Dec 29, 1989
	EQ 1GM BASE/VIAL	A062988 003	Dec 29, 1989
	EQ 5GM BASE/VIAL	A062989 001	Dec 29, 1989
	EQ 10GM BASE/VIAL	A062989 002	Dec 29, 1989
	EQ 20GM BASE/VIAL	A062989 003	Dec 29, 1989
FACTA FARMA	EQ 500MG BASE/VIAL	A063214 001	Dec 27, 1991
	EQ 1GM BASE/VIAL	A063207 001	Dec 27, 1991
	EQ 10GM BASE/VIAL	A063209 001	Dec 27, 1991
	EQ 20GM BASE/VIAL	A063209 002	Apr 30, 1999
FRESENIUS KABI USA	EQ 500MG BASE/VIAL **	A064169 001	Aug 14, 1998
	EQ 1GM BASE/VIAL **	A064169 002	Aug 14, 1998
	EQ 10GM BASE/VIAL **	A064170 001	Mar 18, 1998
	EQ 20GM BASE/VIAL **	A064170 002	Mar 18, 1998
GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	A064033 001	Oct 31, 1993
HIKMA	EQ 250MG BASE/VIAL	A062807 001	Jan 12, 1988
	EQ 500MG BASE/VIAL	A062807 002	Jan 12, 1988
	EQ 1GM BASE/VIAL	A062807 003	Jan 12, 1988
	EQ 5GM BASE/VIAL	A062807 004	Jan 12, 1988
	EQ 10GM BASE/VIAL	A062807 005	Jan 12, 1988
	EQ 20GM BASE/VIAL	A062807 006	Jan 12, 1988
HOSPIRA	EQ 1GM BASE/VIAL	A201654 001	Feb 03, 2016
HOSPIRA INC	EQ 500MG BASE/VIAL	A065226 001	Apr 21, 2005
	EQ 1GM BASE/VIAL	A065226 002	Apr 21, 2005
	EQ 1GM BASE/VIAL	A065244 001	Aug 12, 2005
	EQ 10GM BASE/VIAL	A065247 001	Aug 12, 2005
STERI PHARMA	EQ 500MG BASE/VIAL	A063216 001	Dec 27, 1991
	EQ 1GM BASE/VIAL	A063208 001	Dec 27, 1991
TEVA PHARMS	EQ 250MG BASE/VIAL	A063016 001	Mar 14, 1989
	EQ 500MG BASE/VIAL	A063016 002	Mar 14, 1989
	EQ 1GM BASE/VIAL	A063016 003	Mar 14, 1989
	EQ 5GM BASE/VIAL	A063018 001	Mar 05, 1990
	EQ 10GM BASE/VIAL	A063018 002	Mar 05, 1990
KEFZOL			
ACS DOBFAR	EQ 250MG BASE/VIAL	A061773 001	
	EQ 500MG BASE/VIAL	A061773 002	
	EQ 1GM BASE/VIAL	A061773 003	
	EQ 10GM BASE/VIAL	A061773 004	
	EQ 20GM BASE/VIAL	A061773 005	Sep 08, 1987
LILLY	EQ 500MG BASE/VIAL	A062557 001	Sep 10, 1985
	EQ 1GM BASE/VIAL	A062557 002	Sep 10, 1985

CEFDINIR

CAPSULE; ORAL

OMNICEF

+ ABBVIE

FOR SUSPENSION; ORAL

OMNICEF

+ ABBVIE

+

300MG **	N050739 001	Dec 04, 1997
125MG/5ML **	N050749 001	Dec 04, 1997
250MG/5ML **	N050749 002	Jul 29, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFDITOREN PIVOXILTABLET; ORAL
SPECTRACEF

VANSEN PHARMA	200MG	N021222 001	Aug 29, 2001
	400MG	N021222 002	Jul 21, 2008

CEFEPIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFEPIME HYDROCHLORIDE

FOSUN PHARMA	EQ 500MG BASE/VIAL	A090291 001	Dec 21, 2010
	EQ 1GM BASE/VIAL	A090291 002	Dec 21, 2010
	EQ 2GM BASE/VIAL	A090291 003	Dec 21, 2010
HOSPIRA INC	EQ 500MG BASE/VIAL	A065369 001	Jun 18, 2007
	EQ 1GM BASE/VIAL	A065369 002	Jun 18, 2007
	EQ 1GM BASE/VIAL	A202268 001	Jul 30, 2012
	EQ 2GM BASE/VIAL	A065369 003	Jun 18, 2007
	EQ 2GM BASE/VIAL	A202268 002	Jul 30, 2012

MAXIPIME

+ HOSPIRA INC	EQ 500MG BASE/VIAL	N050679 001	Jan 18, 1996
+	EQ 1GM BASE/VIAL	N050679 002	Jan 18, 1996
+	EQ 2GM BASE/VIAL	N050679 003	Jan 18, 1996

CEFIXIME

FOR SUSPENSION; ORAL

CEFIXIME

SANDOZ	100MG/5ML	A206144 001	Nov 17, 2017
	200MG/5ML	A206144 002	Nov 17, 2017

SUPRAX

+ LEDERLE	100MG/5ML **	N050622 001	Apr 28, 1989
LUPIN PHARMS	100MG/5ML	A065129 001	Feb 23, 2004

TABLET; ORAL

SUPRAX

+ LEDERLE	200MG **	N050621 001	Apr 28, 1989
+	400MG **	N050621 002	Apr 28, 1989
LUPIN PHARMS	400MG	A065130 001	Feb 12, 2004

CEFMENOXIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFMAX

TAP PHARM	EQ 500MG BASE/VIAL	N050571 001	Dec 30, 1987
	EQ 1GM BASE/VIAL	N050571 002	Dec 30, 1987
	EQ 2GM BASE/VIAL	N050571 003	Dec 30, 1987

CEFMETAZOLE SODIUM

INJECTABLE; INJECTION

ZEFAZONE

+ PHARMACIA AND UPJOHN	EQ 1GM BASE/VIAL **	N050637 001	Dec 11, 1989
+	EQ 2GM BASE/VIAL **	N050637 002	Dec 11, 1989

ZEFAZONE IN PLASTIC CONTAINER

+ PHARMACIA AND UPJOHN	EQ 20MG BASE/ML **	N050683 001	Dec 29, 1992
+	EQ 40MG BASE/ML **	N050683 002	Dec 29, 1992

CEFONICID SODIUM

INJECTABLE; INJECTION

MONOCID

GLAXOSMITHKLINE	EQ 500MG BASE/VIAL	N050579 001	May 23, 1984
	EQ 1GM BASE/VIAL	A063295 001	Jul 26, 1993
	EQ 1GM BASE/VIAL	N050579 002	May 23, 1984
	EQ 2GM BASE/VIAL	N050579 003	May 23, 1984
	EQ 10GM BASE/VIAL	N050579 004	May 23, 1984

CEFOPERAZONE SODIUM

INJECTABLE; INJECTION

CEFOBID

PFIZER	EQ 1GM BASE/VIAL	A063333 001	Mar 31, 1995
	EQ 1GM BASE/VIAL	N050551 001	Nov 18, 1982
	EQ 2GM BASE/VIAL	A063333 002	Mar 31, 1995
	EQ 2GM BASE/VIAL	N050551 002	Nov 18, 1982
	EQ 10GM BASE/VIAL	N050551 003	Mar 05, 1990

CEFOBID IN PLASTIC CONTAINER

PFIZER	EQ 20MG BASE/ML	N050613 002	Jul 31, 1987
	EQ 40MG BASE/ML	N050613 001	Jul 23, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFORANIDE

INJECTABLE; INJECTION

PRECEF

APOTHECON	500MG/VIAL	A062579 001	Nov 26, 1984
	1GM/VIAL	A062579 002	Nov 26, 1984
	2GM/VIAL	A062579 003	Nov 26, 1984
	10GM/VIAL	A062579 004	Nov 26, 1984
	20GM/VIAL	A062579 005	Nov 26, 1984
BRISTOL	500MG/VIAL	N050554 001	May 24, 1984
	1GM/VIAL	N050554 002	May 24, 1984
	2GM/VIAL	N050554 003	May 24, 1984
	10GM/VIAL	N050554 004	May 24, 1984
	20GM/VIAL	N050554 005	May 24, 1984

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME

FRESENIUS KABI USA	EQ 500MG BASE/VIAL	A064200 001	Mar 24, 2000
	EQ 1GM BASE/VIAL	A064200 002	Mar 24, 2000
	EQ 2GM BASE/VIAL	A064200 003	Mar 24, 2000
	EQ 10GM BASE/VIAL	A064201 001	Mar 24, 2000
	EQ 20GM BASE/VIAL	A064201 002	Mar 24, 2000
HIKMA	EQ 500MG BASE/VIAL	A065072 001	Nov 20, 2002
	EQ 1GM BASE/VIAL	A065072 002	Nov 20, 2002
	EQ 2GM BASE/VIAL	A065072 003	Nov 20, 2002
	EQ 10GM BASE/VIAL	A065071 001	Nov 20, 2002
WOCKHARDT	EQ 1GM BASE/VIAL	A065197 001	Aug 29, 2006
CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER			
B BRAUN	EQ 2GM BASE	N050792 001	Jul 29, 2004
CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER			
B BRAUN	EQ 1GM BASE	N050792 002	Jul 29, 2004
CEFOTAXIME SODIUM			
AUROBINDO PHARMA	EQ 500MG BASE/VIAL	A065517 001	Nov 06, 2009
	EQ 1GM BASE/VIAL	A065517 002	Nov 06, 2009
	EQ 2GM BASE/VIAL	A065517 003	Nov 06, 2009
AUROBINDO PHARMA LTD	EQ 10GM BASE/VIAL	A065516 001	Nov 06, 2009
CEPHAZONE PHARMA	EQ 10GM BASE/VIAL	A065348 001	Jan 25, 2010
HOSPIRA	EQ 1GM BASE/VIAL	A203132 001	Feb 19, 2016
	EQ 2GM BASE/VIAL	A203132 002	Feb 19, 2016
HOSPIRA INC	EQ 500MG BASE/VIAL	A065290 001	Aug 11, 2006
	EQ 1GM BASE/VIAL	A065290 002	Aug 11, 2006
	EQ 1GM BASE/VIAL	A065293 001	Aug 10, 2006
	EQ 2GM BASE/VIAL	A065290 003	Aug 11, 2006
	EQ 2GM BASE/VIAL	A065293 002	Aug 10, 2006
	EQ 10GM BASE/VIAL	A065292 001	Aug 10, 2006
LUPIN	EQ 500MG BASE/VIAL	A065124 001	Sep 24, 2003
	EQ 1GM BASE/VIAL	A065124 002	Sep 24, 2003
	EQ 2GM BASE/VIAL	A065124 003	Sep 24, 2003
WOCKHARDT	EQ 500MG BASE/VIAL	A065197 002	Jun 20, 2008
	EQ 2GM BASE/VIAL	A065197 003	Jun 20, 2008
CLAFORAN			
SANOFI AVENTIS US	EQ 1GM BASE/VIAL	A062659 001	Jan 13, 1987
	EQ 2GM BASE/VIAL	A062659 002	Jan 13, 1987
+ VALIDUS PHARMS	EQ 500MG BASE/VIAL **	N050547 001	
+	EQ 1GM BASE/VIAL **	N050547 002	
+	EQ 2GM BASE/VIAL **	N050547 003	
+	EQ 10GM BASE/VIAL **	N050547 004	Dec 29, 1983
CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER			
VALIDUS PHARMS	EQ 20MG BASE/ML	N050596 002	May 20, 1985
	EQ 40MG BASE/ML	N050596 004	May 20, 1985
CLAFORAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
VALIDUS PHARMS	EQ 20MG BASE/ML	N050596 001	May 20, 1985
	EQ 40MG BASE/ML	N050596 003	May 20, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

+	PAI HOLDINGS PHARM	EQ 1GM BASE/VIAL **	N050588	001	Dec 27, 1985
+		EQ 2GM BASE/VIAL **	N050588	002	Dec 27, 1985
		EQ 10GM BASE/VIAL	N050588	003	Apr 25, 1988
	TELIGENT	EQ 1GM BASE/VIAL	A063293	001	Apr 29, 1993
		EQ 2GM BASE/VIAL	A063293	002	Apr 29, 1993

CEFOTAN IN PLASTIC CONTAINER

	PAI HOLDINGS PHARM	EQ 20MG BASE/ML	N050694	002	Jul 30, 1993
		EQ 40MG BASE/ML	N050694	001	Jul 30, 1993

CEFOTETAN

	FRESENIUS KABI USA	EQ 10GM BASE/VIAL	A065375	001	Aug 09, 2007
	HIKMA	EQ 1GM BASE/VIAL	A091031	001	Oct 26, 2011
		EQ 2GM BASE/VIAL	A091031	002	Oct 26, 2011
	WEST-WARD PHARM CORP	EQ 10GM BASE/VIAL	A091030	001	Oct 26, 2011

CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER

+	B BRAUN	EQ 1GM BASE/VIAL	N065430	001	Aug 09, 2007
+		EQ 2GM BASE/VIAL	N065430	002	Aug 09, 2007

CEFOTIAM HYDROCHLORIDE

INJECTABLE; INJECTION

CERADON

	TAKEDA	EQ 1GM BASE/VIAL	N050601	001	Dec 30, 1988
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CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

	ACS DOBFAR SPA	EQ 1GM BASE/VIAL	A065467	001	Aug 31, 2011
		EQ 2GM BASE/VIAL	A065467	002	Aug 31, 2011
		EQ 10GM BASE/VIAL	A065464	001	Aug 31, 2011
	FRESENIUS KABI USA	EQ 1GM BASE/VIAL **	A065012	001	Jul 03, 2000
		EQ 2GM BASE/VIAL **	A065012	002	Jul 03, 2000
		EQ 10GM BASE/VIAL	A065011	001	Jul 03, 2000
	HOSPIRA INC	EQ 1GM BASE/VIAL	A065313	001	Jan 23, 2006
		EQ 2GM BASE/VIAL	A065313	002	Jan 23, 2006
		EQ 10GM BASE/VIAL	A065312	001	Feb 13, 2006

MEFOXIN

	MYLAN INSTITUTIONAL	EQ 1GM BASE/VIAL	A062757	001	Jan 08, 1987
+		EQ 1GM BASE/VIAL **	N050517	001	
		EQ 2GM BASE/VIAL	A062757	002	Jan 08, 1987
+		EQ 2GM BASE/VIAL **	N050517	002	
+		EQ 10GM BASE/VIAL **	N050517	003	

MEFOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

+	MERCK	EQ 20MG BASE/ML **	N050581	003	Sep 20, 1984
+		EQ 40MG BASE/ML **	N050581	004	Sep 20, 1984

MEFOXIN IN PLASTIC CONTAINER

	MYLAN INSTITUTIONAL	EQ 20MG BASE/ML	A063182	001	Jan 25, 1993
		EQ 40MG BASE/ML	A063182	002	Jan 25, 1993

MEFOXIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+	MERCK	EQ 20MG BASE/ML **	N050581	002	Sep 20, 1984
+		EQ 40MG BASE/ML **	N050581	001	Sep 20, 1984

CEFPYRAMIDE SODIUM

INJECTABLE; INJECTION

CEFPYRAMIDE SODIUM

	WYETH AYERST	EQ 1GM BASE/VIAL	N050633	002	Jan 31, 1989
		EQ 2GM BASE/VIAL	N050633	003	Jan 31, 1989
		EQ 10GM BASE/VIAL	N050633	005	Jan 31, 1989

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

BANAN

	SANKYO	EQ 50MG BASE/5ML	N050688	002	Aug 07, 1992
		EQ 100MG BASE/5ML	N050688	001	Aug 07, 1992

CEFPODOXIME PROXETIL

	SANDOZ	EQ 50MG BASE/5ML	A090031	001	Jan 14, 2009
		EQ 100MG BASE/5ML	A090031	002	Jan 14, 2009
	SUN PHARM INDS LTD	EQ 50MG BASE/5ML	A065082	001	May 31, 2002
		EQ 100MG BASE/5ML	A065082	002	May 31, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFPODOXIME PROXETIL

FOR SUSPENSION;ORAL

VANTIN

+	PHARMACIA AND UPJOHN	EQ 50MG BASE/5ML **	N050675	001	Aug 07, 1992
+		EQ 100MG BASE/5ML **	N050675	002	Aug 07, 1992

TABLET;ORAL

BANAN

	SANKYO	EQ 100MG BASE	N050687	001	Aug 07, 1992
		EQ 200MG BASE	N050687	002	Aug 07, 1992

CEFPODOXIME PROXETIL

	SUN PHARM INDS LTD	EQ 100MG BASE	A065083	001	Aug 20, 2003
		EQ 200MG BASE	A065083	002	Aug 20, 2003

VANTIN

+	PHARMACIA AND UPJOHN	EQ 100MG BASE **	N050674	001	Aug 07, 1992
+		EQ 200MG BASE **	N050674	002	Aug 07, 1992

CEFPROZIL

FOR SUSPENSION;ORAL

CEFPROZIL

	CHARTWELL RX	125MG/5ML	A065236	001	Dec 08, 2005
		250MG/5ML	A065236	002	Dec 08, 2005
	ORCHID HLTHCARE	125MG/5ML	A065284	002	Dec 30, 2005
		250MG/5ML	A065284	001	Dec 30, 2005
	RANBAXY LABS LTD	125MG/5ML	A065202	001	Jun 30, 2006
		250MG/5ML	A065202	002	Jun 30, 2006
	SANDOZ	125MG/5ML	A065257	001	Dec 08, 2005
		250MG/5ML	A065257	002	Dec 08, 2005

CEFZIL

+	CORDEN PHARMA	125MG/5ML **	N050665	001	Dec 23, 1991
+		250MG/5ML **	N050665	002	Dec 23, 1991

TABLET;ORAL

CEFPROZIL

	ORCHID HLTHCARE	250MG	A065267	001	Dec 19, 2005
		500MG	A065267	002	Dec 19, 2005
	RANBAXY LABS LTD	250MG	A065198	001	Dec 13, 2006
		500MG	A065198	002	Dec 13, 2006
	WOCKHARDT	250MG	A065428	001	Jun 14, 2007
		500MG	A065428	002	Jun 14, 2007

CEFZIL

+	CORDEN PHARMA	250MG **	N050664	001	Dec 23, 1991
+		500MG **	N050664	002	Dec 23, 1991

CEFTAROLINE FOSAMIL

POWDER;INTRAVENOUS

CEFTAROLINE FOSAMIL

	APOTEX	400MG/VIAL	A208075	001	Sep 21, 2021
		600MG/VIAL	A208075	002	Sep 21, 2021

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

	ACS DOBFAR	500MG/VIAL	A062640	001	Nov 20, 1985
	AUROBINDO PHARMA LTD	500MG/VIAL	A065481	001	May 28, 2010
		1GM/VIAL	A065481	002	May 28, 2010
		2GM/VIAL	A065481	003	May 28, 2010
		6GM/VIAL	A065482	001	May 28, 2010
	WOCKHARDT	1GM/VIAL	A065196	001	Oct 15, 2008

CEFTAZIDIME IN DEXTROSE CONTAINER

+	B BRAUN	EQ 1GM BASE	N050823	001	Jun 13, 2011
+		EQ 2GM BASE	N050823	002	Jun 13, 2011

CEPTAZ

	GLAXOSMITHKLINE	500MG/VIAL	N050646	001	Sep 27, 1990
		1GM/VIAL	N050646	002	Sep 27, 1990
		2GM/VIAL	N050646	003	Sep 27, 1990
		10GM/VIAL	N050646	004	Sep 27, 1990

FORTAZ

+	PAI HOLDINGS PHARM	500MG/VIAL	N050578	001	Jul 19, 1985
+		1GM/VIAL	N050578	002	Jul 19, 1985
+		2GM/VIAL	N050578	003	Jul 19, 1985
+		6GM/VIAL	N050578	004	Jul 19, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTAZIDIME

INJECTABLE; INJECTION

PENTACEF

GLAXOSMITHKLINE	1GM/VIAL	A063322 001	Nov 07, 1995
	1GM/VIAL	A064006 001	Mar 31, 1992
	2GM/VIAL	A063322 002	Nov 07, 1995
	2GM/VIAL	A064006 002	Mar 31, 1992
	6GM/VIAL	A064008 001	Mar 31, 1992
	10GM/VIAL	A064008 002	Mar 31, 1992

TAZICEF

HOSPIRA	500MG/VIAL	A062662 001	Mar 06, 1986
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TAZIDIME

LILLY	1GM/VIAL	A062655 001	Nov 20, 1985
	2GM/VIAL	A062655 002	Nov 20, 1985

TAZIDIME IN PLASTIC CONTAINER

LILLY	1GM/VIAL	A062739 001	Jul 10, 1986
	2GM/VIAL	A062739 002	Jul 10, 1986

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

CEFTAZIDIME SODIUM IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 10MG BASE/ML	A063221 001	Apr 29, 1993
	EQ 20MG BASE/ML	A063221 002	Apr 29, 1993
	EQ 40MG BASE/ML	A063221 003	Apr 29, 1993

FORTAZ IN PLASTIC CONTAINER

PAI HOLDINGS PHARM	EQ 10MG BASE/ML	N050634 001	Apr 28, 1989
+	EQ 20MG BASE/ML	N050634 002	Apr 28, 1989
+	EQ 40MG BASE/ML	N050634 003	Apr 28, 1989

CEFTIBUTEN DIHYDRATE

CAPSULE; ORAL

CEDAX

SI PHARMS	EQ 400MG BASE	N050685 002	Dec 20, 1995
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FOR SUSPENSION; ORAL

CEDAX

+	SI PHARMS	EQ 90MG BASE/5ML **	N050686 001	Dec 20, 1995
+		EQ 180MG BASE/5ML **	N050686 002	Dec 20, 1995

CEFTIZOXIME SODIUM

INJECTABLE; INJECTION

CEFIZOX

ASTELLAS	EQ 500MG BASE/VIAL	N050560 001	Sep 15, 1983
	EQ 1GM BASE/VIAL	A063294 002	Mar 31, 1994
	EQ 1GM BASE/VIAL	N050560 002	Sep 15, 1983
	EQ 2GM BASE/VIAL	A063294 003	Mar 31, 1994
	EQ 2GM BASE/VIAL	N050560 003	Sep 15, 1983
	EQ 10GM BASE/VIAL	N050560 005	Mar 19, 1993

CEFIZOX IN DEXTROSE 5% IN PLASTIC CONTAINER

ASTELLAS	EQ 20MG BASE/ML	N050589 001	Oct 03, 1984
	EQ 40MG BASE/ML	N050589 002	Oct 03, 1984

CEFIZOX IN PLASTIC CONTAINER

ASTELLAS	EQ 20MG BASE/ML	N050589 003	Apr 13, 1995
	EQ 40MG BASE/ML	N050589 004	Apr 13, 1995

CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE

AGILA SPECLTS	EQ 10GM BASE/VIAL	A091068 001	Jan 07, 2013
AUROBINDO PHARMA LTD	EQ 10GM BASE/VIAL	A065504 001	Jul 31, 2008
BEDFORD	EQ 10GM BASE/VIAL	A065475 001	Aug 18, 2008
FACTA FARMA	EQ 10GM BASE/VIAL	A065269 001	Feb 28, 2007
FRESENIUS KABI USA	EQ 10GM BASE/VIAL	A065252 001	Feb 15, 2006
HOSPIRA INC	EQ 1GM BASE/VIAL	A065231 001	Aug 02, 2005
	EQ 1GM BASE/VIAL	A202563 001	Aug 20, 2012
	EQ 2GM BASE/VIAL	A065231 002	Aug 02, 2005
	EQ 2GM BASE/VIAL	A202563 002	Aug 20, 2012
	EQ 10GM BASE/VIAL	A065232 001	Aug 02, 2005
LUPIN	EQ 10GM BASE/VIAL	A065263 001	Sep 12, 2006
TEVA	EQ 10GM BASE/VIAL	A065274 001	May 01, 2006

ROCEPHIN

HOFFMANN LA ROCHE	EQ 250MG BASE/VIAL	A063239 001	Aug 13, 1993
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

ROCEPHIN

	EQ 500MG BASE/VIAL	A062654	001	Apr 30, 1987	
	EQ 500MG BASE/VIAL	A063239	002	Aug 13, 1993	
	EQ 1GM BASE/VIAL	A062654	002	Apr 30, 1987	
	EQ 1GM BASE/VIAL	A063239	003	Aug 13, 1993	
	EQ 2GM BASE/VIAL	A062654	003	Apr 30, 1987	
+	EQ 10GM BASE/VIAL **	N050585	005	Dec 21, 1984	
ROCHE	EQ 250MG BASE/VIAL	A062510	001	Mar 12, 1985	
	EQ 500MG BASE/VIAL	A062510	002	Mar 12, 1985	
	EQ 1GM BASE/VIAL	A062510	003	Mar 12, 1985	
ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER					
+	HOFFMANN LA ROCHE	EQ 10MG BASE/ML **	N050624	001	Feb 11, 1987
+		EQ 20MG BASE/ML **	N050624	002	Feb 11, 1987
+		EQ 40MG BASE/ML **	N050624	003	Feb 11, 1987

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAXONE

AUROBINDO PHARMA LTD	EQ 250MG BASE/VIAL	A065505	001	Jul 31, 2008	
	EQ 500MG BASE/VIAL	A065505	002	Jul 31, 2008	
	EQ 1GM BASE/VIAL	A065505	003	Jul 31, 2008	
	EQ 2GM BASE/VIAL	A065505	004	Jul 31, 2008	
BEDFORD	EQ 250MG BASE/VIAL	A065465	001	Aug 18, 2008	
	EQ 500MG BASE/VIAL	A065465	002	Aug 18, 2008	
	EQ 1GM BASE/VIAL	A065465	003	Aug 18, 2008	
	EQ 2GM BASE/VIAL	A065465	004	Aug 18, 2008	
CEPHAZONE PHARMA	EQ 250MG BASE/VIAL	A065294	001	Mar 26, 2007	
	EQ 500MG BASE/VIAL	A065294	002	Mar 26, 2007	
	EQ 1GM BASE/VIAL	A065294	003	Mar 26, 2007	
	EQ 2GM BASE/VIAL	A065294	004	Mar 26, 2007	
FACTA FARMA	EQ 1GM BASE/VIAL	A065268	001	Feb 28, 2007	
	EQ 2GM BASE/VIAL	A065268	002	Feb 28, 2007	
FRESENIUS KABI USA	EQ 250MG BASE/VIAL	A065245	001	Feb 15, 2006	
	EQ 500MG BASE/VIAL	A065245	002	Feb 15, 2006	
	EQ 1GM BASE/VIAL	A065245	003	Feb 15, 2006	
	EQ 2GM BASE/VIAL	A065245	004	Feb 15, 2006	
HOSPIRA INC	EQ 250MG BASE/VIAL	A065230	001	Aug 02, 2005	
	EQ 500MG BASE/VIAL	A065230	002	Aug 02, 2005	
	EQ 1GM BASE/VIAL	A065230	003	Aug 02, 2005	
	EQ 2GM BASE/VIAL	A065230	004	Aug 02, 2005	
LUPIN	EQ 250MG BASE/VIAL	A065125	001	Sep 30, 2003	
	EQ 500MG BASE/VIAL	A065125	002	Sep 30, 2003	
	EQ 1GM BASE/VIAL	A065125	003	Sep 30, 2003	
	EQ 2GM BASE/VIAL	A065125	004	Sep 30, 2003	
TEVA	EQ 1GM BASE/VIAL	A065262	001	Jun 29, 2006	
	EQ 2GM BASE/VIAL	A065262	002	Jun 29, 2006	
TEVA PHARMS USA	EQ 250MG BASE/VIAL	A065227	001	Mar 15, 2007	
	EQ 500MG BASE/VIAL	A065227	002	Mar 15, 2007	
	EQ 1GM BASE/VIAL	A065227	003	Mar 15, 2007	
	EQ 2GM BASE/VIAL	A065227	004	Mar 15, 2007	
ROCEPHIN					
+	HOFFMANN LA ROCHE	EQ 250MG BASE/VIAL **	N050585	001	Dec 21, 1984
+		EQ 500MG BASE/VIAL **	N050585	002	Dec 21, 1984
+		EQ 1GM BASE/VIAL **	N050585	003	Dec 21, 1984
+		EQ 2GM BASE/VIAL **	N050585	004	Dec 21, 1984

CEFTRIAXONE SODIUM; LIDOCAINE

INJECTABLE; INJECTION

ROCEPHIN KIT

HOFFMANN LA ROCHE

EQ 500MG BASE/VIAL, N/A; N/A, 1%	N050585	007	May 08, 1996
EQ 1GM BASE/VIAL, N/A; N/A, 1%	N050585	006	May 08, 1996

CEFUROXIME AXETIL

FOR SUSPENSION; ORAL

CEFTIN

+	GLAXOSMITHKLINE	EQ 125MG BASE/5ML	N050672	001	Jun 30, 1994
+		EQ 250MG BASE/5ML	N050672	002	Apr 29, 1997

CEFUROXIME AXETIL

SUN PHARM INDS LTD

EQ 125MG BASE/5ML	A065323	001	Feb 05, 2008
EQ 250MG BASE/5ML	A065323	002	Feb 05, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFUROXIME AXETIL

TABLET; ORAL

CEFTIN

+	GLAXOSMITHKLINE	EQ 125MG BASE **	N050605 001	Dec 28, 1987
+		EQ 250MG BASE **	N050605 002	Dec 28, 1987
+		EQ 500MG BASE **	N050605 003	Dec 28, 1987

CEFUROXIME AXETIL

	ANI PHARMS	EQ 250MG BASE	A065190 001	Oct 18, 2004
		EQ 500MG BASE	A065190 002	Oct 18, 2004
	FOSUN PHARMA	EQ 250MG BASE	A065126 001	Oct 28, 2003
		EQ 500MG BASE	A065126 002	Oct 28, 2003
	RANBAXY LABS LTD	EQ 125MG BASE	A065043 003	Feb 15, 2002
		EQ 250MG BASE	A065043 002	Feb 15, 2002
		EQ 500MG BASE	A065043 001	Feb 15, 2002
	SUN PHARM INDS LTD	EQ 125MG BASE	A065118 001	Apr 25, 2003
		EQ 250MG BASE	A065118 002	Apr 25, 2003
		EQ 500MG BASE	A065118 003	Apr 25, 2003

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER

+	B BRAUN	EQ 750MG BASE/VIAL	N050780 001	Feb 21, 2001
+		EQ 1.5GM BASE/VIAL	N050780 002	Feb 21, 2001

CEFUROXIME SODIUM

	ACS DOBFAR SPA	EQ 7.5GM BASE/VIAL	A064124 001	May 30, 1997
	FRESENIUS KABI USA	EQ 1.5GM BASE/VIAL	A065001 002	May 30, 2001
		EQ 7.5GM BASE/VIAL	A065002 001	Sep 28, 1998
	HIKMA	EQ 7.5GM BASE/VIAL	A065046 001	Jan 09, 2004
	HOSPIRA INC	EQ 1.5GM BASE/VIAL	A065483 002	Oct 15, 2008
		EQ 1.5GM BASE/VIAL	A065503 001	Oct 15, 2008
		EQ 7.5GM BASE/VIAL	A065484 001	Oct 15, 2008
	TEVA PHARMS	EQ 7.5GM BASE/VIAL	A064191 001	Apr 16, 1998
	WATSON LABS INC	EQ 1.5GM BASE/VIAL	A064035 002	Feb 26, 1993
		EQ 7.5GM BASE/VIAL	A064036 001	Feb 26, 1993

CEFUROXIME SODIUM IN PLASTIC CONTAINER

	SAMSON MEDCL	EQ 75GM BASE/VIAL	A065251 001	Dec 30, 2009
		EQ 225GM BASE/VIAL	A065251 002	Dec 30, 2009

KEFUROX

	ACS DOBFAR	EQ 1.5GM BASE/VIAL	A062591 002	Jan 10, 1986
		EQ 7.5GM BASE/VIAL	A062591 003	Dec 17, 1987
	LILLY	EQ 1.5GM BASE/VIAL	A062592 002	Jan 10, 1986

KEFUROX IN PLASTIC CONTAINER

	LILLY	EQ 1.5GM BASE/VIAL	A062590 002	Jan 10, 1986
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ZINACEF

+	PAI HOLDINGS PHARM	EQ 1.5GM BASE/VIAL	N050558 003	Oct 19, 1983
+		EQ 7.5GM BASE/VIAL	N050558 004	Oct 23, 1986

ZINACEF IN PLASTIC CONTAINER

	PAI HOLDINGS PHARM	EQ 15MG BASE/ML	N050643 001	Apr 28, 1989
+		EQ 30MG BASE/ML	N050643 002	Apr 28, 1989

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

	FRESENIUS KABI USA	EQ 750MG BASE/VIAL	A065001 001	May 30, 2001
	HOSPIRA INC	EQ 750MG BASE/VIAL	A065483 001	Oct 15, 2008
	TEVA PHARMS	EQ 750MG BASE/VIAL	A064192 002	Apr 16, 1998
		EQ 1.5GM BASE/VIAL	A064192 001	Apr 16, 1998
	WATSON LABS INC	EQ 750MG BASE/VIAL	A064035 001	Feb 26, 1993

KEFUROX

	ACS DOBFAR	EQ 750MG BASE/VIAL	A062591 001	Jan 10, 1986
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ZINACEF

+	PAI HOLDINGS PHARM	EQ 750MG BASE/VIAL	N050558 002	Oct 19, 1983
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INJECTABLE; INTRAVENOUS

KEFUROX

	LILLY	EQ 750MG BASE/VIAL	A062592 001	Jan 10, 1986
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KEFUROX IN PLASTIC CONTAINER

	LILLY	EQ 750MG BASE/VIAL	A062590 001	Jan 10, 1986
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CELECOXIB

CAPSULE; ORAL

CELECOXIB

ACIC PHARMS	200MG	A212925 001	Dec 09, 2020
	400MG	A212925 002	Dec 09, 2020
JUBILANT GENERICS	50MG	A207061 001	Apr 04, 2017
	100MG	A207061 002	Apr 04, 2017
	200MG	A207061 003	Apr 04, 2017
	400MG	A207061 004	Apr 04, 2017
MYLAN	50MG	A078857 001	May 30, 2014
	100MG	A078857 002	Feb 11, 2015
	200MG	A078857 003	Feb 11, 2015
	400MG	A078857 004	Feb 11, 2015
UNICHEM	50MG	A213301 001	Jan 12, 2021
	100MG	A213301 002	Jan 12, 2021
	200MG	A213301 003	Jan 12, 2021
	400MG	A213301 004	Jan 12, 2021
YABAO PHARM	100MG	A212564 001	Apr 10, 2023
	200MG	A212564 002	Apr 10, 2023
	400MG	A212564 003	Apr 10, 2023

CELLULOSE SODIUM PHOSPHATE

POWDER; ORAL

CALCIBIND

MISSION PHARMA	2.5GM/PACKET	N018757 002	Dec 28, 1982
	300GM/BOT	N018757 003	Oct 16, 1984

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

APOTHECON	EQ 250MG BASE	A062973 001	Nov 08, 1988
	EQ 250MG BASE	A063063 001	Sep 29, 1989
	EQ 250MG BASE	A063186 001	Dec 30, 1994
	EQ 500MG BASE	A062974 001	Nov 23, 1988
	EQ 500MG BASE	A063063 002	Sep 29, 1989
	EQ 500MG BASE	A063186 002	Dec 30, 1994
BARR	EQ 250MG BASE	A062773 001	Jun 26, 1987
	EQ 500MG BASE	A062775 001	Apr 22, 1987
FACTA FARMA	EQ 250MG BASE	A062118 001	
	EQ 500MG BASE	A062118 002	
HIKMA	EQ 250MG BASE	A065215 001	Jan 24, 2006
	EQ 500MG BASE	A065215 002	Jan 24, 2006
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A061969 001	
	EQ 500MG BASE	A061969 002	
PUREPAC PHARM	EQ 250MG BASE	A062809 001	Apr 22, 1987
	EQ 500MG BASE	A062809 002	Apr 22, 1987
STEVENS J	EQ 250MG BASE	A062870 001	Mar 17, 1988
	EQ 500MG BASE	A062869 001	Mar 17, 1988
SUN PHARM INDS (IN)	EQ 250MG BASE	A062791 001	Jun 11, 1987
	EQ 500MG BASE	A062791 002	Jun 11, 1987
SUN PHARM INDS LTD	EQ 250MG BASE	A065007 001	Sep 16, 1999
	EQ 500MG BASE	A065007 002	Sep 16, 1999
TEVA	EQ 250MG BASE	A062760 001	Apr 24, 1987
	EQ 250MG BASE	A062821 001	Feb 05, 1988
	EQ 500MG BASE	A062761 001	Apr 24, 1987
	EQ 500MG BASE	A062823 001	Feb 05, 1988
YOSHITOMI	EQ 250MG BASE	A062872 001	Jun 20, 1988
	EQ 500MG BASE	A062871 001	Jul 05, 1988
KEFLEX			
+	PRAGMA	EQ 250MG BASE **	N050405 002
+		EQ 333MG BASE **	N050405 004
+		EQ 500MG BASE **	N050405 003
+		EQ 750MG BASE **	N050405 005

FOR SUSPENSION; ORAL

CEPHALEXIN

APOTHECON	EQ 125MG BASE/5ML	A062986 001	Apr 18, 1991
	EQ 250MG BASE/5ML	A062987 001	Jul 25, 1989
BARR	EQ 125MG BASE/5ML	A062778 001	Aug 06, 1987
	EQ 250MG BASE/5ML	A062777 001	Aug 06, 1987
CHARTWELL RX	EQ 125MG BASE/5ML	A065326 001	Jul 10, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHALEXINFOR SUSPENSION;ORAL
CEPHALEXIN

	EQ 250MG BASE/5ML	A065326 002	Jul 10, 2006
FACTA FARMA	EQ 100MG BASE/ML **	A062117 001	
	EQ 125MG BASE/5ML **	A062117 002	
	EQ 250MG BASE/5ML **	A062117 003	
HIKMA PHARMS	EQ 125MG BASE/5ML	A065444 001	Aug 28, 2009
	EQ 250MG BASE/5ML	A065444 002	Aug 28, 2009
SUN PHARM INDS LTD	EQ 125MG BASE/5ML	A065081 001	Jul 27, 2001
	EQ 250MG BASE/5ML	A065081 002	Jul 27, 2001
TEVA	EQ 125MG BASE/5ML	A062767 001	Jun 16, 1987
	EQ 125MG BASE/5ML	A062873 001	May 23, 1988
	EQ 250MG BASE/5ML	A062768 001	Jun 16, 1987
	EQ 250MG BASE/5ML	A062867 001	Apr 15, 1988
VITARINE	EQ 125MG BASE/5ML	A062779 001	Dec 22, 1987
	EQ 250MG BASE/5ML	A062781 001	Dec 22, 1987
KEFLEX			
+ PRAGMA	EQ 100MG BASE/ML **	N050406 003	
+	EQ 125MG BASE/5ML **	N050406 001	
+	EQ 250MG BASE/5ML **	N050406 002	

TABLET;ORAL

CEPHALEXIN

BARR	EQ 250MG BASE	A062826 001	Aug 17, 1987
	EQ 500MG BASE	A062827 001	Aug 17, 1987
VITARINE	EQ 250MG BASE	A062863 001	Aug 11, 1988
	EQ 500MG BASE	A062863 002	Aug 11, 1988
	EQ 1GM BASE	A062863 003	Aug 11, 1988

KEFLET

LILLY	EQ 250MG BASE	A062745 001	Dec 01, 1986
	EQ 250MG BASE	N050440 003	Feb 26, 1987
	EQ 500MG BASE	A062745 002	Dec 01, 1986
	EQ 500MG BASE	N050440 001	
	EQ 1GM BASE	N050440 002	

TABLET, FOR SUSPENSION;ORAL

PANIXINE DISPERDOSE

RANBAXY LABS LTD	EQ 125MG BASE	A065100 002	Sep 11, 2003
	EQ 250MG BASE	A065100 001	Sep 11, 2003

CEPHALEXIN HYDROCHLORIDE

TABLET;ORAL

KEFTAB

LILLY	EQ 250MG BASE	N050614 001	Oct 29, 1987
	EQ 333MG BASE	N050614 003	May 16, 1988
	EQ 500MG BASE	N050614 002	Oct 29, 1987

CEPHALOGLYCIN

CAPSULE;ORAL

KAFOCIN

LILLY	250MG	N050219 001	
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CEPHALOTHIN SODIUM

INJECTABLE;INJECTION

CEPHALOTHIN

INTL MEDICATION	EQ 500MG BASE/VIAL	A062426 001	May 03, 1985
	EQ 1GM BASE/VIAL	A062426 002	May 03, 1985
	EQ 2GM BASE/VIAL	A062426 003	May 03, 1985
	EQ 4GM BASE/VIAL	A062426 004	May 03, 1985

CEPHALOTHIN SODIUM

ABBOTT	EQ 1GM BASE/VIAL	A062547 001	Sep 11, 1985
	EQ 1GM BASE/VIAL	A062548 001	Sep 11, 1985
	EQ 2GM BASE/VIAL	A062547 002	Sep 11, 1985
	EQ 2GM BASE/VIAL	A062548 002	Sep 11, 1985
ABRAXIS PHARM	EQ 1GM BASE/VIAL	A062666 002	Jun 10, 1987
	EQ 2GM BASE/VIAL	A062666 001	Jun 10, 1987
BRISTOL	EQ 1GM BASE/VIAL	A062464 001	May 07, 1984
	EQ 2GM BASE/VIAL	A062464 002	May 07, 1984
	EQ 4GM BASE/VIAL	A062464 003	May 07, 1984

CEPHALOTHIN SODIUM W/ DEXTROSE IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 20MG BASE/ML	A062422 003	Jan 31, 1984
	EQ 20MG BASE/ML	A062422 005	Jul 16, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION

CEPHALOTHIN SODIUM W/ DEXTROSE IN PLASTIC CONTAINER

EQ 20MG BASE/ML	A062730 001	Mar 05, 1987
EQ 40MG BASE/ML	A062422 004	Jan 31, 1984
EQ 40MG BASE/ML	A062422 006	Jul 16, 1991
EQ 40MG BASE/ML	A062730 002	Mar 05, 1987

CEPHALOTHIN SODIUM W/ SODIUM CHLORIDE IN PLASTIC CONTAINER

BAXTER HLTHCARE EQ 20MG BASE/ML	A062422 001	Jan 31, 1984
EQ 40MG BASE/ML	A062422 002	Jan 31, 1984

KEFLIN

LILLY EQ 1GM BASE/VIAL	N050482 001	
EQ 2GM BASE/VIAL	N050482 002	
EQ 4GM BASE/VIAL	N050482 003	
EQ 20GM BASE/VIAL	N050482 007	

KEFLIN IN PLASTIC CONTAINER

LILLY EQ 1GM BASE/VIAL	A062549 001	Sep 10, 1985
EQ 2GM BASE/VIAL	A062549 002	Sep 10, 1985

SEFFIN

GLAXOSMITHKLINE EQ 1GM BASE/VIAL	A062435 001	Nov 15, 1983
EQ 2GM BASE/VIAL	A062435 002	Nov 15, 1983
EQ 10GM BASE/VIAL	A062435 003	Nov 15, 1983

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

CEFADYL

APOTHECON EQ 500MG BASE/VIAL	A062961 001	Sep 20, 1988
EQ 500MG BASE/VIAL	N050446 005	
EQ 1GM BASE/VIAL	A061769 001	
EQ 1GM BASE/VIAL	A062724 001	Dec 23, 1986
EQ 1GM BASE/VIAL	A062961 002	Sep 20, 1988
EQ 1GM BASE/VIAL	N050446 001	
EQ 2GM BASE/VIAL	A061769 002	
EQ 2GM BASE/VIAL	A062724 002	Dec 23, 1986
EQ 2GM BASE/VIAL	A062961 003	Sep 20, 1988
EQ 2GM BASE/VIAL	N050446 002	
EQ 4GM BASE/VIAL	A061769 003	
EQ 4GM BASE/VIAL	A062961 004	Sep 20, 1988
EQ 4GM BASE/VIAL	N050446 003	
EQ 20GM BASE/VIAL	N050446 004	

CEPHAPIRIN SODIUM

ABRAXIS PHARM EQ 500MG BASE/VIAL	A062723 001	Nov 17, 1986
EQ 1GM BASE/VIAL	A062723 002	Nov 17, 1986
EQ 2GM BASE/VIAL	A062723 003	Nov 17, 1986
EQ 4GM BASE/VIAL	A062723 004	Nov 17, 1986
EQ 20GM BASE/VIAL	A062723 005	Nov 17, 1986
HIKMA EQ 500MG BASE/VIAL	A062720 001	Jul 02, 1987
EQ 1GM BASE/VIAL	A062720 002	Jul 02, 1987
EQ 2GM BASE/VIAL	A062720 003	Jul 02, 1987
EQ 20GM BASE/VIAL	A062720 004	Jul 02, 1987

CEPHRADINE

CAPSULE; ORAL

ANSPOR

GLAXOSMITHKLINE 250MG	A061859 001	
500MG	A061859 002	

CEPHRADINE

BARR 250MG	A062850 001	Apr 22, 1988
500MG	A062851 001	Apr 22, 1988
IVAX SUB TEVA PHARMS 250MG	A062762 001	Mar 06, 1987
500MG	A062762 002	Mar 06, 1987
TEVA 250MG	A062683 001	Jan 09, 1987
500MG	A062683 002	Jan 09, 1987
VITARINE 250MG	A062813 001	Feb 25, 1988
500MG	A062813 002	Feb 25, 1988

VELOSEF

APOTHECON 250MG	A061764 001	
500MG	A061764 002	

VELOSEF '250'

ERSANA 250MG	N050548 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHRADINE

CAPSULE; ORAL

VELOSEF '500'

ERSANA

500MG

N050548 002

FOR SUSPENSION; ORAL

ANSPOR

GLAXOSMITHKLINE

125MG/5ML

A061866 001

250MG/5ML

A061866 002

CEPHRADINE

BARR

125MG/5ML

A062858 001 May 19, 1988

250MG/5ML

A062859 001 May 19, 1988

TEVA

125MG/5ML

A062693 001 Jan 09, 1987

250MG/5ML

A062693 002 Jan 09, 1987

VELOSEF '125'

APOTHECON

125MG/5ML

A061763 001

VELOSEF '250'

APOTHECON

250MG/5ML

A061763 002

INJECTABLE; INJECTION

VELOSEF

APOTHECON

250MG/VIAL

A061976 001

500MG/VIAL

A061976 002

1GM/VIAL

A061976 004

2GM/VIAL

A061976 003

4GM/VIAL

A061976 005

TABLET; ORAL

VELOSEF

BRISTOL MYERS SQUIBB 1GM

N050530 001

CERITINIB

CAPSULE; ORAL

ZYKADIA

+

NOVARTIS

150MG

N205755 001 Apr 29, 2014

CERIVASTATIN SODIUM

TABLET; ORAL

BAYCOL

BAYER PHARMS

0.05MG

N020740 001 Jun 26, 1997

0.1MG

N020740 002 Jun 26, 1997

0.2MG

N020740 003 Jun 26, 1997

0.3MG

N020740 004 Jun 26, 1997

0.4MG

N020740 005 May 24, 1999

0.8MG

N020740 006 Jul 24, 2000

CERULETIDE DIETHYLAMINE

INJECTABLE; INJECTION

TYMTRAN

PHARMACIA AND UPJOHN 0.02MG/ML

N018296 001

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

APOTEX

10MG

A207235 001 Aug 12, 2016

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 5MG/5ML

A078617 001 Feb 02, 2010

APOZEAL PHARMS 5MG/5ML

A078870 001 Apr 27, 2009

AUROBINDO PHARMA LTD 5MG/5ML

A090751 001 Dec 16, 2009

LANNETT CO INC 5MG/5ML

A078496 001 Sep 25, 2009

PHARM ASSOC 5MG/5ML

A078412 001 Jun 18, 2008

RANBAXY LABS LTD 5MG/5ML

A077472 001 Jun 18, 2008

WOCKHARDT 5MG/5ML

A078757 001 Aug 28, 2009

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

APOZEAL PHARMS 5MG/5ML

A090474 002 Mar 30, 2009

CHARTWELL RX 5MG/5ML

A090378 002 May 09, 2008

CYPRESS PHARM 5MG/5ML

A090300 001 Oct 10, 2008

PHARM ASSOC 5MG/5ML

A090188 002 Apr 22, 2008

RANBAXY LABS LTD 5MG/5ML

A090183 002 Apr 24, 2008

TRIS PHARMA INC 5MG/5ML

A090572 001 Nov 16, 2012

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

APOZEAL PHARMS 5MG/5ML

A090474 001 Mar 30, 2009

CHARTWELL RX 5MG/5ML

A090378 001 May 09, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CETIRIZINE HYDROCHLORIDE

SYRUP;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

CYPRESS PHARM	5MG/5ML	A090300	002	Oct 10, 2008
PHARM ASSOC	5MG/5ML	A090188	001	Apr 22, 2008
RANBAXY LABS LTD	5MG/5ML	A090183	001	Apr 24, 2008
TRIS PHARMA INC	5MG/5ML	A090572	002	Nov 16, 2012

Zyrtec

J AND J CONSUMER INC	5MG/5ML **	N020346	001	Sep 27, 1996
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TABLET;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

CIPLA LTD	5MG	A077318	001	Jul 25, 2013
	10MG	A077318	002	Jul 25, 2013
HERITAGE PHARMA	5MG	A078615	003	Dec 28, 2007
	10MG	A078615	004	Dec 28, 2007
SUN PHARM INDS INC	5MG	A077499	001	Dec 27, 2007
	10MG	A077499	002	Dec 27, 2007
TORRENT PHARMS LLC	5MG	A079191	001	Apr 15, 2010
	10MG	A079191	004	Apr 15, 2010
UNICHEM	5MG	A078680	003	Jun 26, 2009
	10MG	A078680	004	Jun 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES

SUN PHARM INDS INC	5MG	A077499	003	Dec 27, 2007
	10MG	A077499	004	Dec 27, 2007
UNICHEM	5MG	A078680	001	Jun 26, 2009
	10MG	A078680	002	Jun 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

TORRENT PHARMS LLC	5MG	A079191	003	Apr 15, 2010
	10MG	A079191	002	Apr 15, 2010

Zyrtec Allergy

+ J AND J CONSUMER INC	5MG	N019835	003	Nov 16, 2007
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Zyrtec Hives Relief

+ J AND J CONSUMER INC	5MG	N019835	005	Nov 16, 2007
+	10MG	N019835	006	Nov 16, 2007

TABLET, CHEWABLE;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

SUN PHARM INDS INC	5MG	A077631	004	Jan 11, 2008
	10MG	A077631	003	Jan 11, 2008

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SUN PHARM INDS INC	5MG	A077631	001	Jan 11, 2008
	10MG	A077631	002	Jan 11, 2008

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

SUN PHARM	10MG	A090142	002	Aug 30, 2011
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CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SUN PHARM	10MG	A090142	004	Aug 30, 2011
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CHILDREN'S Zyrtec Allergy

+ J AND J CONSUMER INC	5MG **	N021621	003	Nov 16, 2007
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CHILDREN'S Zyrtec Hives Relief

+ J AND J CONSUMER INC	5MG **	N021621	005	Nov 16, 2007
+	10MG **	N021621	006	Nov 16, 2007

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA LTD	5MG;120MG	A212409	001	Mar 08, 2023
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CETRORELIX ACETATE

POWDER;SUBCUTANEOUS

Cetrotide

+ EMD SERONO INC	EQ 3MG BASE/VIAL	N021197	002	Aug 11, 2000
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CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL

FOR SUSPENSION;INTRATRACHEAL

EXOSURF NEONATAL

GLAXOSMITHKLINE	12MG/VIAL;108MG/VIAL;8MG/VIAL	N020044	001	Aug 02, 1990
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

CEVIMELINE HYDROCHLORIDE

APOTEX INC

30MG

A091260 001 Aug 25, 2011

CHENODIOL

TABLET; ORAL

CHENIX

+ LEADIANT BIOSCI INC

250MG **

N018513 002 Jul 28, 1983

CHLOPHEDIANOL HYDROCHLORIDE

SYRUP; ORAL

ULO

3M

25MG/5ML

N012126 001

CHLORAMPHENICOL

CREAM; TOPICAL

CHLOROMYCETIN

PARKE DAVIS

1%

N050183 001

FOR SOLUTION; OPHTHALMIC

CHLOROMYCETIN

PARKEDALE

25MG/VIAL

N050143 001

INJECTABLE; INJECTION

CHLOROMYCETIN

PARKE DAVIS

250MG/ML

N050153 001

OINTMENT; OPHTHALMIC

CHLORAMPHENICOL

ALTANA

1%

A060133 001

CHLOROFAIR

PHARMAFAIR

1%

A062439 001 Apr 21, 1983

CHLOROMYCETIN

PARKEDALE

1%

N050156 001

CHLOROPTIC S.O.P.

ALLERGAN

1%

A061187 001

ECONOCHLOR

ALCON

1%

A061648 001

SOLUTION/DROPS; OPHTHALMIC

CHLORAMPHENICOL

AKORN

0.5%

A062042 001

ALCON

0.5%

A062628 001 Sep 25, 1985

CHLOROFAIR

PHARMAFAIR

0.5%

A062437 001 Apr 14, 1983

CHLOROPTIC

ALLERGAN

0.5%

N050091 001

ECONOCHLOR

ALCON

0.5%

A061645 001

OPHTHOCHLOR

PARKEDALE

0.5%

A061220 001

OPTOMYCIN

OPTOPICS

0.5%

A062171 001 Mar 31, 1982

SOLUTION/DROPS; OTIC

CHLOROMYCETIN

PARKEDALE

0.5%

N050205 001

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL

ELKINS SINN

EQ 1GM BASE/VIAL

A062406 001 Nov 09, 1982

CHLORAMPHENICOL SODIUM SUCCINATE

GRUPPO LEPETIT

EQ 1GM BASE/VIAL

A062278 001

CHLOROMYCETIN

+ PARKEDALE

EQ 1GM BASE/VIAL

N050155 001

MYCHEL-S

ANGUS

EQ 1GM BASE/VIAL

A060132 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE

FOR SUSPENSION; OPHTHALMIC

CHLOROMYCETIN HYDROCORTISONE

PARKEDALE

12.5MG/VIAL; 25MG/VIAL

N050202 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORAMPHENICOL; HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

OPHTHOCORT

PARKEDALE

10MG/GM;5MG/GM;10,000 UNITS/GM

N050201 002

CHLORAMPHENICOL; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

CHLOROMYXIN

PARKE DAVIS

1%;10,000 UNITS/GM

N050203 002

CHLORAMPHENICOL; PREDNISOLONE

OINTMENT;OPHTHALMIC

CHLOROPTIC-P S.O.P.

ALLERGAN

1%;0.5%

A061188 001

CHLORDIAZEPOXIDE

CAPSULE, EXTENDED RELEASE;ORAL

LIBRELEASE

VALEANT PHARM INTL

30MG

N017813 001 Sep 12, 1983

TABLET;ORAL

LIBRITABS

VALEANT PHARM INTL

5MG

A085482 001

10MG

A085481 001

25MG

A085488 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE;ORAL

A-POXIDE

ABBOTT

5MG

A085447 001

5MG

A085517 001

10MG

A085447 002

10MG

A085518 001

25MG

A085447 003

25MG

A085513 001

CHLORDIAZACHEL

RACHELLE

5MG

A085086 001

10MG

A084639 001

25MG

A085087 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

ASCOT

5MG

A087525 001 Jan 07, 1982

10MG

A087524 001 Jan 07, 1982

25MG

A087512 001 Jan 07, 1982

FERRANTE

5MG

A085118 001

10MG

A085119 001

25MG

A085120 001

HALSEY

5MG

A085340 001

10MG

A085339 001

25MG

A084685 001

IMPAX LABS

5MG

A086213 001

10MG

A085113 001

25MG

A086212 001

IVAX SUB TEVA PHARMS

5MG

A083741 001

10MG

A083742 001

25MG

A083570 001

LEDERLE

5MG

A086892 001

5MG

A087234 001

10MG

A086876 001

10MG

A087037 001

25MG

A086893 001

25MG

A087231 001

MAST MM

10MG

A086217 001

MYLAN

5MG

A084886 001

10MG

A084601 001

25MG

A084887 001

PARKE DAVIS

5MG

A085163 001

10MG

A084598 001

25MG

A085164 001

PIONEER PHARMS

10MG

A089533 001 Jul 15, 1988

25MG

A089558 001 Jul 15, 1988

PUREPAC PHARM

5MG

A085155 001

10MG

A084939 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

	25MG	A085144	001	
ROXANE	5MG	A084706	001	
	10MG	A084700	001	
	25MG	A084705	001	
SUPERPHARM	5MG	A088987	001	Apr 25, 1985
	10MG	A088986	001	Apr 25, 1985
	25MG	A088988	001	Apr 25, 1985
TEVA	5MG	A088705	001	Jan 18, 1985
	10MG	A088706	001	Jan 18, 1985
	25MG	A086494	001	
	25MG	A088707	001	Jan 18, 1985
UPSHER SMITH LABS	5MG	A084919	001	
+	10MG	A084920	001	
	25MG	A084823	001	
USL PHARMA	5MG	A084644	001	
	10MG	A084623	001	
	25MG	A084645	001	
VANGARD	5MG	A088129	001	Mar 28, 1983
	10MG	A088010	001	Mar 28, 1983
	25MG	A088130	001	Mar 28, 1983
WATSON LABS	5MG	A086383	001	
	10MG	A086294	001	
	25MG	A086382	001	
WEST WARD	5MG	A085014	001	
	10MG	A085000	001	
	25MG	A085294	001	
LIBRIUM				
+	VALEANT PHARM INTL	5MG **	N012249	002
+		10MG **	N012249	001
+		25MG **	N012249	003
LYGEN				
ALRA	5MG	A085107	001	
	10MG	A085009	001	
	25MG	A085108	001	

INJECTABLE; INJECTION

LIBRIUM

BAUSCH 100MG/AMP N012301 001

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE

TEVA PHARMS USA	5MG; 2.5MG	A211476	001	Nov 02, 2021
TORRENT	5MG; 2.5MG	A217385	001	Dec 14, 2023

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET; ORAL

MENRIUM 10-4

ROCHE 10MG; 0.4MG N014740 006

MENRIUM 5-2

ROCHE 5MG; 0.2MG N014740 002

MENRIUM 5-4

ROCHE 5MG; 0.4MG N014740 004

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

AKORN	0.12%	A074356	001	May 07, 1996
PAROEX				
SUNSTAR AMERICAS	0.12%	A076434	001	Nov 29, 2005
PERIOGARD				
COLGATE PALMOLIVE CO	0.12%	A073695	001	Jan 14, 1994

SOLUTION; TOPICAL

BRIAN CARE

SOAPCO 4% A071419 001 Dec 17, 1987

EXIDINE

XTTRIUM 2.5% N019421 001 Dec 17, 1985

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORHEXIDINE GLUCONATESOLUTION; TOPICAL
MICRODERM

J AND J 4% A072255 001 Apr 15, 1991

PREVACARE R

J AND J 0.5% A072292 001 Jan 28, 1992

STERI-STAT

MATRIX MEDCL 4% A070104 001 Jul 24, 1986

SPONGE; TOPICAL

CHLORHEXIDINE GLUCONATE

KENDALL IL 4% N019490 001 Mar 27, 1987

E-Z SCRUB

BECTON DICKINSON 4% A073416 001 Mar 14, 2000

HIBICLENS

+ MOLNLYCKE HLTH 4% ** N018423 001

MICRODERM

J AND J 4% A072295 001 Feb 28, 1991

PHARMASEAL SCRUB CARE

CAREFUSION 2200 4% N019793 001 Dec 02, 1988

TABLET; DENTAL

PERIOCHIP

+ DEXCEL PHARMA 2.5MG N020774 001 May 15, 1998

CHLORMERODRIN HG-197

INJECTABLE; INJECTION

CHLORMERODRIN HG 197

BRACCO 0.6-1.4mCi/ML N017269 001

CHLORMEZANONE

TABLET; ORAL

TRANCOPAL

SANOFI AVENTIS US 100MG N011467 003

200MG N011467 005

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

HOSPIRA 2% A087447 001 Apr 16, 1982

3% A087446 001 Apr 16, 1982

NESACAINE-MPF

FRESENIUS KABI USA 2% N009435 003

3% N009435 004

SOLUTION; INTRATHECAL

CLOROTEKAL

+ B BRAUN MEDICAL INC 50MG/5ML (10MG/ML) N208791 001 Sep 26, 2017

CHLOROQUINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARALEN HYDROCHLORIDE

SANOFI AVENTIS US EQ 40MG BASE/ML N006002 002

CHLOROQUINE PHOSPHATE

TABLET; ORAL

ARALEN

+ SANOFI AVENTIS US 500MG ** N006002 001

CHLOROQUINE PHOSPHATE

HIKMA PHARMS 250MG ** A083082 001

500MG ** A083082 002 Sep 17, 1999

MD PHARM 250MG A087228 001

PUREPAC PHARM 250MG A080886 001

TEVA 250MG A087504 001 Jan 13, 1982

WATSON LABS 250MG A087979 001 Dec 21, 1982

500MG A088030 001 Dec 21, 1982

CHLOROQUINE PHOSPHATE; PRIMAQUINE PHOSPHATE

TABLET; ORAL

ARALEN PHOSPHATE W/ PRIMAQUINE PHOSPHATE

SANOFI AVENTIS US EQ 300MG BASE; EQ 45MG BASE N014860 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLOROTHIAZIDE

TABLET; ORAL

CHLOROTHIAZIDE

ABC HOLDING	250MG	A085569	001	
HIKMA INTL PHARMS	250MG	A086028	001	Jul 14, 1982
	500MG	A087736	001	Jul 14, 1982
LEDERLE	250MG	A086940	001	
	500MG	A086938	001	
MYLAN	250MG	A084217	002	
	500MG	A084217	001	
SANDOZ	250MG	A085485	001	
WATSON LABS	250MG	A085165	001	
	250MG	A085173	001	
	250MG	A086795	001	Aug 15, 1983
	500MG	A084026	001	Sep 01, 1982
	500MG	A086796	001	Aug 15, 1983
DIURIL				
+ RISING	250MG **	N011145	004	
+	500MG **	N011145	002	

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

DIURIL

+ RISING	EQ 500MG BASE/VIAL	N011145	005	
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CHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDOCLOR-150

MERCK	150MG;250MG	N016016	001	
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ALDOCLOR-250

MERCK	250MG;250MG	N016016	002	
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METHYLDOPA AND CHLOROTHIAZIDE

PAR PHARM	150MG;250MG	A070783	001	Nov 06, 1987
	250MG;250MG	A070654	001	Nov 06, 1987

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CHLOROTHIAZIDE AND RESERPINE

HIKMA	250MG;0.125MG	A088557	001	Dec 22, 1983
	500MG;0.125MG	A088365	001	Dec 22, 1983

CHLOROTHIAZIDE W/ RESERPINE

WATSON LABS	250MG;0.125MG	A084853	001	
	500MG;0.125MG	A088151	001	Jun 09, 1983

CHLOROTHIAZIDE-RESERPINE

MYLAN	250MG;0.125MG	A087744	001	May 06, 1982
	500MG;0.125MG	A087745	001	May 06, 1982

DIUPRES-250

MERCK	250MG;0.125MG	N011635	003	Aug 26, 1987
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DIUPRES-500

MERCK	500MG;0.125MG	N011635	006	Aug 26, 1987
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CHLOROTRIANISENE

CAPSULE; ORAL

CHLOROTRIANISENE

BANNER PHARMACAPS	12MG	A084652	001	
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TACE

SANOFI AVENTIS US	12MG	N008102	004	
	25MG	N011444	001	
	72MG	N016235	001	

CHLOROXINE

SHAMPOO; TOPICAL

CAPITROL

WESTWOOD SQUIBB	2%	N017594	001	
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CHLORPHENESIN CARBAMATE

TABLET; ORAL

MAOLATE

PAMLAB LLC	400MG	N014217	002	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE;ORAL

CHLORPHENIRAMINE MALEATE

AUROLIFE PHARMA LLC 12MG

A070797 001 Aug 12, 1988

TELDRIN

GLAXOSMITHKLINE 8MG

N017369 001

12MG

N017369 002

INJECTABLE;INJECTION

CHLOR-TRIMETON

SCHERING PLOUGH 10MG/ML

N008826 001

100MG/ML

N008794 001

CHLORPHENIRAMINE MALEATE

BEL MAR 10MG/ML

A080821 001

ELKINS SINN 10MG/ML

A080797 001

WATSON LABS 10MG/ML

A083593 001

10MG/ML

A086096 001

100MG/ML

A086095 001

PYRIDAMAL 100

BEL MAR 100MG/ML

A083733 001

SYRUP;ORAL

CHLOR-TRIMETON

SCHERING 2MG/5ML

N006921 006

CHLORPHENIRAMINE MALEATE

PHARM ASSOC 2MG/5ML

A087520 001 Feb 10, 1982

TABLET;ORAL

ANTAGONATE

BAYER PHARMS 4MG

A083381 001

CHLOR-TRIMETON

SCHERING 4MG

N006921 002

CHLORPHENIRAMINE MALEATE

ANABOLIC 4MG

A083078 001

AUROLIFE PHARMA LLC 4MG

A080961 001

BELL PHARMA 4MG

A083062 001

ELKINS SINN 4MG

A080938 001

IMPAX LABS 4MG

A080809 001

IVAX SUB TEVA PHARMS 4MG

A080779 001

KV PHARM 4MG

A087164 001

LEDERLE 4MG

A086941 001

NEWTRON PHARMS 4MG

A086519 001

PANRAY 4MG

A083243 001

PHARMAVITE 4MG

A085104 001

PHARMERAL 4MG

A083753 001

PIONEER PHARMS 4MG

A088556 001 Jul 13, 1984

PUREPAC PHARM 4MG

A086306 001

PVT FORM 4MG

A080786 001

ROXANE 4MG

A080626 001

SUN PHARM INDUSTRIES 4MG

A080700 001

VITARINE 4MG

A085837 001

WATSON LABS 4MG

A080696 001

4MG

A080791 001

4MG

A085139 001

4MG

A083787 001

Kloromin

HALSEY 4MG

A083629 001

Phenetron

LANNETT 4MG

A080846 001

TABLET, EXTENDED RELEASE;ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC 8MG **

N007638 001

+ 12MG **

N007638 002

EFIDAC 24 CHLORPHENIRAMINE MALEATE

ALZA 16MG

N019746 002 Nov 18, 1994

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

ACELLA 4MG/5ML;5MG/5ML

A206891 001 Jun 09, 2017

TRIS PHARMA INC 4MG/5ML;5MG/5ML

A206438 001 Jan 27, 2015

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION;ORAL

VITUZ

+ PERSION 4MG/5ML;5MG/5ML N204307 001 Feb 20, 2013

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

APOZEAL PHARMS 4MG/5ML;5MG/5ML;60MG/5ML A206660 001 May 15, 2017

MAYNE PHARMA INC 4MG/5ML;5MG/5ML;60MG/5ML A205657 001 Aug 03, 2015

TRIS PHARMA INC 4MG/5ML;5MG/5ML;60MG/5ML A203838 001 Nov 26, 2014

ZUTRIPRO

+ PERSION 4MG/5ML;5MG/5ML;60MG/5ML ** N022439 001 Jun 08, 2011

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

COLD CAPSULE IV

GRAHAM DM 12MG;75MG N018793 001 Apr 25, 1985

COLD CAPSULE V

GRAHAM DM 8MG;75MG N018794 001 Apr 23, 1985

TABLET, EXTENDED RELEASE;ORAL

TRIAMINIC-12

NOVARTIS 12MG;75MG N018115 001

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

CODIMAL-L.A. 12

SCHWARZ PHARMA 12MG;120MG N018935 001 Apr 15, 1985

ISOCOLOR

FISONS 8MG;120MG N018747 001 Mar 06, 1986

PSEUDOEPHEDRINE HYDROCHLORIDE AND CHLORPHENIRAMINE MALEATE

CENT PHARMS 8MG;120MG N019428 001 Aug 02, 1988

GRAHAM DM 8MG;120MG N018844 001 Mar 20, 1985

12MG;120MG N018843 001 Mar 18, 1985

KV PHARM 12MG;120MG A071455 001 Mar 01, 1989

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC 8MG;120MG N018397 001

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

CODEPREX

LANNETT CO INC EQ 4MG MALEATE/5ML;EQ 20MG BASE/5ML N021369 001 Jun 21, 2004

PENNTUSS

FISONS EQ 4MG MALEATE/5ML;EQ 10MG BASE/5ML N018928 001 Aug 14, 1985

TUZISTRA XR

+ TRIS PHARMA INC EQ 2.8MG BASE/5ML;EQ 14.7MG BASE/5ML N207768 001 Apr 30, 2015

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

CAPSULE, EXTENDED RELEASE;ORAL

TUSSICAPS

ECR PHARMA EQ 4MG MALEATE;EQ 5MG BITARTRATE A077273 002 Sep 24, 2007

EQ 8MG MALEATE;EQ 10MG BITARTRATE A077273 001 Sep 24, 2007

SUSPENSION, EXTENDED RELEASE;ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

NEOS THERAPS INC EQ 8MG MALEATE/5ML;EQ 10MG A091671 001 Jun 29, 2012

BITARTRATE/5ML

TUSSIONEX PENNKINETIC

+ UCB INC EQ 8MG MALEATE/5ML;EQ 10MG N019111 001 Dec 31, 1987

BITARTRATE/5ML **

CHLORPHENTERMINE HYDROCHLORIDE

TABLET;ORAL

PRE-SATE

PARKE DAVIS EQ 65MG BASE N014696 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROMAZINE

SUPPOSITORY;RECTAL

THORAZINE

+	GLAXOSMITHKLINE	25MG **	N009149	024
+		100MG **	N009149	033

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

THORAZINE

	GLAXOSMITHKLINE	30MG	N011120	016
		75MG	N011120	017
		150MG	N011120	018
		200MG	N011120	019
		300MG	N011120	020

CONCENTRATE;ORAL

CHLORPROMAZINE HYDROCHLORIDE

	ACTAVIS MID ATLANTIC	100MG/ML	A086863	001
	PHARM ASSOC	30MG/ML	A040231	001 Dec 30, 1999
		100MG/ML	A040224	001 Jan 26, 1999
	WOCKHARDT	30MG/ML	A087032	001 Jul 08, 1982
		100MG/ML	A087053	001

CHLORPROMAZINE HYDROCHLORIDE INTENSOL

	HIKMA	30MG/ML	A088157	001 Apr 27, 1983
		100MG/ML	A088158	001 Apr 27, 1983

SONAZINE

	FOSUN PHARMA	30MG/ML	A080983	004
		100MG/ML	A080983	005

THORAZINE

+	GLAXOSMITHKLINE	30MG/ML **	N009149	032
+		100MG/ML **	N009149	043

INJECTABLE;INJECTION

CHLORPROMAZINE HYDROCHLORIDE

	ABRAXIS PHARM	25MG/ML	A084911	001
	DR REDDYS	25MG/ML	A080365	001
	MARSAM PHARMS LLC	25MG/ML	A089563	001 Apr 15, 1988
	WATSON LABS	25MG/ML	A085591	001
	WYETH AYERST	25MG/ML	A080370	001

THORAZINE

+	GLAXOSMITHKLINE	25MG/ML **	N009149	011
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SYRUP;ORAL

CHLORPROMAZINE HYDROCHLORIDE

	ALPHARMA US PHARMS	10MG/5ML	A086712	001
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SONAZINE

	FOSUN PHARMA	10MG/5ML	A083040	001
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THORAZINE

+	GLAXOSMITHKLINE	10MG/5ML **	N009149	022
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TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

	ABBOTT	10MG	A084414	001
		25MG	A084415	001
		50MG	A084411	001
		100MG	A084412	001
		200MG	A084413	001
	CYCLE	10MG	A085331	001
		25MG	A085331	002
		50MG	A085331	003
		100MG	A085331	004
		200MG	A085331	005
	IVAX SUB TEVA PHARMS	10MG	A083549	001
		25MG	A083549	002
		50MG	A083549	003
		100MG	A083574	001
		200MG	A083575	001
	KV PHARM	10MG	A085750	002 Jan 04, 1982
		25MG	A085751	001
		50MG	A085484	001
		100MG	A085752	001
		200MG	A085748	002 Jan 04, 1982
	LEDERLE	10MG	A084803	001
		25MG	A084801	001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

	50MG	A084800	001	
	100MG	A084789	001	
	200MG	A084802	001	
PUREPAC PHARM	10MG	A080403	004	
	25MG	A080403	001	
	50MG	A080403	002	
	100MG	A080403	003	
	200MG	A080403	005	
PVT FORM	25MG	A080340	001	
	50MG	A080340	002	
	200MG	A080340	003	
SANDOZ	10MG **	A080439	001	
	25MG **	A080439	002	
	50MG **	A080439	003	
	100MG **	A080439	004	
	200MG **	A080439	005	
VANGARD	10MG	A088038	001	Aug 16, 1982
	25MG	A087645	001	
	50MG	A087646	001	
WATSON LABS	10MG	A085959	001	
	25MG	A085956	001	
	50MG	A085960	001	
	100MG	A085957	001	
	200MG	A085958	001	
WEST WARD	10MG	A087783	001	Sep 16, 1982
	25MG	A087865	001	Sep 16, 1982
	50MG	A087878	001	Sep 15, 1982
	100MG	A087884	001	Sep 15, 1982
	200MG	A087880	001	Sep 16, 1982
PROMAPAR				
PARKE DAVIS	10MG	A086886	001	
	25MG	A084423	001	
	50MG	A086887	001	
	100MG	A086888	001	
	200MG	A086885	001	
THORAZINE				
GLAXOSMITHKLINE	10MG **	N009149	002	
	25MG **	N009149	007	
	50MG **	N009149	013	
	100MG **	N009149	018	
	200MG **	N009149	020	

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

ANI PHARMS	100MG	A088768	001	Oct 11, 1984
	100MG	A088812	001	Oct 19, 1984
	100MG	A088840	001	Oct 25, 1984
	100MG	A088918	001	Oct 16, 1984
	100MG	A088921	001	Apr 12, 1985
	100MG	A089446	001	Nov 17, 1986
	250MG	A087353	001	
	250MG	A088813	001	Oct 19, 1984
	250MG	A088826	001	Sep 26, 1984
	250MG	A088919	001	Oct 16, 1984
	250MG	A088922	001	Apr 12, 1985
	250MG	A089447	001	Nov 17, 1986
DAVA PHARMS INC	100MG	A089561	001	Sep 04, 1987
	250MG	A089562	001	Sep 04, 1987
HALSEY	100MG	A089321	001	Jan 16, 1986
	250MG	A088662	001	Jan 09, 1986
MYLAN	100MG	A088549	002	Jun 01, 1984
	250MG	A088549	001	Jun 01, 1984
PAR PHARM	100MG	A088175	001	Feb 27, 1984
	250MG	A088176	001	Feb 27, 1984
RISING	100MG	A088725	001	Aug 31, 1984
	250MG	A088726	001	Aug 31, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

SANDOZ	250MG	A084669	001	
SUPERPHARM	100MG	A088694	001	Sep 17, 1984
	250MG	A088695	001	Sep 17, 1984
USL PHARMA	100MG	A088708	001	Aug 30, 1984
	250MG	A088709	001	Aug 30, 1984
WATSON LABS	100MG	A086865	001	Sep 24, 1984
	100MG	A088608	001	Apr 12, 1984
	250MG	A086866	001	
	250MG	A088568	001	Apr 12, 1984
WATSON LABS TEVA	100MG	A088852	001	Sep 26, 1984
DIABINESE				
+ PFIZER	100MG	N011641	003	
+	250MG	N011641	006	
GLUCAMIDE				
ANI PHARMS	250MG	A088641	001	Oct 11, 1984

CHLORPROTHIXENE

CONCENTRATE; ORAL

TARACTAN

ROCHE	100MG/5ML	N016149	002	
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INJECTABLE; INJECTION

TARACTAN

ROCHE	12.5MG/ML	N012487	001	
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TABLET; ORAL

TARACTAN

ROCHE	10MG	N012486	005	
	25MG	N012486	004	
	50MG	N012486	003	
	100MG	N012486	001	

CHLORTETRACYCLINE HYDROCHLORIDE

OINTMENT; OPHTHALMIC

AUREOMYCIN

LEDERLE	1%	N050404	001	
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CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

ABBOTT	25MG	A087364	001	
	50MG	A087384	001	
ANI PHARMS	25MG	A087296	001	
	25MG	A087706	001	
	25MG	A088164	001	Jan 09, 1984
	50MG	A087689	001	
ASCOT	25MG	A087698	001	Oct 20, 1982
	50MG	A087699	001	Oct 20, 1982
BARR LABS INC	25MG	A088902	001	Sep 19, 1985
	50MG	A088903	001	Sep 19, 1985
COSETTE	50MG	A088651	001	May 30, 1985
DAVA PHARMS INC	25MG	A087451	001	
	50MG	A087450	001	
IVAX PHARMS	25MG	A087555	001	
	50MG	A087176	001	
	50MG	A087947	001	Feb 27, 1984
KV PHARM	25MG	A087311	001	
	50MG	A087312	001	
MUTUAL PHARM	25MG	A087292	001	
	25MG	A089738	001	Sep 19, 1988
	50MG	A087293	001	
	50MG	A089739	001	Sep 19, 1988
PIONEER PHARMS	50MG	A089591	001	Jul 21, 1988
PUREPAC PHARM	25MG	A088139	001	Jul 16, 1986
	50MG	A088140	001	Aug 11, 1983
SANDOZ	25MG	A087380	001	
	50MG	A087118	001	
	50MG	A087381	001	
SUNNY	25MG	A209068	001	Jan 25, 2022
	50MG	A209068	002	Jan 25, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

SUPERPHARM	25MG	A087473 001	Feb 09, 1983
	50MG	A087247 001	Feb 09, 1983
USL PHARMA	25MG	A089051 001	Jun 01, 1987
	50MG	A089052 001	Jun 01, 1987
VANGARD	25MG	A088012 001	Jul 14, 1982
	50MG	A088073 001	Mar 25, 1983
WARNER CHILCOTT	25MG	A087515 001	Jan 24, 1983
	50MG	A087516 001	Feb 09, 1983
WATSON LABS	25MG	A087050 001	
	25MG	A087100 001	
	50MG	A087029 001	
	50MG	A087082 001	
	50MG	A087521 001	
HYGROTON			
+ SANOFI AVENTIS US	25MG **	N012283 004	
+	50MG **	N012283 003	
THALITONE			
MONARCH PHARMS	25MG	A088051 001	Nov 12, 1982

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE AND CHLORTHALIDONE

PAR PHARM	15MG; 0.1MG	A071179 001	Dec 16, 1987
	15MG; 0.2MG	A071178 001	Dec 16, 1987
	15MG; 0.3MG	A071142 001	Dec 16, 1987
CLORPRES			
MYLAN	15MG; 0.1MG	A071325 003	Feb 09, 1987
	15MG; 0.2MG	A071325 002	Feb 09, 1987
	15MG; 0.3MG	A071325 001	Feb 09, 1987
COMBIPRES			
+ BOEHRINGER INGELHEIM	15MG; 0.1MG **	N017503 001	
+	15MG; 0.2MG **	N017503 002	
+	15MG; 0.3MG **	N017503 003	Apr 10, 1984

CHLORTHALIDONE; METOPROLOL TARTRATE

CAPSULE; ORAL

LOPRESSIDONE

NOVARTIS	25MG; 100MG	N019451 001	Dec 31, 1987
	25MG; 200MG	N019451 002	Dec 31, 1987

CHLORTHALIDONE; RESERPINE

TABLET; ORAL

DEMI-REGROTON

SANOFI AVENTIS US	25MG; 0.125MG	N015103 002	
REGROTON			
SANOFI AVENTIS US	50MG; 0.25MG	N015103 001	

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

ACME LABS	500MG	A214365 001	Apr 27, 2023
ACTAVIS ELIZABETH	250MG	A088928 001	May 08, 1987
	500MG	A040113 001	Sep 29, 1995
BARR	500MG	A089895 001	May 04, 1988
GLENMARK PHARMS LTD	375MG	A212185 001	May 26, 2020
	750MG	A212185 002	May 26, 2020
OHM LABS	250MG	A081298 001	Dec 29, 1993
	500MG	A081299 001	Dec 29, 1993
PIONEER PHARMS	250MG	A089592 001	Jan 06, 1989
	500MG	A089948 001	Jan 06, 1989
RISING	250MG	A089852 001	May 04, 1988
STRIDES PHARMA	250MG	A087981 001	Sep 20, 1983
SUN PHARM INDUSTRIES	500MG	A089970 001	Sep 27, 1990
WATSON LABS	250MG	A086901 001	
	250MG	A086948 001	Aug 09, 1982
	500MG	A040137 001	Aug 09, 1996
	500MG	A081019 001	Jul 29, 1991
	500MG	A081040 001	Aug 22, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORZOXAZONE

TABLET; ORAL

PARAFLEX

+ ORTHO MCNEIL PHARM 250MG **

N011300 003

PARAFON FORTE DSC

+ JANSSEN R AND D 500MG **

N011529 002 Jun 15, 1987

STRIFON FORTE DSC

FERNDAL LABS 500MG

A081008 001 Dec 23, 1988

CHOLESTYRAMINE

BAR, CHEWABLE; ORAL

CHOLYBAR

PARKE DAVIS EQ 4GM RESIN/BAR

A071621 001 May 26, 1988

EQ 4GM RESIN/BAR

A071739 001 May 26, 1988

POWDER; ORAL

CHOLESTYRAMINE

ANI PHARMS EQ 4GM RESIN/PACKET

A074554 001 Oct 02, 1996

EQ 4GM RESIN/SCOOPFUL

A074554 002 Oct 02, 1996

IVAX SUB TEVA PHARMS EQ 4GM RESIN/PACKET

A074771 001 Jul 09, 1997

EQ 4GM RESIN/SCOOPFUL

A074771 002 Jul 09, 1997

TEVA EQ 4GM RESIN/PACKET

A074347 001 May 28, 1998

EQ 4GM RESIN/SCOOPFUL

A074347 002 May 28, 1998

UPSHER SMITH LABS EQ 4GM RESIN/PACKET

A214877 001 Jan 21, 2022

EQ 4GM RESIN/SCOOPFUL

A214877 002 Jan 21, 2022

CHOLESTYRAMINE LIGHT

TEVA EQ 4GM RESIN/PACKET

A074348 001 May 28, 1998

EQ 4GM RESIN/SCOOPFUL

A074348 002 May 28, 1998

TEVA PHARMS EQ 4GM RESIN/PACKET

A074555 001 Sep 30, 1998

EQ 4GM RESIN/SCOOPFUL

A074555 002 Sep 30, 1998

QUESTRAN

+ BRISTOL MYERS EQ 4GM RESIN/PACKET **

N016640 001

+ EQ 4GM RESIN/SCOOPFUL **

N016640 003

QUESTRAN LIGHT

+ BRISTOL MYERS EQ 4GM RESIN/PACKET **

N019669 001 Dec 05, 1988

+ EQ 4GM RESIN/SCOOPFUL **

N019669 003 Dec 05, 1988

TABLET; ORAL

QUESTRAN

APOTHECON EQ 800MG RESIN

A073403 002 Dec 27, 1999

EQ 1GM RESIN

A073403 001 Apr 28, 1994

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11

UCSF RODIOPHARM 4-33.1mCi/ML

A208444 001 Nov 20, 2017

UNIV TX MD ANDERSON 4-100mCi/ML

A205690 001 Oct 29, 2015

WA UNIV SCH MED 4-33.1mCi/ML

A208413 001 Jan 10, 2017

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID

MYLAN PHARMS INC EQ 45MG FENOFIBRIC ACID

A200913 001 Mar 25, 2013

EQ 135MG FENOFIBRIC ACID

A200913 002 Mar 25, 2013

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE

ABRAXIS PHARM EQ 0.004MG CHROMIUM/ML

N019271 001 May 05, 1987

CHROMIC PHOSPHATE P-32

INJECTABLE; INJECTION

PHOSPHOCOL P32

CURIUM 5mCi/ML

N017084 001

CICLOPIROX

GEL; TOPICAL

CICLOPIROX

+ ALVOGEN 0.77% **

N020519 001 Jul 21, 1997

SHAMPOO; TOPICAL

CICLOPIROX

VERTICE 1%

A209975 001 Apr 05, 2018

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CICLOPIROX

SOLUTION; TOPICAL

CICLOPIROX

EPIC PHARMA LLC	8%	A078975	001	Feb 17, 2010
MYLAN PHARMS INC	8%	A078567	001	Sep 18, 2007
TEVA PHARMS	8%	A078079	001	Sep 18, 2007

PENLAC

+ VALEANT BERMUDA	8% **	N021022	001	Dec 17, 1999
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CIDOFOVIR

INJECTABLE; INJECTION

VISTIDE

+ GILEAD SCIENCES INC	EQ 75MG BASE/ML **	N020638	001	Jun 26, 1996
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CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION

PRIMAXIN

MERCK	EQ 250MG BASE/VIAL; 250MG/VIAL	A062756	001	Jan 08, 1987
	EQ 500MG BASE/VIAL; 500MG/VIAL	A062756	002	Jan 08, 1987

POWDER; INTRAMUSCULAR

PRIMAXIN

MERCK	EQ 500MG BASE/VIAL; 500MG/VIAL	N050630	001	Dec 14, 1990
	EQ 750MG BASE/VIAL; 750MG/VIAL	N050630	002	Dec 14, 1990

POWDER; INTRAVENOUS

IMIPENEM AND CILASTATIN

HOSPIRA INC	EQ 250MG BASE/VIAL; 250MG/VIAL	A090825	001	Nov 16, 2011
	EQ 500MG BASE/VIAL; 500MG/VIAL	A090825	002	Nov 16, 2011
	EQ 500MG BASE/VIAL; 500MG/VIAL	A091007	001	Nov 16, 2011

PRIMAXIN

+ MERCK	EQ 250MG BASE/VIAL; 250MG/VIAL **	N050587	001	Nov 26, 1985
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CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

ACTAVIS ELIZABETH	100MG	A077028	002	Nov 26, 2004
AUROBINDO PHARMA USA	50MG	A077019	001	Nov 23, 2004
	100MG	A077019	002	Nov 23, 2004
CASI PHARMS INC	50MG	A077310	001	Nov 08, 2005
	100MG	A077021	001	Nov 23, 2004
HIKMA	50MG	A077024	001	May 17, 2005
	100MG	A077024	002	May 17, 2005
IVAX SUB TEVA PHARMS	100MG	A077020	002	Mar 01, 2005
MYLAN	50MG	A077323	002	Apr 20, 2006
	100MG	A077323	001	Apr 20, 2006
NOSTRUM LABS INC	50MG	A077708	001	Sep 28, 2009
	100MG	A077708	002	Sep 28, 2009
PLIVA HRVATSKA DOO	50MG	A077898	001	Oct 29, 2007
	100MG	A077898	002	Oct 29, 2007

PLETAL

+ OTSUKA	50MG **	N020863	001	Jan 15, 1999
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+	100MG **	N020863	002	Jan 15, 1999
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CIMETIDINE

SUSPENSION; ORAL

TAGAMET HB 200

GLAXOSMITHKLINE	200MG/20ML	N020951	001	Jul 09, 1999
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TABLET; ORAL

CIMETIDINE

CHARTWELL MOLECULES	200MG	A074329	002	May 17, 1994
	300MG	A074329	003	May 17, 1994
	400MG	A074329	004	May 17, 1994
	800MG	A074329	001	May 17, 1994
CHARTWELL RX	200MG	A074100	001	Jan 31, 1995
	300MG	A074100	002	Jan 31, 1995
	400MG	A074100	003	Jan 31, 1995
	800MG	A074100	004	Jan 31, 1995
CONTRACT PHARMACAL	200MG	A074961	001	Jun 19, 1998
	200MG	A074963	001	Jun 19, 1998
CYCLE	300MG	A074361	001	Dec 23, 1994
	400MG	A074361	002	Dec 23, 1994
	800MG	A074371	001	Dec 23, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIMETIDINETABLET; ORAL
CIMETIDINE

HIKMA	200MG	A074890 001	Dec 18, 1998
	300MG	A074890 002	Dec 18, 1998
	400MG	A074890 003	Dec 18, 1998
	800MG	A074890 004	Dec 18, 1998
IVAX SUB TEVA PHARMS	200MG	A074401 001	May 30, 1995
	200MG	A074424 001	Jul 28, 1995
	200MG	A075345 001	Jun 16, 1999
	300MG	A074401 002	May 30, 1995
	300MG	A074424 002	Jul 28, 1995
	400MG	A074401 003	May 30, 1995
	400MG	A074424 003	Jul 28, 1995
	800MG	A074402 001	May 30, 1995
	800MG	A074424 004	Jul 28, 1995
NOVITIUM PHARMA	300MG	A074340 001	Jun 23, 1995
	400MG	A074340 002	Jun 23, 1995
	800MG	A074339 001	Jun 23, 1995
PERRIGO	100MG	A074972 001	Jun 19, 1998
PLIVA	200MG	A074568 001	Feb 27, 1997
	300MG	A074568 002	Feb 27, 1997
	400MG	A074568 003	Feb 27, 1997
	800MG	A074566 001	Feb 27, 1997
SANDOZ	100MG	A075122 001	Jun 19, 1998
	200MG	A074250 001	Jun 29, 1995
	200MG	A075122 002	Jun 19, 1998
	300MG	A074250 002	Jun 29, 1995
	400MG	A074250 003	Jun 29, 1995
	800MG	A074250 004	Jun 29, 1995
TEVA	200MG	A074365 001	Feb 28, 1995
	300MG	A074365 002	Feb 28, 1995
	400MG	A074365 003	Feb 28, 1995
	800MG	A074365 004	Feb 28, 1995
UPSHER SMITH LABS	200MG	A074506 001	Jan 24, 1996
	300MG	A074506 002	Jan 24, 1996
	400MG	A074506 003	Jan 24, 1996
	800MG	A074506 004	Jan 24, 1996
WATSON LABS INC	200MG	A074349 001	Aug 30, 1996
	300MG	A074349 002	Aug 30, 1996
	400MG	A074349 003	Aug 30, 1996
	800MG	A074316 001	Feb 28, 1996
WATSON LABS TEVA	200MG	A075425 001	Jul 29, 1999
TAGAMET			
+ GLAXOSMITHKLINE	200MG **	N017920 002	
+	300MG **	N017920 003	
+	400MG **	N017920 004	Dec 14, 1983
+	800MG **	N017920 005	Apr 30, 1986
TAGAMET HB			
+ MEDTECH PRODUCTS	100MG **	N020238 001	Jun 19, 1995

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

DAVA PHARMS INC	EQ 300MG BASE/2ML	A074428 001	Apr 25, 1996
HOSPIRA	EQ 300MG BASE/2ML	A074296 001	Mar 28, 1997
	EQ 300MG BASE/2ML	A074344 001	Jan 31, 1995
	EQ 300MG BASE/2ML	A074345 001	Jan 31, 1995
	EQ 300MG BASE/2ML	A074412 001	Mar 28, 1997
	EQ 300MG BASE/2ML	A074422 001	Jan 31, 1995
LUITPOLD	EQ 300MG BASE/2ML	A074353 001	Dec 20, 1994
TEVA PARENTERAL	EQ 300MG BASE/2ML	A074252 001	Nov 26, 1997
VINTAGE PHARMS LLC	EQ 300MG BASE/2ML	A074005 001	Aug 31, 1994
CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
HOSPIRA	EQ 6MG BASE/ML	A074269 001	Dec 27, 1994
	EQ 90MG BASE/100ML	A074468 005	Dec 29, 1994
	EQ 120MG BASE/100ML	A074468 006	Dec 29, 1994
	EQ 180MG BASE/100ML	A074468 003	Dec 29, 1994
	EQ 240MG BASE/100ML	A074468 004	Dec 29, 1994
	EQ 360MG BASE/100ML	A074468 001	Dec 29, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
EQ 480MG BASE/100ML

A074468 002 Dec 29, 1994

TAGAMET

GLAXOSMITHKLINE EQ 300MG BASE/2ML **

N017939 002

TAGAMET HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+ GLAXOSMITHKLINE EQ 6MG BASE/ML **

N019434 001 Oct 31, 1985

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

ANI PHARMS

EQ 300MG BASE/5ML

A074610 001 Sep 26, 1996

EQ 300MG BASE/5ML

A074859 001 Jul 09, 1998

EQ 300MG BASE/5ML

A075110 001 Jun 18, 1998

CHARTWELL MOLECULAR

EQ 300MG BASE/5ML

A074251 001 Dec 22, 1994

CYCLE

EQ 300MG BASE/5ML

A074541 001 Aug 05, 1997

G AND W LABS INC

EQ 300MG BASE/5ML

A074176 001 Jun 01, 1994

PHARM ASSOC

EQ 300MG BASE/5ML

A075560 001 Mar 15, 2000

PHARMOBEDIANT CNSLTG

EQ 300MG BASE/5ML

A074757 001 Oct 17, 1997

TAGAMET

GLAXOSMITHKLINE EQ 300MG BASE/5ML **

N017924 001

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

LUPIN LTD

EQ 30MG BASE

A210548 001 Jun 28, 2019

EQ 60MG BASE

A210548 002 Jun 28, 2019

EQ 90MG BASE

A210548 003 Jun 28, 2019

MYLAN

EQ 30MG BASE

A203422 001 Oct 16, 2018

EQ 60MG BASE

A203422 002 Oct 16, 2018

EQ 90MG BASE

A203422 003 Oct 16, 2018

STEVENS J

30MG

A204364 001 Dec 02, 2021

60MG

A204364 002 Dec 02, 2021

90MG

A204364 003 Dec 02, 2021

CINOXACIN

CAPSULE; ORAL

CINOXACIN

LILLY

250MG

N018067 001

500MG

N018067 002

CINOXACIN

TEVA

250MG

A073005 001 Feb 28, 1992

500MG

A073006 001 Feb 28, 1992

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO

+ BAYER HLTHCARE

400MG/40ML (10MG/ML) **

N019847 001 Dec 26, 1990

+ BAYER HLTHCARE

200MG/20ML (10MG/ML) **

N019847 002 Dec 26, 1990

1200MG/120ML (10MG/ML) **

N019847 003 Dec 26, 1990

CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER

+ BAYER HLTHCARE

200MG/100ML **

N019857 001 Dec 26, 1990

+ BAYER HLTHCARE

400MG/200ML **

N019857 002 Dec 26, 1990

CIPRO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

BAYER PHARMS

200MG/100ML

N019858 001 Dec 26, 1990

CIPROFLOXACIN

BEDFORD LABS

200MG/20ML (10MG/ML)

A076992 001 Aug 28, 2006

400MG/40ML (10MG/ML)

A076992 002 Aug 28, 2006

1200MG/120ML (10MG/ML)

A076993 001 Aug 28, 2006

DR REDDYS

200MG/20ML (10MG/ML)

A077782 001 Aug 28, 2006

400MG/40ML (10MG/ML)

A077782 002 Aug 28, 2006

FRESENIUS KABI USA

200MG/20ML (10MG/ML)

A076484 001 Aug 28, 2006

400MG/40ML (10MG/ML)

A076484 002 Aug 28, 2006

HOSPIRA

200MG/20ML (10MG/ML)

A077245 001 Aug 28, 2006

400MG/40ML (10MG/ML)

A077245 002 Aug 28, 2006

CIPROFLOXACIN IN DEXTROSE 5%

HIKMA FARMACEUTICA

200MG/100ML

A076757 001 Apr 21, 2008

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE

200MG/100ML

A077888 001 Mar 18, 2008

400MG/200ML

A077888 002 Mar 18, 2008

BEDFORD

200MG/100ML

A078114 001 Mar 18, 2008

400MG/200ML

A078114 002 Mar 18, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

TEVA PHARMS

200MG/100ML

A077138 001 Mar 18, 2008

400MG/200ML

A077138 002 Mar 18, 2008

INJECTABLE, SUSPENSION; OTIC

OTIPRIO

+

ALK ABELLO

6% (60MG/ML)

N207986 001 Dec 10, 2015

CIPROFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CIPROFLOXACIN HYDROCHLORIDE

AKORN

EQ 0.3% BASE

A076555 001 Dec 11, 2008

AMRING PHARMS

EQ 0.3% BASE

A078598 001 Jan 16, 2008

PAI HOLDINGS PHARM

EQ 0.3% BASE

A076754 001 Jun 09, 2004

TABLET; ORAL

CIPRO

+

BAYER HLTHCARE

EQ 100MG BASE **

N019537 001 Apr 08, 1996

+

EQ 750MG BASE **

N019537 004 Oct 22, 1987

CIPROFLOXACIN HYDROCHLORIDE

AIPING PHARM INC

EQ 250MG BASE

A076593 002 Jun 09, 2004

EQ 500MG BASE

A076593 003 Jun 09, 2004

EQ 750MG BASE

A076593 004 Jun 09, 2004

AMNEAL

EQ 100MG BASE

A075939 001 Mar 03, 2005

BARR

EQ 250MG BASE

A074124 001 Jun 09, 2004

EQ 500MG BASE

A074124 002 Jun 09, 2004

EQ 750MG BASE

A074124 003 Jun 09, 2004

DR REDDYS LABS LTD

EQ 100MG BASE

A075593 002 Jun 09, 2004

MYLAN

EQ 250MG BASE

A075685 002 Jun 09, 2004

EQ 500MG BASE

A075685 003 Jun 09, 2004

EQ 750MG BASE

A075685 001 Jun 09, 2004

NOSTRUM LABS

EQ 250MG BASE

A076138 001 Jun 09, 2004

EQ 500MG BASE

A076138 002 Jun 09, 2004

EQ 750MG BASE

A076138 003 Jun 09, 2004

PLIVA

EQ 100MG BASE

A076426 001 Jun 15, 2005

EQ 250MG BASE

A076426 002 Jun 15, 2005

EQ 500MG BASE

A076426 003 Jun 15, 2005

EQ 750MG BASE

A076426 004 Jun 15, 2005

RISING

EQ 100MG BASE

A075817 001 Jun 25, 2007

EQ 250MG BASE

A075817 002 Jun 09, 2004

EQ 750MG BASE

A075817 004 Jun 09, 2004

SUN PHARM INDS LTD

EQ 250MG BASE

A075747 001 Jun 09, 2004

EQ 500MG BASE

A075747 002 Jun 09, 2004

EQ 750MG BASE

A075747 003 Jun 09, 2004

TARO

EQ 100MG BASE

A076912 001 Feb 18, 2005

EQ 250MG BASE

A076912 002 Oct 06, 2004

EQ 500MG BASE

A076912 003 Oct 06, 2004

EQ 750MG BASE

A076912 004 Oct 06, 2004

TEVA

EQ 250MG BASE

A076136 001 Jun 09, 2004

EQ 500MG BASE

A076136 002 Jun 09, 2004

EQ 750MG BASE

A076136 003 Jun 09, 2004

WATSON LABS

EQ 100MG BASE

A076794 001 Feb 10, 2005

TABLET, EXTENDED RELEASE; ORAL

PROQUIN XR

DEPOMED INC

EQ 500MG BASE

N021744 001 May 19, 2005

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPRO XR

BAYER HLTHCARE

212.6MG;EQ 287.5MG BASE **

N021473 001 Dec 13, 2002

425.2MG;EQ 574.9MG BASE **

N021473 002 Aug 28, 2003

CIPROFLOXACIN EXTENDED RELEASE

ANI PHARMS

212.6MG;EQ 287.5MG BASE

A077809 002 Nov 30, 2010

425.2MG;EQ 574.9MG BASE

A077809 001 Nov 30, 2010

DR REDDYS LABS LTD

212.6MG;EQ 287.5MG BASE

A077701 002 Oct 31, 2007

FOSUN PHARMA

212.6MG;EQ 287.5MG BASE

A078712 001 Dec 11, 2007

RISING

212.6MG;EQ 287.5MG BASE

A078183 001 Mar 22, 2007

425.2MG;EQ 574.9MG BASE

A078183 002 Mar 22, 2007

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CISAPRIDE MONOHYDRATE

SUSPENSION;ORAL

PROPULSID

JANSSEN PHARMS

EQ 1MG BASE/ML

N020398 001 Sep 15, 1995

TABLET;ORAL

PROPULSID

JANSSEN PHARMS

EQ 10MG BASE

N020210 001 Jul 29, 1993

EQ 20MG BASE

N020210 002 Dec 23, 1993

TABLET, ORALLY DISINTEGRATING;ORAL

PROPULSID QUICKSOLV

JANSSEN PHARMA

EQ 20MG BASE

N020767 001 Nov 07, 1997

CISATRACURIUM BESYLATE

INJECTABLE;INJECTION

CISATRACURIUM BESYLATE

ACCORD HLTHCARE

EQ 2MG BASE/ML

A205873 001 Jun 16, 2017

PIRAMAL HLTHCARE UK

EQ 2MG BASE/ML

A212190 001 May 04, 2020

EQ 2MG BASE/ML

A212432 001 Mar 11, 2022

EQ 10MG BASE/ML

A212432 002 Mar 11, 2022

ZYDUS PHARMS

EQ 2MG BASE/ML

A213527 001 Aug 31, 2020

CISATRACURIUM BESYLATE PRESERVATIVE FREE

ACCORD HLTHCARE

EQ 2MG BASE/ML

A205872 001 Jun 16, 2017

EQ 10MG BASE/ML

A205872 002 Jun 16, 2017

CISPLATIN

INJECTABLE;INJECTION

CISPLATIN

BEDFORD

10MG/VIAL

A074713 001 Nov 14, 2000

50MG/VIAL

A074713 002 Nov 14, 2000

MYLAN LABS LTD

1MG/ML

A091062 001 Apr 18, 2012

TEVA PHARMS USA

1MG/ML

A074814 001 May 16, 2000

PLATINOL

+ HQ SPCLT PHARMA

10MG/VIAL

N018057 001

PLATINOL-AQ

+ HQ SPCLT PHARMA

0.5MG/ML

N018057 003 Jul 18, 1984

CITALOPRAM HYDROBROMIDE

CAPSULE;ORAL

CITALOPRAM HYDROBROMIDE

MYLAN PHARMS INC

EQ 10MG BASE

A077668 001 Feb 28, 2007

EQ 20MG BASE

A077668 002 Feb 28, 2007

EQ 40MG BASE

A077668 003 Feb 28, 2007

SOLUTION;ORAL

CELEXA

+ FOREST LABS

EQ 10MG BASE/5ML **

N021046 001 Dec 22, 1999

CITALOPRAM HYDROBROMIDE

PHARM ASSOC

EQ 10MG BASE/5ML

A077601 001 Nov 15, 2005

TABLET;ORAL

CELEXA

ABBVIE

EQ 60MG BASE

N020822 004 Jul 17, 1998

CITALOPRAM HYDROBROMIDE

EPIC PHARMA LLC

EQ 10MG BASE

A077036 001 Oct 28, 2004

EQ 20MG BASE

A077036 002 Oct 28, 2004

EQ 40MG BASE

A077036 003 Oct 28, 2004

FOSUN PHARMA

EQ 10MG BASE

A077035 001 Oct 28, 2004

EQ 10MG BASE

A077040 001 Aug 17, 2005

EQ 20MG BASE

A077035 002 Oct 28, 2004

EQ 20MG BASE

A077040 002 Aug 17, 2005

EQ 40MG BASE

A077035 003 Oct 28, 2004

EQ 40MG BASE

A077040 003 Aug 17, 2005

HERITAGE PHARMA

EQ 10MG BASE

A077033 001 Oct 28, 2004

EQ 10MG BASE

A077034 001 Jun 30, 2005

EQ 10MG BASE

A077213 001 Mar 31, 2006

EQ 10MG BASE

A077232 001 Oct 31, 2005

EQ 20MG BASE

A077033 002 Oct 28, 2004

EQ 20MG BASE

A077034 002 Jun 30, 2005

EQ 20MG BASE

A077213 002 Mar 31, 2006

EQ 20MG BASE

A077232 002 Oct 31, 2005

EQ 40MG BASE

A077033 003 Oct 28, 2004

EQ 40MG BASE

A077034 003 Jun 30, 2005

EQ 40MG BASE

A077213 003 Mar 31, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CITALOPRAM HYDROBROMIDE

TABLET;ORAL

CITALOPRAM HYDROBROMIDE

	EQ 40MG BASE	A077232 003	Oct 31, 2005
JUBILANT GENERICS	EQ 10MG BASE	A205407 001	Dec 23, 2015
	EQ 20MG BASE	A205407 002	Dec 23, 2015
	EQ 40MG BASE	A205407 003	Dec 23, 2015
MYLAN	EQ 10MG BASE	A077039 001	Feb 03, 2005
	EQ 20MG BASE	A077039 002	Feb 03, 2005
	EQ 40MG BASE	A077039 003	Feb 03, 2005
MYLAN PHARMS INC	EQ 10MG BASE	A077037 001	Nov 05, 2004
	EQ 20MG BASE	A077037 002	Nov 05, 2004
	EQ 40MG BASE	A077037 003	Nov 05, 2004
NATCO PHARMA LTD	EQ 20MG BASE	A077141 002	Apr 10, 2008
	EQ 40MG BASE	A077141 001	Apr 10, 2008
ROXANE	EQ 10MG BASE	A077041 001	Nov 23, 2004
	EQ 20MG BASE	A077041 002	Nov 23, 2004
	EQ 40MG BASE	A077041 003	Nov 23, 2004
SUN PHARM INDS INC	EQ 10MG BASE	A077032 001	Nov 12, 2004
	EQ 20MG BASE	A077032 002	Nov 12, 2004
	EQ 40MG BASE	A077032 003	Nov 12, 2004
SUN PHARM INDUSTRIES	EQ 10MG BASE	A077052 001	Jul 03, 2006
	EQ 20MG BASE	A077052 002	Jul 03, 2006
	EQ 40MG BASE	A077052 003	Jul 03, 2006
TARO	EQ 10MG BASE	A077278 001	Mar 22, 2006
	EQ 20MG BASE	A077278 002	Mar 22, 2006
	EQ 40MG BASE	A077278 003	Mar 22, 2006

TABLET, ORALLY DISINTEGRATING;ORAL

CITALOPRAM HYDROBROMIDE

+	BIOVAIL LABS INTL	EQ 10MG BASE	N021763 001	Dec 20, 2005
+		EQ 20MG BASE	N021763 002	Dec 20, 2005
+		EQ 40MG BASE	N021763 003	Dec 20, 2005

CITRIC ACID; MAGNESIUM OXIDE; SODIUM CARBONATE

SOLUTION;IRRIGATION

IRRIGATING SOLUTION G IN PLASTIC CONTAINER

BAXTER HLTHCARE	3.24GM/100ML;380MG/100ML;430MG/100ML	N018519 001	Jun 22, 1982
UROLOGIC G IN PLASTIC CONTAINER			
HOSPIRA	3.24GM/100ML;380MG/100ML;430MG/100ML	N018904 001	May 27, 1983

CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

FOR SOLUTION;ORAL

PREPOPIK

+	FERRING PHARMS INC	12GM/PACKET;3.5GM/PACKET;10MG/PACKET **	N202535 001	Jul 16, 2012
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CLADRIBINE

INJECTABLE;INJECTION

CLADRIBINE

MYLAN LABS LTD	1MG/ML	A200510 001	Oct 06, 2011	
LEUSTATIN				
+	JANSSEN PHARMS	1MG/ML **	N020229 001	Feb 26, 1993

CLARITHROMYCIN

FOR SUSPENSION;ORAL

BIAXIN

+	ABBVIE	125MG/5ML	N050698 001	Dec 23, 1993
		187MG/5ML	N050698 003	Sep 30, 1998
+		250MG/5ML	N050698 002	Dec 23, 1993

CLARITHROMYCIN

SUN PHARM INDS LTD	125MG/5ML	A065382 001	Aug 30, 2007
	250MG/5ML	A065382 002	Aug 30, 2007

TABLET;ORAL

BIAXIN

+	ABBVIE	250MG **	N050662 001	Oct 31, 1991
+		500MG **	N050662 002	Oct 31, 1991

CLARITHROMYCIN

AJANTA PHARMA LTD	250MG	A206714 001	Apr 25, 2019
	500MG	A206714 002	Apr 25, 2019
HIKMA	250MG	A065178 002	May 25, 2004
	500MG	A065178 001	May 25, 2004
IVAX SUB TEVA PHARMS	250MG	A065137 001	May 31, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLARITHROMYCIN

TABLET; ORAL

CLARITHROMYCIN

	500MG	A065137 002	May 31, 2005
MYLAN	250MG	A065195 001	Mar 11, 2005
	500MG	A065195 002	Mar 11, 2005
SUN PHARM INDS LTD	250MG	A065174 001	Sep 24, 2004
	500MG	A065174 002	Sep 24, 2004
TEVA	250MG	A065155 001	May 31, 2005
	500MG	A065155 002	May 31, 2005
WOCKHARDT	250MG	A065266 001	May 31, 2006
	500MG	A065266 002	May 31, 2006

TABLET, EXTENDED RELEASE; ORAL

BIAXIN XL

+ ABBVIE	500MG **	N050775 001	Mar 03, 2000
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CLARITHROMYCIN

ANI PHARMS	500MG	A065250 001	Aug 25, 2005
LUPIN LTD	500MG	A202532 001	Sep 15, 2015
RANBAXY	1GM	A065210 001	Jan 26, 2005

CLAVULANATE POTASSIUM; TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TIMENTIN

GLAXOSMITHKLINE	EQ 100MG BASE/VIAL; EQ 3GM BASE/VIAL	A062691 001	Dec 19, 1986
	EQ 100MG BASE/VIAL; EQ 3GM BASE/VIAL	N050590 001	Apr 01, 1985
	EQ 200MG BASE/VIAL; EQ 3GM BASE/VIAL	N050590 002	Apr 01, 1985
	EQ 1GM BASE/VIAL; EQ 30GM BASE/VIAL	N050590 003	Aug 18, 1987

TIMENTIN IN PLASTIC CONTAINER

GLAXOSMITHKLINE	EQ 100MG BASE/100ML; EQ 3GM BASE/100ML	N050658 001	Dec 15, 1989
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CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

ACTAVIS MID ATLANTIC	EQ 0.5MG BASE/5ML	A074075 001	Oct 31, 1993
APOTEX INC	EQ 0.5MG BASE/5ML	A075703 001	Nov 27, 2000
CHARTWELL MOLECULAR	EQ 0.5MG BASE/5ML	A074884 001	Dec 17, 1997
TEVA PHARMS	EQ 0.5MG BASE/5ML	A073095 001	Apr 21, 1992
WOCKHARDT BIO AG	EQ 0.5MG BASE/5ML	A074863 001	Mar 13, 1998

TAVIST

+ NOVARTIS	EQ 0.5MG BASE/5ML **	N018675 001	Jun 28, 1985
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TABLET; ORAL

CLEMASTINE FUMARATE

ANI PHARMS	1.34MG	A073282 001	Jan 31, 1992
	1.34MG	A073282 002	Dec 03, 1992
L PERRIGO CO	1.34MG	A074512 001	Nov 22, 1995
PLD ACQUISITIONS LLC	1.34MG	A073458 001	Oct 31, 1993
SANDOZ	2.68MG	A073459 001	Oct 31, 1993

TAVIST

+ NOVARTIS	2.68MG	N017661 001	
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TAVIST-1

+ GLAXOSMITHKLINE CONS	1.34MG **	N020925 001	Aug 21, 1992
NOVARTIS	1.34MG	N017661 002	
	1.34MG	N017661 003	Aug 21, 1992

CLEVIDIPINE

EMULSION; INTRAVENOUS

CLEVIPREX

+ CHIESI	125MG/250ML (0.5MG/ML)	N022156 003	Nov 08, 2013
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CLIDINIUM BROMIDE

CAPSULE; ORAL

QUARZAN

ROCHE	2.5MG	N010355 001	
	5MG	N010355 002	

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN

PHARMACIA AND UPJOHN	EQ 75MG BASE	A061809 001	
	EQ 150MG BASE	A061809 002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLINDAMYCIN HYDROCHLORIDE

CAPSULE;ORAL

CLINDAMYCIN HYDROCHLORIDE

CHARTWELL MOLECULAR	EQ 150MG BASE	A065243 003	Aug 12, 2005
	EQ 300MG BASE	A065243 001	Aug 12, 2005
MYLAN PHARMS INC	EQ 75MG BASE	A091225 001	May 31, 2011
	EQ 150MG BASE	A091225 002	May 31, 2011
	EQ 300MG BASE	A091225 003	May 31, 2011
TEVA	EQ 75MG BASE	A063027 001	Sep 20, 1989
WATSON LABS	EQ 75MG BASE	A063082 001	Jul 31, 1991
	EQ 150MG BASE	A063083 001	Jul 31, 1991
	EQ 300MG BASE	A063083 002	Mar 18, 2003

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION;ORAL

CLEOCIN

PHARMACIA AND UPJOHN	EQ 75MG BASE/5ML **	A061827 001	
CLINDAMYCIN PALMITATE HYDROCHLORIDE			
MYLAN	EQ 75MG BASE/5ML	A203063 001	May 25, 2016

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM;TOPICAL

EVOCLIN

+ MYLAN	1%	N050801 001	Oct 22, 2004
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CREAM;VAGINAL

CLEOCIN

PFIZER	EQ 2% BASE	N050680 001	Aug 11, 1992
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INJECTABLE;INJECTION

CLEOCIN PHOSPHATE

PHARMACIA AND UPJOHN	EQ 150MG BASE/ML	A061839 001	
CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER			
+ PFIZER	EQ 6MG BASE/ML **	N050639 001	Aug 30, 1989
+	EQ 12MG BASE/ML **	N050639 002	Aug 30, 1989
+	EQ 18MG BASE/ML **	N050639 003	Apr 10, 1991

CLINDAMYCIN PHOSPHATE

ABRAXIS PHARM	EQ 150MG BASE/ML	A062747 001	Jun 03, 1988
ALMAJECT	EQ 150MG BASE/ML	A062801 001	Jul 24, 1987
BEDFORD	EQ 150MG BASE/ML	A063163 001	Jun 30, 1994
BRISTOL MYERS SQUIBB	EQ 150MG BASE/ML	A062908 001	Feb 01, 1989
HIKMA	EQ 150MG BASE/ML	A062806 001	Oct 15, 1987
	EQ 150MG BASE/ML	A062953 001	Apr 21, 1988
	EQ 150MG BASE/ML	A063068 001	Aug 28, 1989
IGI LABS INC	EQ 150MG BASE/ML	A062928 001	Feb 13, 1989
LOCH	EQ 150MG BASE/ML	A062905 001	May 09, 1988
MARSAM PHARMS LLC	EQ 150MG BASE/ML	A062913 001	Oct 20, 1988
RISING	EQ 150MG BASE/ML	A204748 001	Oct 10, 2017
	EQ 150MG BASE/ML	A204749 001	Oct 10, 2017
SOLOPAK	EQ 150MG BASE/ML	A062819 001	Mar 15, 1988
	EQ 150MG BASE/ML	A062852 001	Mar 17, 1988
TEVA PARENTERAL	EQ 150MG BASE/ML	A063041 001	Dec 29, 1989
	EQ 150MG BASE/ML	A063282 001	May 29, 1992
WATSON LABS	EQ 150MG BASE/ML	A062900 001	Jun 08, 1988
	EQ 150MG BASE/ML	A063079 001	Mar 05, 1990

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

ABRAXIS PHARM	EQ 12MG BASE/ML	N050636 001	Dec 22, 1989
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CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT LABS	EQ 6MG BASE/ML	A065027 001	Jun 29, 2001
	EQ 12MG BASE/ML	A065027 002	Jun 29, 2001
	EQ 18MG BASE/ML	A065027 003	Jun 29, 2001
BAXTER HLTHCARE	EQ 6MG BASE/ML	N050648 001	Dec 29, 1989
	EQ 12MG BASE/ML	N050648 002	Dec 29, 1989
	EQ 900MG BASE/100ML	N050648 003	Dec 29, 1989

SOLUTION;TOPICAL

CLEOCIN T

+ PFIZER	EQ 1% BASE **	N050537 001	
PHARMACIA AND UPJOHN	EQ 1% BASE	A062363 001	Feb 08, 1982

CLINDAMYCIN PHOSPHATE

BOCA PHARMA LLC	EQ 1% BASE	A062944 001	Jan 11, 1989
FOUGERA PHARMS	EQ 1% BASE	A065254 001	Feb 14, 2006
G AND W LABS INC	EQ 1% BASE	A062811 001	Sep 01, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLINDAMYCIN PHOSPHATE

SOLUTION; TOPICAL

CLINDAMYCIN PHOSPHATE

NOVAST LABS	EQ 1% BASE	A064108	001	Sep 27, 1996
PAI HOLDINGS PHARM	EQ 1% BASE	A206945	001	Dec 30, 2016
VINTAGE PHARMS	EQ 1% BASE	A062930	001	Jun 28, 1989
	EQ 1% BASE	A203343	001	May 29, 2015
XTRTRIUM LABS INC	EQ 1% BASE	A063304	001	Jul 15, 1997

SWAB; TOPICAL

CLEOCIN

+ PFIZER	EQ 1% BASE	N050537	002	Feb 22, 1994
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CLIOQUINOL; NYSTATIN

OINTMENT; TOPICAL

NYSTAFORM

BAYER PHARMS	10MG/GM; 100,000 UNITS/GM	N050235	001	
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CLOBAZAM

SUSPENSION; ORAL

CLOBAZAM

ACCORD HLTHCARE	2.5MG/ML	A216008	001	Sep 06, 2022
HIKMA	2.5MG/ML	A209715	001	Oct 22, 2018
MYLAN	2.5MG/ML	A211259	001	Oct 22, 2018
TEVA PHARMS USA	2.5MG/ML	A211032	001	Jan 31, 2020
UPSHER SMITH LABS	2.5MG/ML	A210569	001	Oct 22, 2018
VISTAPHARM	2.5MG/ML	A210746	001	Jul 10, 2019

TABLET; ORAL

CLOBAZAM

ACCORD HLTHCARE	10MG	A212398	001	May 23, 2019
	20MG	A212398	002	May 23, 2019
APOTEX	10MG	A209853	001	Jun 09, 2020
	20MG	A209853	002	Jun 09, 2020
BRECKENRIDGE	5MG	A209308	003	Oct 19, 2021
CELLTRION	10MG	A211959	001	Dec 09, 2020
	20MG	A211959	002	Dec 09, 2020
CHARTWELL MOLECULAR	10MG	A212092	001	Oct 30, 2019
	20MG	A212092	002	Oct 30, 2019
HIKMA	10MG	A208785	001	Oct 22, 2018
	20MG	A208785	002	Oct 22, 2018
TARO	10MG	A209440	001	Oct 22, 2018
	20MG	A209440	002	Oct 22, 2018

ONFI

+ LUNDBECK PHARMS LLC	5MG **	N202067	001	Oct 21, 2011
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CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

NOVAST LABS	0.05%	A206805	001	Jul 31, 2017
OLUX				
+ MYLAN	0.05%	N021142	001	May 26, 2000
OLUX E				
+ MYLAN	0.05%	N022013	001	Jan 12, 2007

CREAM; TOPICAL

CLOBETASOL PROPIONATE

CHARTWELL RX	0.05%	A211207	001	Mar 26, 2021
PAI HOLDINGS PHARM	0.05%	A209974	001	Apr 17, 2018
TEVA PHARMS USA	0.05%	A074087	001	Feb 16, 1994
TORRENT	0.05%	A211836	001	Dec 30, 2019

CLOBETASOL PROPIONATE (EMOLLIENT)

ANI PHARMS	0.05%	A075733	001	Aug 22, 2001
PAI HOLDINGS PHARM	0.05%	A209411	001	Aug 21, 2017

EMBELINE E

HIKMA	0.05%	A075325	001	Dec 24, 1998
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TEMOVATE

+ FOUGERA PHARMS	0.05% **	N019322	001	Dec 27, 1985
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TEMOVATE E

+ FOUGERA PHARMS	0.05% **	N020340	001	Jun 17, 1994
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GEL; TOPICAL

CLOBETASOL PROPIONATE

PAI HOLDINGS PHARM	0.05%	A208881	001	Mar 06, 2017
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOBETASOL PROPIONATE

GEL;TOPICAL

TEMOVATE

+ FOUGERA PHARMS 0.05% ** N020337 001 Apr 29, 1994

LOTION;TOPICAL

CLOBETASOL PROPIONATE

EPIC PHARMA LLC 0.05% A211348 001 Oct 26, 2018

PAI HOLDINGS PHARM 0.05% A208667 001 Nov 29, 2016

IMPEKLO

+ MYLAN 0.05% N213691 001 May 19, 2020

OINTMENT;TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC 0.05% A074128 001 Aug 03, 1994

AMNEAL 0.05% A210551 001 Aug 21, 2018

PAI HOLDINGS PHARM 0.05% A208589 001 Jan 23, 2019

TORRENT 0.05% A212926 001 Oct 25, 2019

TEMOVATE

+ FOUGERA PHARMS 0.05% ** N019323 001 Dec 27, 1985

SHAMPOO;TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC 0.05% A078854 001 Jun 07, 2011

SOLUTION;TOPICAL

CLOBETASOL PROPIONATE

ENCUBE 0.05% A076977 001 Aug 05, 2005

PRINSTON INC 0.05% A213139 001 Feb 08, 2021

WOCKHARDT BIO AG 0.05% A075205 001 Nov 13, 1998

TEMOVATE

+ FOUGERA PHARMS 0.05% ** N019966 001 Feb 22, 1990

SPRAY;TOPICAL

CLOBETASOL PROPIONATE

AKORN 0.05% A207218 001 Apr 28, 2017

APOTEX 0.05% A210446 001 Apr 17, 2018

CLOFARABINE

SOLUTION;INTRAVENOUS

CLOFARABINE

EUGIA PHARMA 20MG/20ML (1MG/ML) A212457 001 Oct 03, 2022

HOSPIRA 20MG/20ML (1MG/ML) A210283 001 Dec 27, 2018

NOVAST LABS 20MG/20ML (1MG/ML) A210270 001 Sep 14, 2018

CLOFAZIMINE

CAPSULE;ORAL

LAMPRENE

+ NOVARTIS 50MG N019500 002 Dec 15, 1986

100MG N019500 001 Dec 15, 1986

CLOFIBRATE

CAPSULE;ORAL

ATROMID-S

WYETH AYERST 500MG N016099 002

CLOFIBRATE

BANNER PHARMACAPS 500MG A073396 001 Mar 20, 1992

SANDOZ 500MG A072191 001 May 02, 1988

TEVA 500MG A072600 001 Jul 25, 1991

USL PHARMA 500MG A070531 001 Jun 16, 1986

WATSON LABS 500MG A071603 001 Sep 18, 1987

CLOMIPHENE CITRATE

TABLET;ORAL

CLOMID

+ SANOFI AVENTIS US 50MG ** N016131 002

MILOPHENE

GRANATA BIO 50MG A072196 001 Dec 20, 1988

SEROPHENE

EMD SERONO 50MG N018361 001 Mar 22, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

TEVA	25MG	A074849 001	Apr 04, 1997
	25MG	A074958 001	Aug 26, 1997
	50MG	A074849 002	Apr 04, 1997
	50MG	A074958 002	Aug 26, 1997
	75MG	A074849 003	Apr 04, 1997
	75MG	A074958 003	Aug 26, 1997
WATSON LABS	25MG	A074600 001	Nov 27, 1996
	25MG	A074751 001	Sep 30, 1998
	50MG	A074600 002	Nov 27, 1996
	50MG	A074751 002	Sep 30, 1998
	75MG	A074600 003	Nov 27, 1996
	75MG	A074751 003	Sep 30, 1998

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

CHARTWELL RX	0.5MG	A074925 001	Sep 30, 1997
	1MG	A074925 002	Sep 30, 1997
	2MG	A074925 003	Sep 30, 1997
MYLAN PHARMS INC	0.5MG	A074940 001	Oct 30, 1997
	1MG	A074940 002	Oct 30, 1997
	2MG	A074940 003	Oct 30, 1997
SUN PHARM INDS INC	0.5MG	A075423 001	Apr 27, 2001
	1MG	A075423 002	Apr 27, 2001
	2MG	A075423 003	Apr 27, 2001
TEVA	0.5MG	A074920 001	Aug 04, 1998
	1MG	A074920 002	Aug 04, 1998
	2MG	A074920 003	Aug 04, 1998
WATSON LABS	0.5MG	A074964 001	Dec 30, 1997
	1MG	A074964 002	Dec 30, 1997
	2MG	A074964 003	Dec 30, 1997
KLONOPIN			
CHEPLAPHARM	0.125MG	N017533 005	Apr 09, 1997
	0.25MG	N017533 006	Apr 09, 1997
TABLET, ORALLY DISINTEGRATING; ORAL			
KLONOPIN RAPIDLY DISINTEGRATING			
+ ROCHE	0.125MG **	N020813 001	Dec 23, 1997
+	0.25MG **	N020813 002	Dec 23, 1997
+	0.5MG **	N020813 003	Dec 23, 1997
+	1MG **	N020813 004	Dec 23, 1997
+	2MG **	N020813 005	Dec 23, 1997

CLONIDINE

SUSPENSION, EXTENDED RELEASE; ORAL

CLONIDINE

TRIS PHARMA INC	EQ 0.09MG BASE/ML	N022499 001	Dec 03, 2009
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TABLET, EXTENDED RELEASE; ORAL

NEXICLON XR

ATHENA	EQ 0.26MG BASE	N022500 002	Dec 03, 2009
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CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

AM REGENT	1MG/10ML (0.1MG/ML)	A091104 001	Oct 08, 2009
	5MG/10ML (0.5MG/ML)	A091104 002	Oct 08, 2009

DURACLON

+ MYLAN INSTITUTIONAL	5MG/10ML (0.5MG/ML) **	N020615 002	Apr 27, 1999
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TABLET; ORAL

CATAPRES

+ BOEHRINGER INGELHEIM	0.1MG **	N017407 001	
+	0.2MG **	N017407 002	
+	0.3MG **	N017407 003	

CLONIDINE HYDROCHLORIDE

AM THERAP	0.1MG	A070881 001	Jul 08, 1986
	0.2MG	A070882 001	Jul 08, 1986
	0.3MG	A070883 001	Jul 08, 1986
DURAMED PHARMS BARR	0.1MG	A071103 001	Aug 14, 1986
	0.2MG	A071102 001	Aug 14, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

	0.3MG	A071101	001	Aug 14, 1986
INTERPHARM	0.1MG	A071252	001	Oct 01, 1986
	0.2MG	A071253	001	Oct 01, 1986
	0.3MG	A071254	001	Oct 01, 1986
PAR PHARM	0.1MG	A070461	001	Jul 08, 1986
	0.2MG	A070460	001	Jul 08, 1986
	0.3MG	A070459	001	Jul 08, 1986
RISING	0.1MG	A070317	002	Jul 09, 1987
	0.2MG	A070317	003	Jun 09, 1987
	0.3MG	A070317	001	Jun 09, 1987
SUN PHARM INDS INC	0.1MG	A090329	001	Jul 03, 2014
	0.2MG	A090329	002	Jul 03, 2014
	0.3MG	A090329	003	Jul 03, 2014
TEVA	0.1MG	A070747	001	Jul 08, 1986
	0.2MG	A070702	001	Jul 08, 1986
	0.3MG	A070659	001	Jul 08, 1986
WARNER CHILCOTT	0.1MG	A072138	001	Jun 13, 1988
	0.2MG	A072139	001	Jun 13, 1988
	0.3MG	A072140	001	Jun 13, 1988
WATSON LABS	0.1MG	A070395	001	Mar 23, 1987
	0.1MG	A070965	001	Jul 08, 1986
	0.2MG	A070396	001	Mar 23, 1987
	0.2MG	A070964	001	Jul 08, 1986
	0.3MG	A070397	001	Mar 23, 1987
	0.3MG	A070963	001	Jul 08, 1986

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

ACTAVIS ELIZABETH	0.1MG	A202792	001	May 15, 2015
	0.2MG	A202792	002	May 15, 2015
	0.2MG	A203320	002	May 15, 2015
AMNEAL PHARMS NY	0.1MG	A210052	001	Nov 20, 2017
ANCHEN PHARMS	0.1MG	A202983	001	Apr 02, 2014
	0.2MG	A202983	002	Apr 02, 2014
	0.2MG	A202984	002	Sep 30, 2013
DR REDDYS LABS SA	0.1MG	A210680	001	Apr 30, 2018
UPSHER SMITH LABS	0.1MG	A211433	001	Oct 12, 2018
JENLOGA				
+ CONCORDIA PHARMS INC	0.1MG **	N022331	001	Sep 30, 2009
+	0.2MG **	N022331	002	May 25, 2010
KAPVAY				
+ CONCORDIA PHARMS INC	0.1MG	N022331	003	Sep 28, 2010
+	0.2MG **	N022331	004	Sep 28, 2010

CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

ACTAVIS TOTOWA	EQ 75MG BASE	A090307	001	May 28, 2013
ANI PHARMS	EQ 300MG BASE	A090625	001	May 17, 2012
	EQ 300MG BASE	A091216	001	May 17, 2012
CADILA	EQ 75MG BASE	A201686	001	Oct 10, 2012
CHARTWELL RX	EQ 75MG BASE	A202266	001	Aug 14, 2012
	EQ 300MG BASE	A202266	002	Nov 20, 2012
RISING	EQ 75MG BASE	A077665	001	May 17, 2012
	EQ 300MG BASE	A077665	002	May 17, 2012
SUN PHARM	EQ 75MG BASE	A090494	001	May 17, 2012
SUN PHARM INDUSTRIES	EQ 75MG BASE	A078133	001	Jun 10, 2013

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

ABLE	3.75MG	A071777	001	Jul 14, 1987
	7.5MG	A071778	001	Jul 14, 1987
	15MG	A071779	001	Jul 14, 1987
AM THERAP	3.75MG	A071429	001	Jun 23, 1987
	7.5MG	A071430	001	Jun 23, 1987
	15MG	A071431	001	Jun 23, 1987
DASH PHARMS	3.75MG	A071509	001	Oct 19, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

	7.5MG		A071510	001	Oct 19, 1987
	15MG		A071511	001	Oct 19, 1987
DAVA PHARMS INC	3.75MG		A071742	001	Dec 14, 1987
	7.5MG		A071743	001	Dec 14, 1987
	15MG		A071744	001	Dec 14, 1987
GD SEARLE LLC	3.75MG		A071727	001	Dec 18, 1987
	7.5MG		A071728	001	Dec 18, 1987
	15MG		A071729	001	Dec 18, 1987
PUREPAC PHARM	3.75MG		A071924	001	Apr 25, 1988
	7.5MG		A071925	001	Apr 25, 1988
	15MG		A071926	001	Apr 25, 1988
QUANTUM PHARMICS	3.75MG		A071549	001	Sep 12, 1988
	7.5MG		A071550	001	Sep 12, 1988
	15MG		A071522	001	Sep 12, 1988
RISING	3.75MG		A072112	002	Aug 26, 1988
	7.5MG		A072112	003	Aug 26, 1988
	15MG		A072112	001	Aug 26, 1988
USL PHARMA	3.75MG		A071242	001	Jun 23, 1987
	7.5MG		A071243	001	Jun 23, 1987
	15MG		A071244	001	Jun 23, 1987
WARNER CHILCOTT	3.75MG		A071774	001	Mar 01, 1988
	7.5MG		A071775	001	Mar 01, 1988
	15MG		A071776	001	Mar 01, 1988
WATSON LABS	3.75MG		A071878	001	Mar 15, 1988
	7.5MG		A071879	001	Mar 15, 1988
	15MG		A071860	001	Mar 15, 1988
TRANXENE					
+ AJENAT PHARMS	3.75MG **		N017105	001	
+	7.5MG **		N017105	002	
+	15MG **		N017105	003	
TABLET;ORAL					
CLORAZEPATE DIPOTASSIUM					
ABLE	3.75MG		A071780	001	Jun 26, 1987
	7.5MG		A071781	001	Jun 26, 1987
	15MG		A071782	001	Jun 26, 1987
AM THERAP	3.75MG		A071747	001	Jun 23, 1987
	7.5MG		A071748	001	Jun 23, 1987
	15MG		A071749	001	Jun 23, 1987
AUROLIFE PHARMA LLC	3.75MG		A072514	002	May 11, 1990
	7.5MG		A072514	003	May 11, 1990
	15MG		A072514	001	May 11, 1990
LEDERLE	3.75MG		A072013	001	Dec 15, 1987
	7.5MG		A072014	001	Dec 15, 1987
	15MG		A072015	001	Dec 15, 1987
PUREPAC PHARM	3.75MG		A072330	001	Aug 08, 1988
	7.5MG		A072331	001	Aug 08, 1988
	15MG		A072332	001	Aug 08, 1988
QUANTUM PHARMICS	3.75MG		A071730	001	Oct 26, 1987
	7.5MG		A071731	001	Oct 26, 1987
	15MG		A071702	001	Oct 26, 1987
SUN PHARM INDS LTD	3.75MG		A076911	001	Sep 29, 2004
	7.5MG		A076911	002	Sep 29, 2004
	15MG		A076911	003	Sep 29, 2004
WARNER CHILCOTT	3.75MG		A071828	001	Mar 03, 1988
	7.5MG		A071829	001	Mar 03, 1988
	15MG		A071830	001	Mar 03, 1988
WATSON LABS	3.75MG		A071852	001	Feb 09, 1988
	7.5MG		A071853	001	Feb 09, 1988
	15MG		A071854	001	Feb 09, 1988
GEN-XENE					
ALRA	3.75MG		A071787	001	Apr 26, 1988
	7.5MG		A071788	001	Apr 26, 1988
	15MG		A071789	001	Apr 26, 1988
TRANXENE					
+ AJENAT PHARMS	3.75MG **		N017105	006	
+	15MG **		N017105	008	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLORAZEPATE DIPOTASSIUM

TABLET;ORAL

TRANXENE SD

+	AJENAT PHARMS	11.25MG **	N017105 005
+		22.5MG **	N017105 004

CLOTRIMAZOLE

CREAM;TOPICAL

LOTRIMIN

+	SCHERING PLOUGH	1% **	N017619 001
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MYCELEX

	BAYER HEALTHCARE LLC	1%	N018183 001
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CREAM;VAGINAL

GYNE-LOTRIMIN

+	BAYER HEALTHCARE LLC	1% **	N018052 002	Nov 30, 1990
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GYNE-LOTRIMIN 3

+	BAYER HEALTHCARE LLC	2%	N020574 001	Nov 24, 1998
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MYCELEX-7

	BAYER HEALTHCARE LLC	1%	N018230 002	Dec 26, 1991
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CREAM, TABLET;TOPICAL, VAGINAL

GYNE-LOTRIMIN 3 COMBINATION PACK

+	BAYER HEALTHCARE LLC	1%,200MG	N020526 002	Jul 29, 1996
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GYNE-LOTRIMIN COMBINATION PACK

+	BAYER HEALTHCARE LLC	1%,100MG	N020289 002	Apr 26, 1993
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MYCELEX-7 COMBINATION PACK

	BAYER HEALTHCARE LLC	1%,100MG	N020389 002	Jun 23, 1994
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LOTION;TOPICAL

LOTRIMIN

	SCHERING	1%	N018813 001	Feb 17, 1984
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SOLUTION;TOPICAL

LOTRIMIN

+	SCHERING PLOUGH	1% **	N017613 001
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MYCELEX

+	BAYER HLTHCARE	1% **	N018181 001
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TABLET;VAGINAL

GYNE-LOTRIMIN

+	BAYER HEALTHCARE LLC	100MG	N017717 002	Nov 30, 1990
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GYNE-LOTRIMIN 3

+	BAYER HEALTHCARE LLC	200MG	N020525 001	Jul 29, 1996
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GYNIX

	TEVA PHARMS	100MG	A073249 001	Feb 13, 1998
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MYCELEX-7

	BAYER HEALTHCARE LLC	100MG	N018182 002	Dec 26, 1991
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MYCELEX-G

	BAYER PHARMS	500MG	N019069 001	Apr 19, 1985
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TROCHE/LOZENGE;ORAL

MYCELEX

+	BAYER HLTHCARE	10MG **	N018713 001	Jun 17, 1983
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CLOXACILLIN SODIUM

CAPSULE;ORAL

CLOXACILLIN SODIUM

	APOTHECON	EQ 250MG BASE	A061452 001
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		EQ 500MG BASE	A061452 002
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	TEVA	EQ 250MG BASE	A062240 001
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		EQ 500MG BASE	A062240 002
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CLOXAPEN

	GLAXOSMITHKLINE	EQ 250MG BASE	A061806 001
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		EQ 250MG BASE	A062233 001
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		EQ 500MG BASE	A061806 002
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		EQ 500MG BASE	A062233 002
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FOR SOLUTION;ORAL

CLOXACILLIN SODIUM

	TEVA	EQ 125MG BASE/5ML	A062268 001
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		EQ 125MG BASE/5ML	A062978 001	Apr 06, 1989
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TEGOPEN

	APOTHECON	EQ 125MG BASE/5ML	A061453 001
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		EQ 125MG BASE/5ML	N050192 001
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOZAPINE

TABLET; ORAL

CLOZAPINE

DR REDDYS LABS SA	25MG	A203807 001	Sep 17, 2015
	50MG	A203807 003	Aug 22, 2017
	100MG	A203807 002	Sep 17, 2015
	200MG	A203807 004	Aug 22, 2017
MYLAN	12.5MG	A075417 003	Apr 15, 2010
PAR PHARM	25MG	A075162 001	Apr 26, 2005
	100MG	A075162 002	Apr 26, 2005
SANDOZ	25MG	A074546 001	Aug 30, 1996
	100MG	A074546 002	Aug 30, 1996
ZYDUS PHARMS	25MG	A209480 001	Dec 06, 2017
	50MG	A209480 002	Dec 06, 2017
	100MG	A209480 003	Dec 06, 2017
	200MG	A209480 004	Dec 06, 2017

TABLET, ORALLY DISINTEGRATING; ORAL

FAZACLO ODT

+ JAZZ	12.5MG **	N021590 004	May 30, 2007
+	25MG **	N021590 001	Feb 10, 2004
	50MG **	N021590 003	Jun 03, 2005
+	100MG **	N021590 002	Feb 10, 2004
+	150MG **	N021590 005	Jul 09, 2010
+	200MG **	N021590 006	Jul 09, 2010

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENERGAN VC W/ CODEINE			
+ ANI PHARMS	10MG/5ML; 5MG/5ML; 6.25MG/5ML **	N008306 005	Apr 02, 1984
PHERAZINE VC W/ CODEINE			
HALSEY	10MG/5ML; 5MG/5ML; 6.25MG/5ML	A088870 001	Mar 02, 1987
PROMETH VC W/ CODEINE			
NOSTRUM LABS INC	10MG/5ML; 5MG/5ML; 6.25MG/5ML	A088764 001	Oct 31, 1984
PROMETHAZINE VC W/ CODEINE			
CENCI	10MG/5ML; 5MG/5ML; 6.25MG/5ML	A088816 001	Nov 22, 1985
WOCKHARDT	10MG/5ML; 5MG/5ML; 6.25MG/5ML	A088896 001	Jan 04, 1985

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENERGAN W/ CODEINE			
+ ANI PHARMS	10MG/5ML; 6.25MG/5ML **	N008306 004	Apr 02, 1984
PHERAZINE W/ CODEINE			
HALSEY	10MG/5ML; 6.25MG/5ML	A088739 001	Dec 23, 1988
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE			
+ ACTAVIS MID ATLANTIC	10MG/5ML; 6.25MG/5ML	A088763 001	Oct 31, 1984
PHARM ASSOC	10MG/5ML; 6.25MG/5ML	A089647 001	Dec 22, 1988
PROMETHAZINE W/ CODEINE			
CENCI	10MG/5ML; 6.25MG/5ML	A088814 001	Nov 22, 1985

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

ACTIFED W/ CODEINE			
GLAXOSMITHKLINE	10MG/5ML; 30MG/5ML; 1.25MG/5ML **	N012575 003	Apr 04, 1984
TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES W/ CODEINE			
CENCI	10MG/5ML; 30MG/5ML; 1.25MG/5ML	A089018 001	Jul 23, 1986
TRIPROLIDINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE AND CODEINE PHOSPHATE			
WOCKHARDT	10MG/5ML; 30MG/5ML; 1.25MG/5ML	A088833 001	Nov 16, 1984

CODEINE SULFATE

SOLUTION; ORAL

CODEINE SULFATE			
HIKMA	30MG/5ML	N202245 001	Jun 30, 2011

COLCHICINE; PROBENECID

TABLET; ORAL

COLBENEMID			
+ MERCK	0.5MG; 500MG **	N012383 001	
PROBEN-C			
WATSON LABS	0.5MG; 500MG	A085552 001	
PROBENECID AND COLCHICINE			
ANI PHARMS	0.5MG; 500MG	A083734 001	
BEECHAM	0.5MG; 500MG	A084321 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

COLCHICINE; PROBENECID

TABLET; ORAL

PROBENECID AND COLCHICINE

IMPAX LABS

0.5MG;500MG

A083720 002

SANDOZ

0.5MG;500MG

A086130 001

PROBENECID W/ COLCHICINE

LEDERLE

0.5MG;500MG

A086954 001

WATSON LABS

0.5MG;500MG

A083221 001

COLESEVELAM HYDROCHLORIDE

BAR, CHEWABLE; ORAL

WELCHOL

+ COSETTE

3.75GM

N210895 001 Apr 03, 2019

CAPSULE; ORAL

WELCHOL

COSETTE

375MG

N021141 001 May 26, 2000

FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

WATSON LABS INC

1.875GM/PACKET

A202178 001 Sep 01, 2020

3.75GM/PACKET

A202178 002 Sep 01, 2020

WELCHOL

+ COSETTE

1.875GM/PACKET **

N022362 001 Oct 02, 2009

TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

CHARTWELL RX

625MG

A201354 001 Dec 17, 2020

INVAGEN PHARMS

625MG

A212602 001 Apr 20, 2020

UNITED RES LABS

625MG

A213456 001 Jan 21, 2022

WATSON LABS INC

625MG

A200830 001 Sep 02, 2020

COLISTIN SULFATE

SUSPENSION; ORAL

COLY-MYCIN S

PARKE DAVIS

EQ 25MG BASE/5ML

N050355 001

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

VAPRISOL

CUMBERLAND

20MG/4ML (5MG/ML)

N021697 001 Dec 29, 2005

COPANLISIB DIHYDROCHLORIDE

POWDER; INTRAVENOUS

ALIQOPA

+ BAYER HEALTHCARE

60MG/VIAL

N209936 001 Sep 14, 2017

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

CU-7

GD SEARLE LLC

89MG

N017408 001

TATUM-T

GD SEARLE LLC

120MG

N018205 001

CORTICOTROPIN

INJECTABLE; INJECTION

ACTH

PARKEDALE

25 UNITS/VIAL

N008317 002

40 UNITS/VIAL

N008317 004

ACTHAR

SANOFI AVENTIS US

25 UNITS/VIAL

N007504 002

40 UNITS/VIAL

N007504 003

ACTHAR GEL

MALLINCKRODT ARD

40 UNITS/ML

N008372 006

CORTICOTROPIN

ORGANICS LAGRANGE

40 UNITS/ML

N010831 001

80 UNITS/ML

N010831 002

WATSON LABS

40 UNITS/VIAL

A088772 001 Nov 21, 1984

PURIFIED CORTROPHIN GEL

ANI PHARMS

40 UNITS/ML

N008975 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CORTICOTROPIN-ZINC HYDROXIDE

INJECTABLE; INJECTION

CORTROPHIN-ZINC

ANI PHARMS 40 UNITS/ML N009854 001

CORTISONE ACETATE

INJECTABLE; INJECTION

CORTISONE ACETATE

PHARMACIA AND UPJOHN 25MG/ML N008126 002

WATSON LABS 25MG/ML A083147 003

25MG/ML A085677 001

50MG/ML A083147 004

50MG/ML A085677 002

CORTONE

MERCK 25MG/ML N007110 002

50MG/ML N007110 003

TABLET; ORAL

CORTISONE ACETATE

BARR 25MG A083471 001

CHARTWELL MOLECULAR 25MG A080694 001

ELKINS SINN 25MG A080836 001

EVERYLIFE 25MG A084246 001

HEATHER 25MG A085736 001

HIKMA INTL PHARMS 25MG A080776 002

IMPAX LABS 25MG N009458 001

INWOOD LABS 25MG A080731 001

IVAX SUB TEVA PHARMS 25MG A080630 001

25MG A083536 001

PANRAY 5MG N008284 002

25MG N008284 001

PHARMACIA AND UPJOHN 5MG N008126 003

10MG N008126 004

25MG N008126 001

PUREPAC PHARM 25MG A080493 001

VITARINE 25MG A080333 001

WATSON LABS 25MG A085884 001

WHITEWORTH TOWN PLSN 25MG A080341 001

CORTONE

+ MERCK 25MG ** N007750 003

COSYNTROPIN

SOLUTION; INTRAVENOUS

COSYNTROPIN

SANDOZ 0.25MG/ML (0.25MG/ML) N022028 001 Feb 21, 2008

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION

INTAL

KING PHARMS LLC 0.8MG/INH N018887 001 Dec 05, 1985

CAPSULE; INHALATION

INTAL

+ SANOFI AVENTIS US 20MG ** N016990 001

CAPSULE; ORAL

GASTROCROM

UCB INC 100MG N019188 001 Dec 22, 1989

CONCENTRATE; ORAL

CROMOLYN SODIUM

GENERA PHARMS 100MG/5ML A090954 001 Dec 18, 2009

SOLUTION; INHALATION

CROMOLYN SODIUM

ACTAVIS MID ATLANTIC 10MG/ML A075067 001 Jul 19, 1999

BAUSCH 10MG/ML A075585 001 Dec 21, 2000

EUGIA PHARMA 10MG/ML A074209 001 Apr 26, 1994

HIKMA 10MG/ML A075333 001 Apr 30, 2002

ROXANE 10MG/ML A075175 001 Sep 30, 1999

WATSON LABS 10MG/ML A076469 001 Jun 17, 2005

INTAL

+ KING PHARMS LLC 10MG/ML ** N018596 001 May 28, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CROMOLYN SODIUM

SOLUTION/DROPS;OPHTHALMIC

CROLOM					
BAUSCH AND LOMB	4%	A074443	001	Jan 30, 1995	
CROMOLYN SODIUM					
AKORN	4%	A074706	001	Apr 29, 1998	
APOTEX INC	4%	A075615	001	Jan 26, 2001	
CROMOPTIC					
KING PHARMS	4%	A075088	001	Apr 27, 1999	
OPTICROM					
+ ALLERGAN	4% **	N018155	001	Oct 03, 1984	
SPRAY, METERED;NASAL					
CROMOLYN SODIUM					
ACTAVIS MID ATLANTIC	5.2MG/SPRAY	A074800	001	Jul 26, 2001	
HH AND P	5.2MG/SPRAY	A077976	001	Sep 07, 2007	
PERRIGO	5.2MG/SPRAY	A075427	001	Dec 12, 2001	
NASALCROM					
+ BLACKSMITH BRANDS	5.2MG/SPRAY **	N020463	001	Jan 03, 1997	

CROTAMITON

CREAM;TOPICAL

EURAX					
+ JOURNEY	10%	N006927	001		

CRYPTENAMINE ACETATES

INJECTABLE; INJECTION

UNITENSEN					
MEDPOINTE PHARM HLC	260CSR UNIT/ML	N008814	001		

CRYPTENAMINE TANNATES

TABLET;ORAL

UNITENSEN					
MEDPOINTE PHARM HLC	260CSR UNIT	N009217	001		

CUPRIC SULFATE

INJECTABLE; INJECTION

CUPRIC SULFATE					
+ ABRAXIS PHARM	EQ 0.4MG COPPER/ML **	N019350	001	May 05, 1987	

CYANOCOBALAMIN

GEL, METERED;NASAL

NASCOBAL					
PAR PHARM	0.5MG/INH	N019722	001	Nov 05, 1996	

INJECTABLE; INJECTION

BERUBIGEN					
PHARMACIA AND UPJOHN	1MG/ML	N006798	001		
BETALIN 12					
LILLY	0.1MG/ML	A080855	001		
	1MG/ML	A080855	002		
COBAVITE					
WATSON LABS	0.1MG/ML	A083013	001		
	1MG/ML	A083064	001		
CYANOCOBALAMIN					
ABRAXIS PHARM	0.03MG/ML	A080510	003		
	0.1MG/ML	A080510	001		
	1MG/ML	A080510	002		
AKORN	1MG/ML	A087969	001	Nov 10, 1983	
DELL LABS	0.03MG/ML	A080689	001		
	0.1MG/ML	A080689	002		
	1MG/ML	A080689	003		
DR REDDYS	0.1MG/ML	A080573	002		
	1MG/ML	A080573	001		
FRESENIUS KABI USA	0.1MG/ML	A080557	002		
LUITPOLD	0.03MG/ML	A080668	001		
LYPHOMED	1MG/ML	A083075	001		
MYLAN INSTITUTIONAL	1MG/ML	A040451	001	Sep 23, 2003	
SANOFI AVENTIS US	1MG/ML	A080564	001		
SOLOPAK	1MG/ML	A087551	001	Feb 29, 1984	
WARNER CHILCOTT	1MG/ML	N007085	002		
WATSON LABS	0.1MG/ML	A083120	001		
	1MG/ML	A083120	002		
WYETH AYERST	0.1MG/ML	A080554	001		

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN	1MG/ML	A080554	002	
XIROMED	1MG/ML	A215046	001	Aug 20, 2021
REDISOL				
MERCCK	1MG/ML	N006668	010	
RUBIVITE				
BEL MAR	0.03MG/ML	N010791	004	
	0.05MG/ML	N010791	001	
	0.1MG/ML	N010791	002	
	0.12MG/ML	N010791	005	
	1MG/ML	N010791	003	
RUBRAMIN PC				
BRISTOL MYERS SQUIBB	0.1MG/ML	N006799	002	
+	1MG/ML **	N006799	004	
+	1MG/ML **	N006799	010	Apr 28, 1988
RUVITE				
SAVAGE LABS	1MG/ML	A080570	002	
VI-TWEL				
BAYER HLTHCARE	1MG/ML	N007012	002	
SPRAY, METERED; NASAL				
CALOMIST				
PAR PHARM	25MCG/SPRAY	N022102	001	Jul 27, 2007
TABLET; ORAL				
CYANOCOBALAMIN				
WEST WARD	1MG	A084264	001	

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; CYANOCOBALAMIN CO-58

N/A; N/A

DICOPAC KIT				
GE HEALTHCARE	N/A; N/A; N/A	N017406	001	

CYANOCOBALAMIN; TANNIC ACID; ZINC ACETATE

INJECTABLE; INJECTION

DEPINAR				
ARMOUR PHARM	0.5MG/ML; 2.3MG/ML; 1MG/ML	N011208	001	

CYCLACILLIN

FOR SUSPENSION; ORAL

CYCLAPEN-W				
WYETH AYERST	125MG/5ML	N050508	001	
	250MG/5ML	N050508	002	
	500MG/5ML	N050508	003	

TABLET; ORAL

CYCLACILLIN				
TEVA	250MG	A062895	001	Aug 04, 1988
	500MG	A062895	002	Aug 04, 1988
CYCLAPEN-W				
WYETH AYERST	250MG	N050509	001	
	500MG	N050509	002	

CYCLIZINE LACTATE

INJECTABLE; INJECTION

MAREZINE				
GLAXOSMITHKLINE	50MG/ML	N009495	001	

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE				
APOTEX	15MG	A206703	001	Jul 24, 2018
	30MG	A206703	002	Jul 24, 2018

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE				
CHARTWELL RX	5MG	A072854	002	Feb 03, 2006
	10MG	A072854	001	Nov 19, 1991
PLIVA	10MG	A074421	001	Sep 29, 1995
RISING	5MG	A073144	002	Feb 03, 2006
	7.5MG	A073144	003	Mar 25, 2013
	10MG	A073144	001	May 30, 1991
SANDOZ	10MG	A073683	001	Feb 26, 1993
WATSON LABS	10MG	A073143	001	Nov 27, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

	10MG	A074436 001	Nov 30, 1994
FLEXERIL			
+	JANSSEN RES AND DEV 5MG **	N017821 001	
+	10MG **	N017821 002	

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AK-PENTOLATE

AKORN	1%	A085555 001	
CYCLOPENTOLATE HYDROCHLORIDE			
AKORN	0.5%	A205937 001	Dec 09, 2015
ALCON PHARMS LTD	1%	A089162 001	Jan 24, 1991
SOLA BARNES HIND	1%	A084150 001	
	1%	A084863 001	

PENTOLAIR

PHARMAFAIR	0.5%	A088643 001	Feb 09, 1987
	1%	A088150 001	Feb 25, 1983

CYCLOPHOSPHAMIDE

CAPSULE; ORAL

CYCLOPHOSPHAMIDE

ANI PHARMS	25MG	A207014 001	Mar 19, 2018
	50MG	A207014 002	Mar 19, 2018

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

BAXTER HLTHCARE	100MG/VIAL	A088371 001	Jul 03, 1986
	200MG/VIAL	A088372 001	Jul 03, 1986
	500MG/VIAL	A088373 001	Jul 03, 1986
	1GM/VIAL	A088374 001	Sep 24, 1986

CYTOXAN

+	BAXTER HLTHCARE 100MG/VIAL **	N012142 001	
+	200MG/VIAL **	N012142 002	

CYTOXAN (LYOPHILIZED)

+	BAXTER HLTHCARE 500MG/VIAL **	N012142 003	
+	500MG/VIAL **	N012142 008	Jan 04, 1984
+	1GM/VIAL **	N012142 004	Aug 30, 1982
+	1GM/VIAL **	N012142 010	Sep 24, 1985
+	2GM/VIAL **	N012142 005	Aug 30, 1982
+	2GM/VIAL **	N012142 009	Dec 10, 1985

LYOPHILIZED CYTOXAN

+	BAXTER HLTHCARE 100MG/VIAL **	N012142 006	Dec 05, 1985
+	200MG/VIAL **	N012142 007	Dec 10, 1985

NEOSAR

BEDFORD	100MG/VIAL	A087442 001	Feb 16, 1982
	200MG/VIAL	A087442 002	Feb 16, 1982
	500MG/VIAL	A087442 003	Feb 16, 1982
	1GM/VIAL	A087442 004	Jul 08, 1983
	2GM/VIAL	A087442 005	Mar 30, 1989
TEVA PARENTERAL	100MG/VIAL	A040015 001	Apr 29, 1993
	200MG/VIAL	A040015 002	Apr 29, 1993
	500MG/VIAL	A040015 003	Apr 29, 1993
	1GM/VIAL	A040015 004	Apr 29, 1993
	2GM/VIAL	A040015 005	Apr 29, 1993

SOLUTION; INTRAVENOUS

CYCLOPHOSPHAMIDE

+	NEVAKAR INJECTABLES 500MG/2.5ML (200MG/ML)	N217651 001	Jun 28, 2023
+	1GM/5ML (200MG/ML)	N217651 002	Jun 28, 2023

TABLET; ORAL

CYCLOPHOSPHAMIDE

ROXANE	25MG	A040032 001	Aug 17, 1999
	50MG	A040032 002	Aug 17, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYCLOSPORINE

CAPSULE; ORAL

GENGRAF

ABBVIE 50MG A065003 002 May 12, 2000

NEORAL

+ NOVARTIS 50MG ** N050715 003 Jul 14, 1995

SOLUTION; ORAL

CYCLOSPORINE

DR REDDYS LABS SA 100MG/ML A065054 001 Dec 18, 2001

PHARM ASSOC 100MG/ML A065167 001 Jan 05, 2005

PHARMOBEDIANT CNSLTG 100MG/ML A065133 001 Sep 17, 2004

CYCLOTHIAZIDE

TABLET; ORAL

ANHYDRON

LILLY 2MG N013157 002

FLUIDIL

PHARMACIA AND UPJOHN 2MG N018173 001

CYCRIMINE HYDROCHLORIDE

TABLET; ORAL

PAGITANE

LILLY 1.25MG N008951 001

2.5MG N008951 002

CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

+ ACTAVIS MID ATLANTIC 2MG/5ML ** A086833 001

HALSEY 2MG/5ML A089199 001 Jul 03, 1986

MORTON GROVE 2MG/5ML A087001 001 Nov 04, 1982

NASKA 2MG/5ML A089021 001 Dec 21, 1987

TRIS PHARMA INC 2MG/5ML A205431 001 Dec 21, 2021

PERIACTIN

+ MERCK 2MG/5ML ** N013220 002

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

AM THERAP 4MG A088798 001 Feb 15, 1985

ASCOT 4MG A087685 001 Oct 25, 1982

DURAMED PHARMS BARR 4MG A088232 001 Oct 25, 1983

FOSUN PHARMA 4MG A086808 001

HALSEY 4MG A089057 001 Jul 03, 1986

+ HERITAGE PHARMA 4MG A087056 001

KV PHARM 4MG A086737 001

MD PHARM 4MG A087566 001 Nov 10, 1982

MYLAN 4MG A086678 001

PIONEER PHARMS 4MG A087839 001 Feb 08, 1984

PLIVA 4MG A088205 001 Jul 26, 1983

RISING 4MG A207783 001 Dec 29, 2016

STRIDES PHARMA 4MG A087129 001

SUPERPHARM 4MG A087405 001

VITARINE 4MG A087284 001

WATSON LABS 4MG A085245 001

4MG A086165 001

4MG A086580 001

PERIACTIN

+ MERCK 4MG ** N012649 001

CYSTEINE HYDROCHLORIDE

INJECTABLE; INJECTION

CYSTEINE HYDROCHLORIDE

+ HOSPIRA 7.25% ** N019523 001 Oct 22, 1986

SOLUTION; INTRAVENOUS

NOURESS

+ BAXTER HLTHCARE CORP 500MG/10ML (50MG/ML) N212535 001 Dec 13, 2019

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

MEITHEAL	20MG/ML	A206189 001	Jun 26, 2020
	20MG/ML	A206190 001	Nov 09, 2017
MYLAN LABS LTD	20MG/ML	A200916 001	Dec 13, 2011
RISING	20MG/ML	A200914 001	Dec 13, 2011
+ TEVA PARENTERAL	100MG/VIAL **	N016793 001	
+	500MG/VIAL **	N016793 002	
+	1GM/VIAL **	N016793 003	Dec 21, 1987
+	2GM/VIAL **	N016793 004	Dec 21, 1987
CYTOSAR-U			
TEVA PHARMS USA	100MG/VIAL	A075206 001	Dec 30, 1998
	500MG/VIAL	A075206 002	Dec 30, 1998
	1GM/VIAL	A075206 004	Dec 30, 1998
	2GM/VIAL	A075206 003	Dec 30, 1998

INJECTABLE, LIPOSOMAL; INJECTION

DEPOCYT

+ PACIRA PHARMS INC	10MG/ML	N021041 001	Apr 01, 1999
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DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

ABRAXIS PHARM	100MG/VIAL	A070962 001	Aug 28, 1986
	200MG/VIAL	A070990 001	Aug 28, 1986
DTIC-DOME			
+ BAYER HLTHCARE	100MG/VIAL **	N017575 001	
+	200MG/VIAL **	N017575 002	

DACLATASVIR DIHYDROCHLORIDE

TABLET; ORAL

DAKLINZA

+ BRISTOL-MYERS SQUIBB	EQ 30MG BASE	N206843 001	Jul 24, 2015
+	EQ 60MG BASE	N206843 002	Jul 24, 2015
+	EQ 90MG BASE	N206843 003	Apr 13, 2016

DACTINOMYCIN

INJECTABLE; INJECTION

DACTINOMYCIN

AM REGENT	0.5MG/VIAL	A202562 001	Aug 23, 2013
HIKMA	0.5MG/VIAL	A090304 001	Mar 16, 2010

DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

DALFAMPRIDINE

HIKMA	10MG	A206646 001	Oct 24, 2018
RISING	10MG	A206858 001	Jul 06, 2020

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; INTRAVENOUS

SYNERCID

KING PHARMS	420MG/VIAL; 180MG/VIAL	N050748 002	Aug 24, 2000
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DALTEPARIN SODIUM

INJECTABLE; INJECTION

FRAGMIN

PFIZER	7,500 IU/0.75ML	N020287 008	Apr 04, 2002
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INJECTABLE; SUBCUTANEOUS

FRAGMIN

PFIZER	10,000IU/0.4ML (25,000IU/ML)	N020287 002	May 01, 2007
	95,000IU/9.5ML (10,000IU/ML)	N020287 007	Apr 04, 2002

DANAPAROID SODIUM

INJECTABLE; INJECTION

ORGARAN

ASPEN GLOBAL INC	750 UNITS/0.6ML	N020430 001	Dec 24, 1996
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DANAZOL

CAPSULE; ORAL

DANAZOL

AM THERAP 200MG A071569 001 Dec 30, 1987

DANOCRINE

SANOFI AVENTIS US 50MG ** N017557 003

100MG ** N017557 004

200MG ** N017557 002

DANTROLENE SODIUM

INJECTABLE; INJECTION

DANTROLENE SODIUM

EUGIA PHARMA SPECLTS 20MG/VIAL A205239 001 Feb 18, 2016

DAPAGLIFLOZIN

TABLET; ORAL

DAPAGLIFLOZIN

ZYDUS PHARMS 5MG A211582 001 Feb 22, 2022

10MG A211582 002 Feb 22, 2022

DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

QTERNMET XR

+ ASTRAZENECA AB 2.5MG;1GM;EQ 2.5MG BASE N210874 001 May 02, 2019

+ 5MG;1GM;EQ 2.5MG BASE N210874 002 May 02, 2019

+ 5MG;1GM;EQ 5MG BASE N210874 003 May 02, 2019

+ 10MG;1GM;EQ 5MG BASE N210874 004 May 02, 2019

DAPIPRAZOLE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DAPIPRAZOLE HYDROCHLORIDE

+ FERA PHARMS 0.5% ** N019849 001 Dec 31, 1990

WOODWARD 0.5% A204902 001 May 30, 2019

DAPSONE

TABLET; ORAL

DAPSONE

ALVOGEN 25MG A205429 001 Jan 07, 2016

100MG A205429 002 Jan 07, 2016

CHARTWELL RX 25MG A204074 001 May 10, 2016

100MG A204074 002 May 10, 2016

TARO 25MG A209430 001 Mar 01, 2019

100MG A209430 002 Mar 01, 2019

DAPTOMYCIN

POWDER; INTRAVENOUS

CUBICIN

CUBIST PHARMS LLC 250MG/VIAL ** N021572 001 Sep 12, 2003

+ 500MG/VIAL ** N021572 002 Sep 12, 2003

CUBICIN RF

+ CUBIST PHARMS LLC 500MG/VIAL ** N021572 003 Jul 06, 2016

DAPTOMYCIN

HOSPIRA 500MG/VIAL A202857 001 Sep 12, 2014

DAPZURA RT

+ BAXTER HLTHCARE CORP 500MG/VIAL ** N213645 001 Jan 25, 2022

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

DARIFENACIN HYDROBROMIDE

ANCHEN PHARMS EQ 7.5MG BASE A091190 001 Mar 13, 2015

EQ 15MG BASE A091190 002 Mar 13, 2015

JUBILANT GENERICS EQ 7.5MG BASE A205550 001 Oct 12, 2016

EQ 15MG BASE A205550 002 Oct 12, 2016

XIROMED EQ 7.5MG BASE A209571 002 Oct 22, 2019

EQ 15MG BASE A209571 001 Oct 22, 2019

ENABLEX

+ ABBVIE EQ 7.5MG BASE ** N021513 001 Dec 22, 2004

+ EQ 15MG BASE ** N021513 002 Dec 22, 2004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DARUNAVIR

TABLET;ORAL

PREZISTA

+	JANSSEN PRODS	300MG **	N021976	001	Jun 23, 2006
+		400MG **	N021976	003	Oct 21, 2008

DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR

TABLET;ORAL

VIEKIRA PAK (COPACKAGED)

+	ABBVIE	EQ 250MG BASE;12.5MG, 75MG, 50MG	N206619	001	Dec 19, 2014
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DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, EXTENDED RELEASE;ORAL

VIEKIRA XR

+	ABBVIE	EQ 200MG BASE;8.33MG;50MG;33.33MG **	N208624	001	Jul 22, 2016
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DASATINIB

TABLET;ORAL

DASATINIB

APOTEX

		20MG	A202103	001	Jun 10, 2016
		50MG	A202103	002	Jun 10, 2016
		70MG	A202103	003	Jun 10, 2016
		80MG	A203180	001	Nov 23, 2021
		100MG	A202103	004	Jun 10, 2016
		140MG	A203180	002	Nov 23, 2021

DAUNORUBICIN CITRATE

INJECTABLE, LIPOSOMAL;INJECTION

DAUNOXOME

GALEN (UK)

		EQ 2MG BASE/ML	N050704	002	Apr 08, 1996
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DAUNORUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

CERUBIDINE

SANOFI AVENTIS US

		EQ 20MG BASE/VIAL	A061876	001	
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WYETH AYERST

		EQ 20MG BASE/VIAL **	N050484	001	
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DAUNORUBICIN HYDROCHLORIDE

HISUN PHARM HANGZHOU

		EQ 20MG BASE/VIAL	A206195	001	Apr 25, 2019
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TEVA PARENTERAL

		EQ 20MG BASE/VIAL	A064212	001	Jun 23, 1998
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		EQ 50MG BASE/VIAL	A064212	002	May 03, 1999
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DECAMETHONIUM BROMIDE

INJECTABLE;INJECTION

SYNCURINE

GLAXOSMITHKLINE

		1MG/ML	N006931	002	
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DECITABINE

INJECTABLE;INTRAVENOUS

DACOGEN

+ OTSUKA

		50MG/VIAL	N021790	001	May 02, 2006
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DECITABINE

CIPLA

		50MG/VIAL	A208601	001	Nov 16, 2017
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DEFERASIROX

GRANULE;ORAL

DEFERASIROX

AMNEAL

		90MG	A214194	003	Aug 02, 2021
		180MG	A214194	001	Feb 09, 2021
		360MG	A214194	002	Feb 09, 2021

TABLET;ORAL

DEFERASIROX

CIPLA

		90MG	A211852	001	Feb 11, 2020
		180MG	A211852	003	Jun 15, 2020
		360MG	A211852	002	Feb 11, 2020
		90MG	A210727	001	Dec 27, 2019
		180MG	A210727	003	Jun 15, 2020
		360MG	A210727	002	Dec 27, 2019
		90MG	A209223	001	Nov 25, 2019
		180MG	A209223	003	Apr 24, 2020
		360MG	A209223	002	Nov 25, 2019

TABLET, FOR SUSPENSION;ORAL

DEFERASIROX

TEVA PHARMS USA

		125MG	A207124	001	Sep 23, 2022
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEFERASIROXTABLET, FOR SUSPENSION;ORAL
DEFERASIROX

	250MG	A207124 002	Sep 23, 2022
	500MG	A207124 003	Sep 23, 2022
ZYDUS PHARMS	125MG	A208882 001	May 05, 2020
	250MG	A208882 002	May 05, 2020
	500MG	A208882 003	May 05, 2020

DEFERIPRONESOLUTION;ORAL
FERRIPROX

+ CHIESI	80MG/ML	N208030 002	Apr 20, 2018
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DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

DR REDDYS	500MG/VIAL	A076806 001	Mar 31, 2006
	2GM/VIAL	A076806 002	Mar 31, 2006
DESFERAL + NOVARTIS	2GM/VIAL **	N016267 002	May 25, 2000

DELAVIRDINE MESYLATE

TABLET;ORAL

RESCRIPTOR

+ VIIV HLTHCARE	100MG	N020705 001	Apr 04, 1997
+	200MG	N020705 002	Jul 14, 1999

DEMECARIUM BROMIDE

SOLUTION/DROPS;OPHTHALMIC

HUMORSOL

MERCK	0.125%	N011860 002	
	0.25%	N011860 001	

DEMECLOCYCLINE HYDROCHLORIDE

CAPSULE;ORAL

DECLOMYCIN

LEDERLE	150MG	N050262 001	
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SYRUP;ORAL

DECLOMYCIN

LEDERLE	75MG/5ML	N050257 001	
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TABLET;ORAL

DECLOMYCIN

COREPHARMA	75MG **	N050261 001	
	150MG **	N050261 002	
	300MG **	N050261 003	

DEMECLOCYCLINE HYDROCHLORIDE

BARR	150MG	A065171 001	Dec 13, 2004
	300MG	A065171 002	Dec 13, 2004
IMPAX LABS	150MG	A065094 001	Mar 22, 2004
	300MG	A065094 002	Mar 22, 2004

DESERPIDINE

TABLET;ORAL

HARMONYL

ABBVIE	0.1MG	N010796 001	
	0.25MG	N010796 002	

DESERPIDINE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

ORETICYL 25

ABBVIE	0.125MG; 25MG	N012148 001	
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ORETICYL 50

ABBVIE	0.125MG; 50MG	N012148 003	
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ORETICYL FORTE

ABBVIE	0.25MG; 25MG	N012148 002	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESERPIDINE; METHYCLOTHIAZIDE

TABLET; ORAL

ENDURONYL

ABBOTT 0.25MG; 5MG

N012775 001

ENDURONYL FORTE

ABBOTT 0.5MG; 5MG

N012775 002

METHYCLOTHIAZIDE AND DESERPIDINE

WATSON LABS 0.25MG; 5MG

A088486 001 Aug 10, 1984

0.5MG; 5MG

A088452 001 Aug 10, 1984

DESIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

PERTOFRANE

SANOFI AVENTIS US 25MG

N013621 001

50MG

N013621 002

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

ANI PHARMS 10MG

A205153 001 Oct 28, 2016

25MG

A071803 002 Dec 08, 1987

25MG

A205153 002 Oct 28, 2016

50MG

A071803 003 Dec 08, 1987

50MG

A205153 003 Oct 28, 2016

75MG

A071803 004 Dec 08, 1987

75MG

A205153 004 Oct 28, 2016

100MG

A071803 001 May 29, 1997

100MG

A205153 005 Oct 28, 2016

150MG

A071803 005 May 29, 1997

150MG

A205153 006 Oct 28, 2016

USL PHARMA 25MG

A071864 001 Sep 09, 1987

50MG

A071865 001 Sep 09, 1987

75MG

A071866 001 Sep 09, 1987

100MG

A071867 001 Sep 09, 1987

DESLANOSIDE

INJECTABLE; INJECTION

CEDILANID-D

NOVARTIS 0.2MG/ML

N009282 002

DESLORATADINE

SOLUTION; ORAL

CLARINEX

+ MERCK SHARP DOHME 0.5MG/ML **

N021300 001 Sep 01, 2004

DESLORATADINE

TARO 0.5MG/ML

A202592 001 Jun 30, 2015

0.5MG/ML

A202936 001 May 26, 2016

TABLET; ORAL

DESLORATADINE

DASH PHARMS 5MG

A078351 001 Feb 10, 2012

PERRIGO 5MG

A078361 001 Dec 22, 2011

SANDOZ 5MG

A078364 001 Dec 03, 2010

SUN PHARM INDS 5MG

A078359 001 Nov 16, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

CLARINEX

+ ORGANON 2.5MG **

N021312 002 Jul 14, 2005

+ 5MG **

N021312 001 Jun 26, 2002

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX D 24 HOUR

+ ORGANON 5MG; 240MG **

N021605 001 Mar 03, 2005

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

FERRING PHARMS INC 0.015MG/ML

N018938 002 Apr 25, 1995

DESMOPRESSIN ACETATE

AM REGENT 0.004MG/ML

A091374 001 Feb 14, 2019

BEDFORD 0.004MG/ML

A074575 001 Feb 18, 2000

HOSPIRA 0.004MG/ML

A075220 001 Aug 28, 2000

DESMOPRESSIN ACETATE PRESERVATIVE FREE

BEDFORD 0.004MG/ML

A074574 001 Feb 18, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESMOPRESSIN ACETATESOLUTION;NASAL
CONCENTRAID

FERRING	0.01%	N019776 001	Dec 26, 1990
DDAVP			
+ FERRING PHARMS INC	0.01%	N017922 001	
DESMOPRESSIN ACETATE			
SUN PHARM INDS	0.01%	A077212 001	Apr 12, 2012

SPRAY, METERED;NASAL

DDAVP			
+ FERRING PHARMS INC	0.01MG/SPRAY **	N017922 002	Feb 06, 1989
DDAVP (NEEDS NO REFRIGERATION)			
+ FERRING PHARMS INC	0.01MG/SPRAY	N017922 003	Aug 07, 1996
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)			
SUN PHARM	0.01MG/SPRAY	A078271 001	Dec 23, 2013
MINIRIN			
+ FERRING	0.01MG/SPRAY	N021333 001	Sep 16, 2002
NOCTIVA			
+ ACERUS PHARMS	0.00083MG/SPRAY	N201656 001	Mar 03, 2017
+	0.00166MG/SPRAY	N201656 002	Mar 03, 2017
STIMATE			
FERRING PHARMS INC	0.15MG/SPRAY	N020355 001	Mar 07, 1994
STIMATE (NEEDS NO REFRIGERATION)			
+ FERRING PHARMS INC	0.15MG/SPRAY	N020355 002	Oct 24, 2007

TABLET;ORAL

DESMOPRESSIN ACETATE			
FERRING	0.1MG	N021795 001	May 08, 2008
	0.2MG	N021795 002	May 08, 2008
IMPAX LABS INC	0.1MG	A077122 001	Jan 25, 2006
	0.2MG	A077122 002	Jan 25, 2006
NATCO PHARMA USA	0.1MG	A200653 001	Jun 27, 2014
	0.2MG	A200653 002	Jun 27, 2014

TABLET;SUBLINGUAL

NOCDURNA			
+ FERRING PHARMS INC	0.0277MG	N022517 001	Jun 21, 2018
+	0.0553MG	N022517 002	Jun 21, 2018

DESOGESTREL; ETHINYL ESTRADIOL

TABLET;ORAL-21

DESOGEN			
ORGANON USA INC	0.15MG;0.03MG	N020071 001	Dec 10, 1992
DESOGESTREL AND ETHINYL ESTRADIOL			
DURAMED PHARMS BARR	0.15MG;0.03MG	A075256 001	Aug 12, 1999
ORTHO-CEPT			
+ JANSSEN PHARMS	0.15MG;0.03MG **	N020301 001	Dec 14, 1992

TABLET;ORAL-28

DESOGEN			
ORGANON USA INC	0.15MG;0.03MG	N020071 002	Dec 10, 1992
DESOGESTREL AND ETHINYL ESTRADIOL			
DR REDDYS LABS SA	0.15MG,N/A;0.02MG,0.01MG	A076916 001	Dec 29, 2008
	0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG	A077182 001	Jan 24, 2006
MYLAN LABS LTD	0.15MG;0.03MG	A202085 001	May 20, 2015
NAARI PTE LTD	0.15MG;0.03MG	A207067 001	Sep 13, 2018
EMOQUETTE			
VINTAGE PHARMS LLC	0.15MG;0.03MG	A076675 001	Feb 25, 2011
MIRCETTE			
+ TEVA BRANDED PHARM	0.15MG,N/A;0.02MG,0.01MG **	N020713 001	Apr 22, 1998
ORTHO-CEPT			
+ JANSSEN PHARMS	0.15MG;0.03MG **	N020301 002	Dec 14, 1992

DESONIDE

AEROSOL, FOAM;TOPICAL

VERDESO			
+ ALMIRALL	0.05%	N021978 001	Sep 19, 2006
GEL;TOPICAL			
DESONATE			
+ LEO PHARMA AS	0.05% **	N021844 001	Oct 20, 2006

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESONIDE

LOTION; TOPICAL				
DESONIDE				
PAI HOLDINGS PHARM	0.05%	A207855	001	Sep 28, 2017
ointment; TOPICAL				
DESONIDE				
PAI HOLDINGS PHARM	0.05%	A212002	001	Mar 12, 2019

DESOXIMETASONE

CREAM; TOPICAL				
DESOXIMETASONE				
AKORN	0.05%	A203787	001	Jan 06, 2017
	0.25%	A203234	001	Jun 12, 2015
TOPICORT				
+ TARO	0.25% **	N017856	001	
TOPICORT LP				
+ TARO	0.05% **	N018309	001	
GEL; TOPICAL				
TOPICORT				
+ TARO	0.05% **	N018586	001	Mar 29, 1982
ointment; TOPICAL				
DESOXIMETASONE				
ALTANA	0.25%	A073440	001	Apr 01, 1998
PAI HOLDINGS PHARM	0.25%	A208101	001	Feb 25, 2016
TOPICORT				
TARO	0.25%	A074286	001	Jun 07, 1996
+ TARO	0.25% **	N018763	001	Sep 30, 1983

DESOXYCORTICOSTERONE ACETATE

INJECTABLE; INJECTION				
DOCA				
ORGANON USA INC	5MG/ML	N001104	001	
PELLET; IMPLANTATION				
PERCORTEN				
NOVARTIS	125MG	N005151	001	

DESOXYCORTICOSTERONE PIVALATE

INJECTABLE; INJECTION				
PERCORTEN				
NOVARTIS	25MG/ML	N008822	001	

DESVENLAFAXINE

TABLET, EXTENDED RELEASE; ORAL				
KHEDEZLA				
OSMOTICA PHARM CORP	50MG	N204683	001	Jul 10, 2013
	100MG	N204683	002	Jul 10, 2013

DESVENLAFAXINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL				
DESVENLAFAXINE				
+ SUN PHARM	EQ 50MG BASE	N205583	001	Jan 28, 2014
+ SUN PHARM	EQ 100MG BASE	N205583	002	Jan 28, 2014
TEVA PHARMS USA	EQ 50MG BASE	N205208	001	Oct 11, 2013
TEVA PHARMS USA	EQ 100MG BASE	N205208	002	Oct 11, 2013

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL				
DESVENLAFAXINE SUCCINATE				
MYLAN	EQ 50MG BASE	A204095	001	Jun 29, 2015
MYLAN	EQ 100MG BASE	A204095	002	Jun 29, 2015

DEXAMETHASONE

AEROSOL; TOPICAL				
AEROSEB-DEX				
ALLERGAN HERBERT	0.01% **	A083296	002	
DECASPRAY				
+ MERCK	0.04% **	N012731	002	
ELIXIR; ORAL				
DECADRON				
MERCK	0.5MG/5ML	N012376	002	
DEXAMETHASONE				
ALPHARMA US PHARMS	0.5MG/5ML	A088997	001	Oct 10, 1986
+ PHARMOBEDIANT CNSLTG	0.5MG/5ML	A088254	001	Jul 27, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE

ELIXIR; ORAL

HEXADROL

ASPEN GLOBAL INC 0.5MG/5ML N012674 001

GEL; TOPICAL

DECADERM

MERCCK 0.1% N013538 001

SUSPENSION/DROPS; OPHTHALMIC

DEXAMETHASONE

WATSON LABS 0.1% A089170 001 May 09, 1989

TABLET; ORAL

DECADRON

+ MERCCK 0.25MG ** N011664 004

+ 0.5MG ** N011664 001

+ 0.75MG ** N011664 002

+ 1.5MG ** N011664 003

+ 4MG ** N011664 005

+ 6MG ** N011664 006 Jul 30, 1982

DEXAMETHASONE

BAUSCH 1.5MG A040700 001 Aug 15, 2008

CHARTWELL RX 1.5MG A085456 001

IMPAX LABS 0.75MG A085376 001

PHOENIX LABS NY 0.75MG A083806 001

PRASCO 0.75MG A080399 001

PVT FORM 0.75MG A083420 001

ROXANE 0.25MG A084614 001

STRIDES PHARMA 0.25MG A088149 001 Apr 28, 1983

SUN PHARM INDUSTRIES 0.25MG A084013 001

0.25MG A084764 001

0.5MG A084084 001

0.5MG A084766 001

0.75MG A084081 001

0.75MG A084765 001

1.5MG A084086 001

1.5MG A084763 001

UPSHER SMITH 0.75MG A087534 001

1.5MG A087533 001

WATSON LABS 0.25MG A085455 001

0.5MG A085458 001

0.75MG A080968 001

0.75MG A084457 001

0.75MG A085818 001

1.5MG A085840 001

WHITEWORTH TOWN PLSN 0.75MG A084327 001

DEXONE 0.5

SOLVAY 0.5MG A084991 001

DEXONE 0.75

SOLVAY 0.75MG A084993 001

DEXONE 1.5

SOLVAY 1.5MG A084990 001

DEXONE 4

SOLVAY 4MG A084992 001

HEXADROL

ASPEN GLOBAL INC 0.5MG N012675 004

0.75MG N012675 007

1.5MG N012675 009

4MG N012675 010

DEXAMETHASONE ACETATE

INJECTABLE; INJECTION

DECADRON-LA

+ MERCCK EQ 8MG BASE/ML ** N016675 001

DEXAMETHASONE ACETATE

WATSON LABS EQ 8MG BASE/ML A084315 001

WATSON LABS TEVA EQ 16MG BASE/ML A087711 001 May 24, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE SODIUM PHOSPHATE

AEROSOL;NASAL

DEXACORT

UCB INC

EQ 0.1MG PHOSPHATE/INH

N014242 001

AEROSOL, METERED;INHALATION

DEXACORT

UCB INC

EQ 0.1MG PHOSPHATE/INH

N013413 001

CREAM;TOPICAL

DECADRON

MERCCK

EQ 0.1% PHOSPHATE

N011983 002

INJECTABLE;INJECTION

DECADRON

+ MERCCK

EQ 4MG PHOSPHATE/ML **

N012071 002

+

EQ 24MG PHOSPHATE/ML **

N012071 004

DEXACEN-4

CENT PHARMS

EQ 4MG PHOSPHATE/ML

A084342 001

DEXAMETHASONE

ABRAXIS PHARM

EQ 4MG PHOSPHATE/ML

A088448 001 Jan 25, 1984

FRESENIUS KABI USA

EQ 10MG PHOSPHATE/ML

A088469 001 Jan 25, 1984

DEXAMETHASONE SODIUM PHOSPHATE

AKORN

EQ 4MG PHOSPHATE/ML

A084493 001

BEL MAR

EQ 4MG PHOSPHATE/ML

A084752 001

DELL LABS

EQ 4MG PHOSPHATE/ML

A083161 001

DR REDDYS

EQ 4MG PHOSPHATE/ML

A089169 001 Apr 09, 1986

INTL MEDICATION

EQ 20MG PHOSPHATE/ML

A088522 001 Feb 17, 1984

+ LUITPOLD

EQ 4MG PHOSPHATE/ML

A087440 001 Jul 21, 1982

LYPHOMED

EQ 4MG PHOSPHATE/ML

A087065 001

TEVA PARENTERAL

EQ 4MG PHOSPHATE/ML

A081125 001 Aug 31, 1990

EQ 10MG PHOSPHATE/ML

A081126 001 Aug 31, 1990

WATSON LABS

EQ 4MG PHOSPHATE/ML

A083702 001

EQ 4MG PHOSPHATE/ML

A084355 001

EQ 10MG PHOSPHATE/ML

A087668 001 Jul 01, 1982

EQ 24MG PHOSPHATE/ML

A085606 001

WYETH AYERST

EQ 4MG PHOSPHATE/ML

A085641 001

HEXADROL

+ ASPEN GLOBAL INC

EQ 4MG PHOSPHATE/ML **

N014694 002

+

EQ 10MG PHOSPHATE/ML **

N014694 003

EQ 20MG PHOSPHATE/ML

N014694 004

OINTMENT;OPHTHALMIC

DECADRON

MERCCK

EQ 0.05% PHOSPHATE

N011977 001

DEXAIR

PHARMAFAIR

EQ 0.05% PHOSPHATE

A088071 001 Dec 28, 1982

MAXIDEX

ALCON

EQ 0.05% PHOSPHATE

A083342 001

SOLUTION/DROPS;OPHTHALMIC

DEXAIR

PHARMAFAIR

EQ 0.1% PHOSPHATE

A088433 001 Dec 15, 1983

DEXAMETHASONE SODIUM PHOSPHATE

SOLA BARNES HIND

EQ 0.1% PHOSPHATE

A084170 001

EQ 0.1% PHOSPHATE

A084173 001

SOLUTION/DROPS;OPHTHALMIC, OTIC

DECADRON

+ MERCCK

EQ 0.1% PHOSPHATE

N011984 001

DEXAMETHASONE SODIUM PHOSPHATE

SANDOZ

EQ 0.1% PHOSPHATE

A088771 001 Jan 16, 1985

SOLUTION/DROPS;OTIC

DEXAMETHASONE SODIUM PHOSPHATE

AKORN

EQ 0.1% PHOSPHATE

A084855 001

DEXAMETHASONE SODIUM PHOSPHATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

DECADRON W/ XYLOCAINE

MERCCK

EQ 4MG PHOSPHATE/ML;10MG/ML

N013334 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE SODIUM PHOSPHATE; NEOMYCIN SULFATE

OINTMENT;OPHTHALMIC

NEODECADRON

MERCCK

EQ 0.05% PHOSPHATE;EQ 3.5MG BASE/GM

N050324 001

SOLUTION/DROPS;OPHTHALMIC

NEODECADRON

MERCCK

EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML

N050322 001

NEOMYCIN SULFATE AND DEXAMETHASONE SODIUM PHOSPHATE

BAUSCH AND LOMB

EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML

A064055 001 Oct 30, 1995

NEOMYCIN SULFATE-DEXAMETHASONE SODIUM PHOSPHATE

ALCON PHARMS LTD

EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML

A062714 001 Jul 21, 1986

PHARMAFAIR

EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML

A062539 001 Jan 10, 1985

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

DEXACIDIN

NOVARTIS

0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM

A062566 001 Feb 22, 1985

DEXASPORIN

PHARMAFAIR

0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM

A062411 001 May 16, 1983

SUSPENSION/DROPS;OPHTHALMIC

DEXACIDIN

NOVARTIS

0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

A062544 001 Oct 29, 1984

DEXASPORIN

PHARMAFAIR

0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

A062428 001 May 18, 1983

NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE

ALCON PHARMS LTD

0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

A062721 001 Nov 17, 1986

DEXAMETHASONE; TOBRAMYCIN

SUSPENSION/DROPS;OPHTHALMIC

TOBRAMYCIN AND DEXAMETHASONE

AMNEAL

0.1%;0.3%

A212991 001 Jul 15, 2021

DEXBROMPHENIRAMINE MALEATE

SYRUP;ORAL

DISOMER

SCHERING

2MG/5ML

N011814 002

TABLET;ORAL

DISOMER

SCHERING

2MG

N011814 001

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET;ORAL

DISOPHROL

SCHERING

2MG;60MG

N012394 002

TABLET, EXTENDED RELEASE;ORAL

BROMPHERIL

COPLBY PHARM

6MG;120MG

A089116 001 Jan 22, 1987

DEXBROMPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE SULFATE

AVANTHI INC

6MG;120MG

A078648 001 Feb 27, 2013

DISOBROM

SANDOZ

6MG;120MG

A070770 001 Sep 30, 1991

DISOPHROL

SCHERING PLOUGH

6MG;120MG

N013483 004 Sep 13, 1982

DRIXORAL

+

SCHERING PLOUGH

6MG;120MG **

N013483 003 Sep 13, 1982

RESPORAL

PIONEER PHARMS

6MG;120MG

A089139 001 Jun 16, 1988

DEXCHLORPHENIRAMINE MALEATE

SYRUP;ORAL

POLARAMINE

SCHERING

2MG/5ML

A086837 001 Jul 19, 1982

TABLET;ORAL

DEXCHLORPHENIRAMINE MALEATE

ANI PHARMS

2MG

A088682 001 Jan 17, 1986

POLARAMINE

+

SCHERING

2MG

A086835 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXLANSOPRAZOLETABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL
DEXILANT SOLUTAB

+ TAKEDA PHARMS USA 30MG N208056 001 Jan 26, 2016

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

DEXMEDETOMIDINE HYDROCHLORIDE

AM REGENT EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML) A203773 001 May 12, 2017

PIRAMAL CRITICAL EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML) A214794 001 Sep 03, 2021

SLAYBACK PHARMA LLC EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML) A212791 003 May 08, 2020

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

AUROLIFE PHARMA LLC 5MG A204266 001 Aug 25, 2015

10MG A204266 002 Aug 25, 2015

15MG A204266 003 Aug 25, 2015

20MG A204266 004 Dec 21, 2015

25MG A204266 005 May 04, 2021

30MG A202580 001 Aug 28, 2013

35MG A204266 006 May 04, 2021

40MG A204266 007 Aug 25, 2015

TABLET;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

BIONPHARMA 5MG A209754 001 Mar 24, 2020

10MG A209754 002 Mar 24, 2020

LANNETT CO INC 2.5MG A209468 001 Sep 25, 2017

5MG A209468 002 Sep 25, 2017

10MG A209468 003 Sep 25, 2017

TEVA PHARMS 2.5MG A077107 003 Jan 29, 2007

5MG A077107 001 Jan 29, 2007

10MG A077107 002 Jan 29, 2007

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE;INJECTION

TOTECT

+ CLINIGEN EQ 500MG BASE/VIAL ** N022025 001 Sep 06, 2007

ZINECARD

+ PFIZER EQ 250MG BASE/VIAL N020212 001 May 26, 1995

+ EQ 500MG BASE/VIAL N020212 002 May 26, 1995

DEXTROAMPHETAMINE SULFATE

CAPSULE;ORAL

DEXAMPEX

TEVA 15MG A085355 001

CAPSULE, EXTENDED RELEASE;ORAL

DEXTROAMPHETAMINE SULFATE

ABLE 5MG A076814 001 Aug 25, 2004

10MG A076814 002 Aug 25, 2004

15MG A076814 003 Aug 25, 2004

DR REDDYS LABS SA 5MG A076137 001 Jan 18, 2002

10MG A076137 002 Jan 18, 2002

15MG A076137 003 Jan 18, 2002

MYLAN 5MG A206735 001 Jan 27, 2016

10MG A206735 002 Jan 27, 2016

15MG A206735 003 Jan 27, 2016

NESHER PHARMS 5MG A209111 001 Jun 27, 2017

10MG A209111 002 Jun 27, 2017

15MG A209111 003 Jun 27, 2017

STRIDES PHARMA 5MG A205077 001 Jun 21, 2019

10MG A205077 002 Jun 21, 2019

15MG A205077 003 Jun 21, 2019

ELIXIR;ORAL

DEXEDRINE

GLAXOSMITHKLINE 5MG/5ML ** A083902 001

TABLET;ORAL

DEXAMPEX

TEVA 5MG A083735 001

10MG A083735 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXEDRINE

+ GLAXOSMITHKLINE 5MG **

A084935 001

DEXTROAMPHETAMINE SULFATE

ANI PHARMS

5MG

A085370 001

EPIC PHARMA LLC

5MG

A090652 001 Mar 07, 2014

10MG

A090652 002 Mar 07, 2014

HALSEY

10MG

A083930 001

LANNETT

5MG

A083903 001

10MG

A083903 003

15MG

A085652 001

MAST MM

5MG

A086521 001

NESHER PHARMS

5MG

A040365 001 Oct 31, 2002

5MG

A206588 001 Mar 28, 2016

10MG

A040367 001 Oct 31, 2002

10MG

A206588 002 Mar 28, 2016

PUREPAC PHARM

5MG

A084125 001

SANDOZ

10MG

A085371 001

STRIDES PHARMA

5MG

A040299 001 May 13, 1999

VITARINE

5MG

A084986 001

10MG

A085892 001

DEXTROSTAT

+ SHIRE

5MG **

A084051 001

+

10MG **

A084051 002

FERNDEX

FERNDALE LABS

5MG

A084001 001

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHERAZINE DM

HALSEY

15MG/5ML; 6.25MG/5ML

A088913 001 Mar 02, 1987

PROMETH W/ DEXTROMETHORPHAN

G AND W LABS INC

15MG/5ML; 6.25MG/5ML **

A088762 001 Oct 31, 1984

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

+ ANI PHARMS

15MG/5ML; 6.25MG/5ML **

N011265 002 Apr 02, 1984

HIKMA

15MG/5ML; 6.25MG/5ML

A040027 001 Jul 31, 1996

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

B BRAUN

10GM/100ML

N018046 001

MILES

10GM/100ML

N018504 001

DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN

2.5GM/100ML

N018358 001

2.5GM/100ML

N019626 001 Feb 02, 1988

DEXTROSE 20% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

20GM/100ML **

N017521 004

DEXTROSE 30% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

30GM/100ML **

N017521 003

DEXTROSE 38.5% IN PLASTIC CONTAINER

ABBOTT

38.5GM/100ML

N018923 001 Sep 19, 1984

DEXTROSE 40% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

40GM/100ML **

N017521 002

DEXTROSE 5% IN PLASTIC CONTAINER

DHL

5GM/100ML

N019971 001 Sep 28, 1995

+ ICU MEDICAL INC

50MG/ML

N019222 001 Jul 13, 1984

DEXTROSE 50% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

50GM/100ML **

N017521 001

ICU MEDICAL INC

50GM/100ML

N019894 001 Dec 26, 1989

DEXTROSE 60%

B BRAUN

60GM/100ML

N017995 002 Sep 22, 1982

DEXTROSE 60% IN PLASTIC CONTAINER

B BRAUN

60GM/100ML

N017995 001

+ BAXTER HLTHCARE

60GM/100ML **

N017521 005 Mar 26, 1982

60GM/100ML

N020047 002 Jul 02, 1991

HOSPIRA

60GM/100ML

N019346 001 Jan 25, 1985

DEXTROSE 7.7% IN PLASTIC CONTAINER

B BRAUN

7.7GM/100ML

N019626 003 Feb 02, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE

INJECTABLE;INJECTION

DEXTROSE 70% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 70GM/100ML **

N017521 006 Mar 26, 1982

DEXTROSE; MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE;INJECTION

PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5GM/100ML;32MG/100ML;128MG/100ML;234MG/
100ML

N017385 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE;INJECTION

ISOLYTE P W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;31MG/100ML;130MG/100ML;26MG/1
00ML;320MG/100ML

N019025 001 Dec 27, 1984

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE;
SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE;INJECTION

IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA 5GM/100ML;53MG/100ML;100MG/100ML;100MG/
100ML;180MG/100ML;280MG/100ML;16MG/100M
L

N019515 001 May 08, 1986

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE;INJECTION

ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;30MG/100ML;97MG/100ML;220MG/1
00ML;140MG/100ML

N019844 001 Jun 10, 1993

ISOLYTE H W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;30MG/100ML;97MG/100ML;220MG/1
00ML;140MG/100ML

N018273 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM
GLUCONATE

INJECTABLE;INJECTION

ISOLYTE S IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;30MG/100ML;37MG/100ML;370MG/1
00ML;530MG/100ML;500MG/100ML

N019843 001 Aug 09, 1993

ISOLYTE S W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;30MG/100ML;37MG/100ML;370MG/1
00ML;530MG/100ML;500MG/100ML

N018274 001

PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5GM/100ML;30MG/100ML;37MG/100ML;368MG/1
00ML;526MG/100ML;502MG/100ML

N017451 001

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE;INJECTION

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;37MG/100ML

N019699 001 Sep 29, 1989

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;75MG/100ML

N018744 001 Nov 09, 1982

5GM/100ML;75MG/100ML

N019699 002 Sep 29, 1989

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;110MG/100ML

N019699 003 Sep 29, 1989

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;150MG/100ML

N018744 002 Nov 09, 1982

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;220MG/100ML

N018744 003 Nov 09, 1982

5GM/100ML;220MG/100ML

N019699 005 Sep 29, 1989

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 5GM/100ML;300MG/100ML

N018744 004 Nov 09, 1982

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML;224MG/100ML

N018371 003

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML;298MG/100ML

N018371 002

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM LACTATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, MONOBASIC
ANHYDROUS

INJECTABLE;INJECTION

IONOSOL T AND DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA 5GM/100ML;111MG/100ML;256MG/100ML;146MG
/100ML;207MG/100ML

N019514 001 May 08, 1986

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;150MG/100ML;130MG/100ML;280MG	N019870	001	Jun 10, 1993
	/100ML;91MG/100ML			
ISOLYTE M W/ DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;150MG/100ML;130MG/100ML;280MG	N018270	001	
	/100ML;91MG/100ML			

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO. 75 IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5GM/100ML;205MG/100ML;100MG/100ML;120MG	N018840	001	Jun 29, 1983
	/100ML;220MG/100ML			

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.075%				
B BRAUN	5GM/100ML;75MG/100ML;200MG/100ML	N018268	009	
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;150MG/100ML;200MG/100ML	N018268	004	
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;220MG/100ML;200MG/100ML	N018268	005	
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;300MG/100ML;200MG/100ML	N018268	006	
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;75MG/100ML;330MG/100ML	N018268	011	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;150MG/100ML;330MG/100ML	N018268	012	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;220MG/100ML;330MG/100ML	N018268	013	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.30% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;300MG/100ML;330MG/100ML	N018268	014	Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.075%				
B BRAUN	5GM/100ML;75MG/100ML;450MG/100ML	N018268	010	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;150MG/100ML;450MG/100ML	N018268	001	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;220MG/100ML;450MG/100ML	N018268	002	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML;300MG/100ML;450MG/100ML	N018268	003	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5GM/100ML;224MG/100ML;450MG/100ML	N018008	003	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N018008	001	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5GM/100ML;75MG/100ML;450MG/100ML	N018008	002	
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;225MG/100ML **	N018365	002	Jul 05, 1983
+ ICU MEDICAL INC	5GM/100ML;149MG/100ML;225MG/100ML **	N018365	006	Mar 28, 1988
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;300MG/100ML	N018876	001	Jan 17, 1986
	5GM/100ML;149MG/100ML;300MG/100ML	N018876	006	Mar 28, 1988
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;900MG/100ML	N019691	002	Mar 24, 1988
+ ICU MEDICAL INC	5GM/100ML;149MG/100ML;900MG/100ML	N019691	004	Mar 24, 1988
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER				
ICU MEDICAL INC	5GM/100ML;224MG/100ML;225MG/100ML **	N018365	008	Mar 28, 1988
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
ICU MEDICAL INC	5GM/100ML;224MG/100ML;300MG/100ML	N018876	007	Mar 28, 1988
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	5GM/100ML;224MG/100ML;450MG/100ML	N018362	006	Mar 28, 1988
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	5GM/100ML;224MG/100ML;900MG/100ML	N019691	006	Mar 24, 1988
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	5GM/100ML;298MG/100ML;225MG/100ML **	N018365	009	Mar 28, 1988
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
ICU MEDICAL INC	5GM/100ML;298MG/100ML;300MG/100ML	N018876	008	Mar 28, 1988
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	5GM/100ML;298MG/100ML;450MG/100ML	N018362	007	Mar 28, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ ICU MEDICAL INC	5GM/100ML;298MG/100ML;900MG/100ML	N019691 008	Mar 24, 1988
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML;149MG/100ML;300MG/100ML	N018876 002	Jan 17, 1986
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML;224MG/100ML;225MG/100ML **	N018365 003	Jul 05, 1983
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML;224MG/100ML;300MG/100ML	N018876 003	Jan 17, 1986
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ ICU MEDICAL INC	5GM/100ML;224MG/100ML;900MG/100ML	N019691 007	Mar 24, 1988
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML;298MG/100ML;225MG/100ML **	N018365 004	Jul 05, 1983
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML;298MG/100ML;300MG/100ML	N018876 004	Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;225MG/100ML **	N018365 005	Mar 28, 1988
	5GM/100ML;149MG/100ML;225MG/100ML **	N018365 007	Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;300MG/100ML	N018876 005	Mar 28, 1988
	5GM/100ML;149MG/100ML;300MG/100ML	N018876 009	Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
+ ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;450MG/100ML	N018362 008	Mar 28, 1988
+ ICU MEDICAL INC	5GM/100ML;149MG/100ML;450MG/100ML	N018362 004	Mar 28, 1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ ICU MEDICAL INC	5GM/100ML;74.5MG/100ML;900MG/100ML	N019691 001	Mar 24, 1988
+ ICU MEDICAL INC	5GM/100ML;149MG/100ML;900MG/100ML	N019691 003	Mar 24, 1988

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;200MG/100ML	N018386 001	
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;450MG/100ML	N018229 001	
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;900MG/100ML	N018047 001	
BAXTER HLTHCARE	10GM/100ML;900MG/100ML	N016696 001	
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	2.5GM/100ML;450MG/100ML	N018030 001	
HOSPIRA	2.5GM/100ML;450MG/100ML	N018096 001	
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	2.5GM/100ML;900MG/100ML	N018376 001	
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ABBOTT	3.3GM/100ML;300MG/100ML	N018055 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML	N018030 005	
DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;200MG/100ML	N018030 004	
MILES	5GM/100ML;200MG/100ML	N018399 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;225MG/100ML	N019482 001	Oct 04, 1985
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;300MG/100ML	N019486 001	Oct 04, 1985
MILES	5GM/100ML;300MG/100ML	N018501 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;330MG/100ML	N018030 003	
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;450MG/100ML	N019484 001	Oct 04, 1985
B BRAUN	5GM/100ML;450MG/100ML	N018030 002	
MILES	5GM/100ML;450MG/100ML	N018400 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;900MG/100ML	N019483 001	Oct 04, 1985
B BRAUN	5GM/100ML;900MG/100ML	N018026 001	
MILES	5GM/100ML;900MG/100ML	N018500 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROTHYROXINE SODIUM

TABLET; ORAL

CHOLOXIN

ABBVIE	1MG	N012302 005
	2MG	N012302 002
	4MG	N012302 004
	6MG	N012302 006

DEZOCINE

INJECTABLE; INJECTION

DALGAN

ASTRAZENECA	5MG/ML	N019082 001	Dec 29, 1989
	10MG/ML	N019082 002	Dec 29, 1989
	15MG/ML	N019082 003	Dec 29, 1989

DIATRIZOATE MEGLUMINE

INJECTABLE; INJECTION

ANGIOVIST 282			
BAYER HLTHCARE	60%	A087726 001	Sep 23, 1982
CARDIOGRAFIN			
BRACCO	85%	N011620 002	
DIATRIZOATE MEGLUMINE			
BRACCO	76%	N010040 017	
HYPaque			
GE HEALTHCARE	30%	N016403 002	
	60%	N016403 001	
RENO-60			
BRACCO	60%	N010040 016	
RENO-DIP			
BRACCO	30%	N010040 012	
UROVIST MEGLUMINE DIU/CT			
BAYER HLTHCARE	30%	A087739 001	Sep 23, 1982
SOLUTION; URETERAL			
RENO-30			
BRACCO	30%	N010040 021	
UROVIST CYSTO			
BAYER HLTHCARE	30%	A087729 001	Sep 23, 1982
UROVIST CYSTO PEDIATRIC			
BAYER HLTHCARE	30%	A087731 001	Sep 23, 1982
SOLUTION; URETHRAL			
HYPaque-CYSTO			
GE HEALTHCARE	30%	N016403 003	

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

ANGIOVIST 292			
BAYER HLTHCARE	52%;8%	A087724 001	Sep 23, 1982
ANGIOVIST 370			
BAYER HLTHCARE	66%;10%	A087723 001	Sep 23, 1982
DIATRIZOATE-60			
INTL MEDICATION	52%;8%	A088166 001	Jun 17, 1983
HYPaque-76			
GE HEALTHCARE	66%;10%	A086505 001	
HYPaque-M, 75%			
GE HEALTHCARE	50%;25%	N010220 003	
HYPaque-M, 90%			
GE HEALTHCARE	60%;30%	N010220 002	
MD-60			
MALLINCKRODT	52%;8%	A087074 001	
MD-76			
MALLINCKRODT	66%;10%	A087073 001	
MD-76R			
+ LIEBEL-FLARSHEIM	66%;10%	N019292 001	Sep 29, 1989
RENOCAL-76			
BRACCO	66%;10%	A089347 001	Jun 01, 1988
RENOGRAFIN-60			
BRACCO	52%;8%	N010040 006	
RENOGRAFIN-76			
+ BRACCO	66%;10%	N010040 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

RENOVIST

BRACCO 34.3%;35% N010040 020

RENOVIST II

BRACCO 28.5%;29.1% N010040 019

SOLUTION; ORAL, RECTAL

GASTROVIST

BAYER HLTHCARE 66%;10% A087728 001 Sep 23, 1982

DIATRIZOATE MEGLUMINE; IODIPAMIDE MEGLUMINE

SOLUTION; INTRAUTERINE

SINOGRAFIN

+ BRACCO 52.7%;26.8% N011324 002

DIATRIZOATE SODIUM

FOR SOLUTION; ORAL, RECTAL

HYPAQUE

GE HEALTHCARE 100% N011386 001

INJECTABLE; INJECTION

HYPAQUE

GE HEALTHCARE 25% N009561 003

50% N009561 001

MD-50

MALLINCKRODT 50% A087075 001

UROVIST SODIUM 300

BAYER HLTHCARE 50% A087725 001 Sep 23, 1982

SOLUTION; ORAL, RECTAL

HYPAQUE

GE HEALTHCARE 40% N011386 003

SOLUTION; URETERAL

HYPAQUE SODIUM 20%

GE HEALTHCARE 20% N009561 002

DIAZEPAM

CAPSULE, EXTENDED RELEASE; ORAL

VALRELEASE

ROCHE 15MG N018179 001

GEL; RECTAL

DIASTAT

+ BAUSCH 5MG/ML (5MG/ML) ** N020648 002 Jul 29, 1997

+ 10MG/2ML (5MG/ML) ** N020648 003 Jul 29, 1997

+ 15MG/3ML (5MG/ML) ** N020648 004 Jul 29, 1997

+ 20MG/4ML (5MG/ML) ** N020648 005 Jul 29, 1997

INJECTABLE; INJECTION

DIAZEPAM

ABRAXIS PHARM 5MG/ML A070662 001 Jun 25, 1986

HIKMA 5MG/ML A070312 001 Dec 16, 1985

5MG/ML A071308 001 Jul 17, 1987

5MG/ML A071309 001 Jul 17, 1987

5MG/ML A071310 001 Jul 17, 1987

HOSPIRA 5MG/ML A071584 001 Oct 13, 1987

MARSAM PHARMS LLC 5MG/ML A072371 001 Jan 29, 1993

PARENTA PHARMS 5MG/ML A076815 001 Apr 15, 2004

US ARMY 5MG/ML ** N020124 001 Dec 05, 1990

WARNER CHILCOTT 5MG/ML A071613 001 Oct 22, 1987

5MG/ML A071614 001 Oct 22, 1987

WATSON LABS 5MG/ML A070296 001 Feb 12, 1986

5MG/ML A070911 001 Aug 28, 1986

5MG/ML A070912 001 Aug 28, 1986

5MG/ML A070930 001 Dec 01, 1986

WATSON LABS INC 5MG/ML A072370 001 Jan 29, 1993

5MG/ML A072397 001 Jan 29, 1993

DIZAC

PHARMACIA AND UPJOHN 5MG/ML ** N019287 001 Jun 18, 1993

VALIUM

+ ROCHE 5MG/ML ** N016087 001

TABLET; ORAL

DIAZEPAM

ACTAVIS ELIZABETH 2MG A070781 001 Mar 19, 1986

5MG A070706 001 Mar 19, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIAZEPAM

TABLET; ORAL

DIAZEPAM

	10MG	A070707 001	Mar 19, 1986
BARR	2MG	A070152 001	Nov 01, 1985
	10MG	A070154 001	Nov 01, 1985
CHARTWELL RX	2MG	A070302 001	Dec 20, 1985
	5MG	A070303 001	Dec 20, 1985
	10MG	A070304 001	Dec 20, 1985
DAVA PHARMS INC	2MG	A070228 002	Sep 26, 1985
	5MG	A070228 003	Sep 26, 1985
	10MG	A070228 001	Sep 26, 1985
DURAMED PHARMS BARR	2MG	A070894 001	Aug 27, 1986
	5MG	A070895 001	Aug 27, 1986
	10MG	A070896 001	Aug 27, 1986
FERNDALE LABS	2MG	A070903 001	Apr 01, 1987
	5MG	A070904 001	Apr 01, 1987
	10MG	A070905 001	Apr 01, 1987
HALSEY	2MG	A070987 001	Aug 15, 1986
	5MG	A070996 001	Aug 15, 1986
	10MG	A070956 001	Aug 15, 1986
IVAX SUB TEVA PHARMS	2MG	A070360 001	Sep 04, 1985
	5MG	A070361 001	Sep 04, 1985
	10MG	A070362 001	Sep 04, 1985
MARTEC USA LLC	10MG	A072402 001	Apr 25, 1989
PIONEER PHARMS	2MG	A070787 001	Aug 02, 1988
	5MG	A070788 001	Aug 02, 1988
	10MG	A070776 001	Aug 02, 1988
ROXANE	2MG	A070356 001	Jun 17, 1986
	5MG	A070357 001	Jun 17, 1986
	10MG	A070358 001	Jun 17, 1986
TEVA PHARMS	5MG	A070153 001	Nov 01, 1985
VIRTUS	2MG	A070462 001	Feb 25, 1986
	5MG	A070463 001	Feb 25, 1986
WARNER CHILCOTT	2MG	A070209 001	Sep 04, 1985
	5MG	A070210 001	Sep 04, 1985
	10MG	A070222 001	Sep 04, 1985
WATSON LABS	2MG	A070456 001	Nov 01, 1985
	5MG	A070457 001	Nov 01, 1985
	10MG	A070458 001	Nov 01, 1985
Q-PAM			
QUANTUM PHARMICS	2MG	A070423 001	Dec 12, 1985
	2MG	A072431 001	Apr 29, 1988
	5MG	A070424 001	Dec 12, 1985
	5MG	A072432 001	Apr 29, 1988
	10MG	A070425 001	Dec 12, 1985
	10MG	A072433 001	Apr 29, 1988

DIAZOXIDE

CAPSULE; ORAL

PROGLYCEM

TEVA BRANDED PHARM	50MG	N017425 001	
	100MG	N017425 002	

INJECTABLE; INJECTION

DIAZOXIDE

ABRAXIS PHARM	15MG/ML	A071519 001	Aug 26, 1987
HYPERSTAT			
SCHERING	15MG/ML	N016996 001	

DIBUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEAVY SOLUTION NUPERCAINE

NOVARTIS	2.5MG/ML	N006203 001	
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DICHLORPHENAMIDE

TABLET; ORAL

DARANIDE

+ XERIS	50MG **	N011366 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DICLOFENAC

CAPSULE; ORAL

ZORVOLEX

+ ZYLA	18MG **	N204592 001	Oct 18, 2013
+	35MG	N204592 002	Oct 18, 2013

DICLOFENAC POTASSIUM

TABLET; ORAL

CATAFLAM

+ NOVARTIS	25MG **	N020142 001	Nov 24, 1993
+	50MG **	N020142 002	Nov 24, 1993

DICLOFENAC POTASSIUM

CHARTWELL RX	50MG	A075582 001	Feb 23, 2001
SUN PHARM INDUSTRIES	50MG	A075470 001	Feb 21, 2002
WATSON LABS TEVA	50MG	A075152 001	Nov 27, 1998

DICLOFENAC SODIUM

GEL; TOPICAL

SOLARAZE

+ FOUGERA PHARMS	3% **	N021005 001	Oct 16, 2000
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SOLUTION; INTRAVENOUS

DICLOFENAC SODIUM

RISING	37.5MG/ML (37.5MG/ML)	A208786 001	Jun 18, 2019
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DYLOJECT

+ JAVELIN PHARMS INC	37.5MG/ML (37.5MG/ML) **	N022396 001	Dec 23, 2014
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SOLUTION; TOPICAL

DICLOFENAC SODIUM

APOTEX	1.5%	A202027 001	May 27, 2014
PAI HOLDINGS PHARM	1.5%	A202769 001	Jul 08, 2015
RISING	1.5%	A206715 001	Aug 07, 2017
TWI PHARMS	1.5%	A202393 001	Nov 24, 2014

PENNSAID

+ NUVO PHARMS INC	1.5% **	N020947 001	Nov 04, 2009
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SOLUTION/DROPS; OPHTHALMIC

DICLOFENAC SODIUM

AKORN	0.1%	A077845 001	Apr 17, 2008
FALCON PHARMS	0.1%	N020809 001	May 04, 1998

VOLTAREN

+ NOVARTIS	0.1%	N020037 001	Mar 28, 1991
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TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUM

AUROBINDO PHARMA USA	50MG	A075281 002	Feb 12, 2002
	75MG	A075281 003	Feb 12, 2002
CHARTWELL RX	25MG	A074376 001	Sep 28, 1995
	50MG	A074376 002	Sep 28, 1995
	75MG	A074394 001	Nov 30, 1995
MICRO LABS	50MG	A074986 001	Feb 26, 1999
	75MG	A074986 002	Feb 26, 1999
PLIVA	50MG	A074432 002	Jul 29, 1999
	75MG	A074432 003	Jul 29, 1999
ROXANE	25MG	A074391 001	Jun 29, 1995
	50MG	A074391 002	Jun 29, 1995
	75MG	A074391 003	Jun 29, 1995
TEVA	50MG	A074723 001	Mar 30, 1999
	75MG	A074390 001	Aug 15, 1996
TEVA PHARMS	25MG	A074459 001	Jun 25, 1997
	50MG	A074459 002	Jun 25, 1997
	75MG	A074459 003	Jun 25, 1997

VOLTAREN

+ NOVARTIS	25MG **	N019201 001	Jul 28, 1988
+	50MG **	N019201 002	Jul 28, 1988
+	75MG **	N019201 003	Jul 28, 1988

TABLET, EXTENDED RELEASE; ORAL

DICLOFENAC SODIUM

ACTAVIS ELIZABETH	100MG	A075910 001	Jan 07, 2002
AUROBINDO PHARMA USA	100MG	A076152 001	Dec 13, 2001

VOLTAREN-XR

+ NOVARTIS	100MG **	N020254 001	Mar 08, 1996
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM AND MISOPROSTOL

EXELA HOLDINGS	50MG;0.2MG	A200540 001	Mar 14, 2014
	75MG;0.2MG	A200540 002	Mar 14, 2014
ZYDUS PHARMS	50MG;0.2MG	A206771 001	Jun 12, 2023
	75MG;0.2MG	A206771 002	Jun 12, 2023

DICLOXACILLIN SODIUM

CAPSULE;ORAL

DICLOXACILLIN SODIUM

SANDOZ	EQ 125MG BASE	A061454 002	
	EQ 250MG BASE	A061454 001	
	EQ 500MG BASE	A061454 003	

DYCILL

GLAXOSMITHKLINE	EQ 250MG BASE	A060254 002	
	EQ 250MG BASE	A062238 001	
	EQ 500MG BASE	A060254 003	
	EQ 500MG BASE	A062238 002	

PATHOCIL

+ WYETH AYERST	EQ 250MG BASE **	N050011 002	
+	EQ 500MG BASE **	N050011 003	Mar 28, 1983

FOR SUSPENSION;ORAL

DICLOXACILLIN SODIUM

APOTHECON	EQ 62.5MG BASE/5ML	A061455 001	
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DYNAPEN

APOTHECON	EQ 62.5MG BASE/5ML	N050337 002	
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PATHOCIL

WYETH AYERST	EQ 62.5MG BASE/5ML	N050092 001	
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DICUMAROL

CAPSULE;ORAL

DICUMAROL

LILLY	25MG	N005509 003	
	50MG	N005509 001	

TABLET;ORAL

DICUMAROL

ABBVIE	25MG	N005545 003	
	50MG	N005545 004	
	100MG	N005545 005	

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

BENTYL

+ ALLERGAN	10MG **	N007409 003	Oct 15, 1984
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DICYCLOMINE HYDROCHLORIDE

PIONEER PHARMS	10MG	A089361 001	Jan 10, 1989
SUN PHARM INDUSTRIES	10MG	A084505 001	Oct 21, 1986
WATSON LABS	10MG	A083179 001	Feb 12, 1986

INJECTABLE;INJECTION

DICYCLOMINE HYDROCHLORIDE

DR REDDYS	10MG/ML	A080614 001	Feb 11, 1986
HIKMA	10MG/ML	A210788 001	Feb 11, 2019
PRAXGEN PHARMS	10MG/ML	A212058 001	Apr 26, 2019
RENEW PHARMS	10MG/ML	A207084 001	May 04, 2018

SYRUP;ORAL

BENTYL

+ APTALIS PHARMA US	10MG/5ML **	N007961 002	Oct 15, 1984
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DICYCLOMINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	A084479 001	
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TABLET;ORAL

BENTYL

+ ALLERGAN	20MG **	N007409 001	Oct 15, 1984
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DICYCLOMINE HYDROCHLORIDE

PIONEER PHARMS	20MG	A088585 001	Aug 20, 1986
SUN PHARM INDUSTRIES	20MG	A084600 001	Jul 29, 1985
WATSON LABS	20MG	A084361 001	Feb 06, 1986

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIDANOSINE

CAPSULE, DELAYED REL PELLETS;ORAL

DIDANOSINE

AUROBINDO PHARMA	125MG	A090094 001	Sep 24, 2008
	200MG	A090094 002	Sep 24, 2008
	250MG	A090094 003	Sep 24, 2008
	400MG	A090094 004	Sep 24, 2008
BARR	200MG	A077167 001	Dec 03, 2004
	250MG	A077167 002	Dec 03, 2004
	400MG	A077167 003	Dec 03, 2004
MYLAN PHARMS INC	125MG	A090788 001	Apr 08, 2010
	200MG	A090788 002	Apr 08, 2010
	250MG	A090788 003	Apr 08, 2010
	400MG	A090788 004	Apr 08, 2010

VIDEX EC

+ BRISTOL MYERS SQUIBB	125MG **	N021183 001	Oct 31, 2000
+	200MG **	N021183 002	Oct 31, 2000
+	250MG **	N021183 003	Oct 31, 2000
+	400MG **	N021183 004	Oct 31, 2000

FOR SOLUTION;ORAL

DIDANOSINE

AUROBINDO PHARMA	10MG/ML	A078112 001	Mar 08, 2007
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VIDEX

+ BRISTOL	10MG/ML	N020156 001	Oct 09, 1991
BRISTOL MYERS SQUIBB	100MG/PACKET	N020155 003	Oct 09, 1991
	167MG/PACKET	N020155 004	Oct 09, 1991
	250MG/PACKET	N020155 005	Oct 09, 1991
	375MG/PACKET	N020155 006	Oct 09, 1991

TABLET, CHEWABLE;ORAL

VIDEX

+ BRISTOL MYERS SQUIBB	25MG **	N020154 002	Oct 09, 1991
+	50MG **	N020154 003	Oct 09, 1991
+	100MG **	N020154 004	Oct 09, 1991
+	150MG **	N020154 005	Oct 09, 1991
+	200MG **	N020154 006	Oct 28, 1999

TABLET, FOR SUSPENSION;ORAL

DIDANOSINE

AUROBINDO	100MG	A077275 001	Aug 14, 2012
	150MG	A077275 002	Aug 14, 2012
	200MG	A077275 003	Aug 14, 2012

DIENESTROL

CREAM;VAGINAL

DIENESTROL

ORTHO MCNEIL PHARM	0.01%	N006110 005	
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DV

SANOFI AVENTIS US	0.01%	A083518 001	
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ESTRAGUARD

SOLVAY	0.01%	A084436 001	
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SUPPOSITORY;VAGINAL

DV

SANOFI AVENTIS US	0.7MG	A083517 001	
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DIETHYLCARBAMAZINE CITRATE

TABLET;ORAL

HETRAZAN

LEDERLE	50MG	N006459 001	
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DIETHYLPROPION HYDROCHLORIDE

TABLET;ORAL

DIETHYLPROPION HYDROCHLORIDE

CHARTWELL RX	25MG	A088267 001	Aug 25, 1983
	25MG	A088268 001	Aug 25, 1983
EPIC PHARMA LLC	25MG	A040828 001	Nov 05, 2008
SANDOZ	25MG	A085916 001	
TEVA	25MG	A088642 001	Sep 20, 1984
UCB INC	25MG	A085544 001	
WATSON LABS	25MG	A085741 001	

TENUATE

+ NOSTRUM LABS INC	25MG **	N011722 002	
SANOFI AVENTIS US	25MG	N017668 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL		
TEPANIL		
3M	25MG	N011673 001
TABLET, EXTENDED RELEASE; ORAL		
TENUATE		
SANOFI AVENTIS US	75MG	N017669 001
TENUATE DOSPAN		
+ NOSTRUM LABS INC	75MG **	N012546 001
TEPANIL TEN-TAB		
3M	75MG	N017956 001

DIETHYLSTILBESTROL

INJECTABLE; INJECTION		
STILBESTROL		
BRISTOL MYERS SQUIBB	0.2MG/ML	N004056 003
	0.5MG/ML	N004056 004
	1MG/ML	N004056 005
	5MG/ML	N004056 006
SUPPOSITORY; VAGINAL		
DIETHYLSTILBESTROL		
LILLY	0.1MG	N004040 001
	0.5MG	N004040 002
STILBESTROL		
BRISTOL MYERS SQUIBB	0.1MG	N004056 001
	0.5MG	N004056 002
TABLET; ORAL		
DIETHYLSTILBESTROL		
LILLY	0.1MG	N004041 002
	0.5MG	N004041 003
	1MG	N004041 004
	5MG	N004041 005
STILBESTROL		
TABLICAPS	0.5MG	A083004 001
	1MG	A083002 001
	5MG	A083006 001
STILBETIN		
BRISTOL MYERS SQUIBB	0.1MG	N004056 007
	0.25MG	N004056 017
	0.5MG	N004056 008
	1MG	N004056 009
	5MG	N004056 010
TABLET, DELAYED RELEASE; ORAL		
DIETHYLSTILBESTROL		
LILLY	0.1MG	N004039 002
	0.25MG	N004039 005
	0.5MG	N004039 003
	1MG	N004039 004
	5MG	N004039 006
STILBESTROL		
TABLICAPS	0.5MG	A083003 001
	1MG	A083005 001
	5MG	A083007 001
STILBETIN		
BRISTOL MYERS SQUIBB	0.1MG	N004056 011
	0.5MG	N004056 012
	1MG	N004056 013
	5MG	N004056 014

DIETHYLSTILBESTROL DIPHOSPHATE

INJECTABLE; INJECTION		
STILPHOSTROL		
BAYER PHARMS	250MG/5ML	N010010 001
TABLET; ORAL		
STILPHOSTROL		
BAYER PHARMS	50MG	N010010 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE				
AVONDALE PHARMS	0.05%		A075187 001	Mar 30, 1998
FLORONE				
PFIZER	0.05% **		N017741 001	
FLORONE E				
PFIZER	0.05%		N019259 001	Aug 28, 1985
PSORCON				
+ TARO PHARMS NORTH	0.05% **		N020205 001	Nov 20, 1992
OINTMENT; TOPICAL				
DIFLORASONE DIACETATE				
AKORN	0.05%		A206572 001	Jul 24, 2015
PSORCON				
+ PFIZER	0.05%		N019260 001	Aug 28, 1985
PSORCON E				
PFIZER	0.05%		N017994 001	

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL				
ANI PHARMS	500MG		A074604 001	Jun 10, 1996
DASTECH INTL	250MG		A073562 001	Nov 27, 1992
	500MG		A073563 001	Nov 27, 1992
PUREPAC PHARM	250MG		A074285 001	May 07, 1996
	500MG		A074285 002	May 07, 1996
TEVA	250MG		A073679 001	Jul 31, 1992
WATSON LABS	250MG		A074400 001	Jul 17, 1997
	500MG		A074400 002	Jul 17, 1997
DOLOBID				
+ MERCK	250MG **		N018445 001	Apr 19, 1982
+	500MG **		N018445 002	Apr 19, 1982

DIGITOXIN

INJECTABLE; INJECTION

CRYSTODIGIN				
LILLY	0.2MG/ML		A084100 005	

DIGOXIN

CAPSULE; ORAL

LANOXICAPS				
GLAXOSMITHKLINE LLC	0.05MG		N018118 002	Jul 26, 1982
	0.1MG		N018118 003	Jul 26, 1982
	0.15MG		N018118 004	Sep 24, 1984
	0.2MG		N018118 001	Jul 26, 1982

INJECTABLE; INJECTION

DIGOXIN				
ABRAXIS PHARM	0.25MG/ML		A083217 001	
HOSPIRA	0.25MG/ML		A040093 001	May 16, 1996
	0.25MG/ML		A040206 001	Aug 28, 1998
WYETH AYERST	0.25MG/ML		A084386 001	
DIGOXIN PEDIATRIC				
HOSPIRA	0.1MG/ML		A040092 001	Apr 25, 1996

TABLET; ORAL

LANOXIN				
+ CONCORDIA	0.1875MG		N020405 003	Sep 30, 1997
	0.375MG		N020405 005	Sep 30, 1997
	0.5MG		N020405 006	Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45				
+ BAUSCH	1MG/ML **		N005929 001	

DIHYDROERGOTAMINE MESYLATE; HEPARIN SODIUM; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

EMBOLEX				
NOVARTIS	0.5MG/0.5ML; 2,500		N018885 001	Nov 30, 1984
	UNITS/0.5ML; 5.33MG/0.5ML			
	0.5MG/0.7ML; 5,000		N018885 002	Nov 30, 1984
	UNITS/0.7ML; 7.46MG/0.7ML			

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

CARDIZEM SR

+	BIOVAIL	60MG **	N019471 001	Jan 23, 1989
+		90MG **	N019471 002	Jan 23, 1989
+		120MG **	N019471 003	Jan 23, 1989
+		180MG **	N019471 004	Jan 23, 1989

DILACOR XR

+	ALLERGAN	120MG **	N020092 001	May 29, 1992
+		180MG **	N020092 002	May 29, 1992
+		240MG **	N020092 003	May 29, 1992

DILT-CD

	APOTEX	120MG	A076151 001	May 20, 2004
		180MG	A076151 002	May 20, 2004
		240MG	A076151 003	May 20, 2004
		300MG	A076151 004	May 20, 2004

DILTIAZEM HYDROCHLORIDE

	ACTAVIS LABS FL INC	120MG	A074852 001	Oct 10, 1997
		180MG	A074852 002	Oct 10, 1997
		240MG	A074852 003	Oct 10, 1997
	BIOVAIL	60MG	A074845 001	Sep 15, 1999
		90MG	A074845 002	Sep 15, 1999
		120MG	A074845 003	Sep 15, 1999
		120MG	N020939 001	Jan 28, 2000
		180MG	N020939 002	Jan 28, 2000
		240MG	N020939 003	Jan 28, 2000
		300MG	N020939 004	Jan 28, 2000
		360MG	N020939 005	Sep 14, 2001
		420MG	N020939 006	Sep 14, 2001
	MYLAN	120MG	A075124 002	Mar 18, 1998
		180MG	A075124 003	Mar 18, 1998
		240MG	A075124 001	Mar 18, 1998
	NESHER PHARMS	120MG	A076563 002	Sep 12, 2006
		180MG	A076563 003	Sep 12, 2006
		240MG	A076563 004	Sep 12, 2006
		300MG	A076563 005	Sep 12, 2006
		360MG	A076563 006	Sep 12, 2006
		420MG	A076563 001	Sep 12, 2006
	PAR PHARM	360MG	A209766 001	May 30, 2018
	SUN PHARM	120MG	A090492 001	Oct 28, 2011
		180MG	A090492 002	Oct 28, 2011
		240MG	A090492 003	Oct 28, 2011
		300MG	A090492 004	Oct 28, 2011
	TEVA	60MG	A074079 001	Nov 30, 1993
		90MG	A074079 002	Nov 30, 1993
		120MG	A074079 003	Nov 30, 1993

DILTZAC

	APOTEX	120MG	A076395 001	Feb 01, 2006
		180MG	A076395 002	Feb 01, 2006
		240MG	A076395 003	Feb 01, 2006
		300MG	A076395 004	Feb 01, 2006
		360MG	A076395 005	Feb 01, 2006

INJECTABLE; INJECTION

CARDIZEM

	BIOVAIL	100MG/VIAL **	N020792 001	Sep 05, 1997
+	BIOVAIL LABS INTL	5MG/ML **	N020027 001	Oct 24, 1991
+		25MG/VIAL **	N020027 003	Aug 18, 1995

DILTIAZEM HYDROCHLORIDE

	DR REDDYS	5MG/ML	A074894 001	Aug 26, 1997
	HOSPIRA	5MG/ML	A075004 001	Feb 16, 2000
		5MG/ML	A075106 001	Apr 29, 1999
	INTL MEDICATION	5MG/ML	A075749 001	Nov 21, 2001
	MYLAN LABS LTD	5MG/ML	A075375 001	Sep 30, 1999

SOLUTION; INTRAVENOUS

DILTIAZEM HYDROCHLORIDE IN DEXTROSE 5%

+	EXELA PHARMA	125MG/125ML (1MG/ML)	N215252 001	Oct 28, 2021
+		250MG/250ML (1MG/ML)	N215252 002	Oct 28, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DILTIAZEM HYDROCHLORIDE

TABLET; ORAL

DILTIAZEM HYDROCHLORIDE

APOTHECON	30MG	A074051 001	Mar 31, 1993
	60MG	A074051 002	Mar 31, 1993
	90MG	A074051 003	Mar 31, 1993
	120MG	A074051 004	Mar 31, 1993
CHARTWELL MOLECULES	30MG	A074093 001	Nov 05, 1992
	60MG	A074093 002	Nov 05, 1992
	90MG	A074093 003	Nov 05, 1992
	120MG	A074093 004	Nov 05, 1992
EDENBRIDGE PHARMS	30MG	A211596 001	Nov 18, 2019
	60MG	A211596 002	Nov 18, 2019
	90MG	A211596 003	Nov 18, 2019
	120MG	A211596 004	Nov 18, 2019
IVAX SUB TEVA PHARMS	30MG	A074168 001	Mar 03, 1995
	60MG	A074168 002	Mar 03, 1995
	90MG	A074168 003	Mar 03, 1995
	120MG	A074168 004	Mar 03, 1995
RISING	30MG	A072838 004	Nov 05, 1992
	60MG	A072838 003	Nov 05, 1992
	90MG	A072838 002	Nov 05, 1992
	120MG	A072838 001	Nov 05, 1992
TEVA	30MG	A074084 001	Feb 25, 1994
	60MG	A074084 002	Feb 25, 1994
TEVA PHARMS	30MG	A074067 001	Nov 05, 1992
	60MG	A074067 002	Nov 05, 1992
	90MG	A074067 003	Nov 05, 1992
	120MG	A074067 004	Nov 05, 1992

DILTIAZEM MALATE

TABLET, EXTENDED RELEASE; ORAL

TIAMATE

MERCK	EQ 120MG HYDROCHLORIDE	N020506 001	Oct 04, 1996
	EQ 180MG HYDROCHLORIDE	N020506 002	Oct 04, 1996
	EQ 240MG HYDROCHLORIDE	N020506 003	Oct 04, 1996

DILTIAZEM MALATE; ENALAPRIL MALEATE

TABLET, EXTENDED RELEASE; ORAL

TECZEM

BIOVAIL	EQ 180MG HYDROCHLORIDE; 5MG	N020507 001	Oct 04, 1996
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DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

BAXTER HLTHCARE	50MG/ML	A084767 001	
WATSON LABS	50MG/ML	A083531 001	
+ WATSON LABS TEVA	50MG/ML	A080615 001	
WYETH AYERST	50MG/ML	A084316 001	

LIQUID; ORAL

DIMENHYDRINATE

ALRA	12.5MG/4ML	A080715 001	
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TABLET; ORAL

DIMENHYDRINATE

HEATHER	50MG	A080841 001	
NEXGEN PHARMA INC	50MG	A085985 001	
+ WATSON LABS	50MG	A085166 001	

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL

DIMETHYL FUMARATE

SAWAI USA	120MG	A210285 001	Dec 21, 2021
	240MG	A210285 002	Dec 21, 2021
ZYDUS PHARMS	120MG	A210538 001	Sep 24, 2020
	240MG	A210538 002	Sep 24, 2020

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

DIMETHYL SULFOXIDE

MYLAN INSTITUTIONAL 50%

A076185 001 Nov 29, 2002

DIMYRISTOYL LECITHIN; PERFLEXANE

INJECTABLE; INTRAVENOUS

IMAGENT

VESSELON SPV LLC 0.92MG/VIAL;0.092MG/VIAL

N021191 001 May 31, 2002

DINOPROST TROMETHAMINE

INJECTABLE; INJECTION

PROSTIN F2 ALPHA

PHARMACIA AND UPJOHN EQ 5MG BASE/ML

N017434 001

DINOPROSTONE

SUPPOSITORY; VAGINAL

PROSTIN E2

+ PFIZER 20MG

N017810 001

DIPHEMANIL METHYLSULFATE

TABLET; ORAL

PRANTAL

SCHERING 100MG

N008114 004

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

BENADRYL

MCNEIL CONS 25MG

N005845 007

50MG

N005845 001

DIPHENHYDRAMINE HYDROCHLORIDE

ALRA 25MG

A080519 004

50MG

A080519 003

ANABOLIC 50MG

A083275 001

ELKINS SINN 25MG

A085701 001

50MG

A085701 002

FOSUN PHARMA 25MG

A080832 001

25MG

A080845 002

50MG

A080832 002

50MG

A080845 001

HALSEY 50MG

A087914 001

Jun 04, 1984

HEATHER 25MG

A084524 001

50MG

A083953 001

HERITAGE PHARMA 50MG

A080727 001

50MG

A080738 001

HIKMA INTL PHARMS 50MG

A083567 001

IMPAX LABS 25MG

A080807 001

50MG

A080807 002

IVAX SUB TEVA PHARMS 25MG

A080762 001

50MG

A080762 002

LANNETT 25MG

A080868 001

50MG

A080868 002

LEDERLE 25MG

A086874 001

50MG

A086875 001

LNK 25MG

A087977 001

Jan 27, 1983

50MG

A087978 001

Jan 27, 1983

MK LABS 25MG

A083087 001

50MG

A083087 002

MUTUAL PHARM 25MG

A084506 001

NEWTRON PHARMS 25MG

A086543 001

50MG

A086544 001

NEXGEN PHARMA INC 25MG

A083634 001

PERRIGO 25MG

A083061 001

50MG

A083061 002

PIONEER PHARMS 25MG

A089101 001

Dec 20, 1985

50MG

A088880 001

Dec 20, 1985

PUREPAC PHARM 25MG

A085156 001

50MG

A085150 001

PVT FORM 25MG

A083027 001

50MG

A083027 002

ROXANE 50MG

A080635 001

SUN PHARM INDUSTRIES 25MG

A089488 001

Jan 02, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

	50MG	A089489	001	Jan 02, 1987
SUPERPHARM	25MG	A089040	001	May 15, 1985
	50MG	A089041	001	May 15, 1985
TEVA	25MG	A085874	001	
	50MG	A085874	002	
VALEANT PHARM INTL	25MG	A080596	001	
	50MG	A080592	001	
VANGARD	25MG	A088034	001	Oct 27, 1982
	50MG	A087630	001	
WATSON LABS	25MG	A080728	001	
	25MG	A083797	001	
	25MG	A085138	001	
	50MG	A083797	002	
	50MG	A085083	001	
WHITEWORTH TOWN PLSN	25MG	A083441	001	
	50MG	A080800	001	

ELIXIR; ORAL

BELIX

HALSEY	12.5MG/5ML	A086586	001	Oct 03, 1983
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BENADRYL

MCNEIL CONS	12.5MG/5ML	N005845	004	
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DIBENIL

CENCI	12.5MG/5ML	A088304	001	Dec 16, 1983
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DIPHEN

USL PHARMA	12.5MG/5ML	A084640	001	
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DIPHENHYDRAMINE HYDROCHLORIDE

BUNDY	12.5MG/5ML	A083674	001	
CENCI	12.5MG/5ML	A087941	001	Dec 17, 1982
KV PHARM	12.5MG/5ML	A085621	001	
LANNETT	12.5MG/5ML	A080939	002	
LEDERLE	12.5MG/5ML	A086937	001	
MK LABS	12.5MG/5ML	A083088	002	
NASKA	12.5MG/5ML	A088680	001	May 31, 1985
PERRIGO	12.5MG/5ML	A083063	001	
PUREPAC PHARM	12.5MG/5ML	A083237	001	Jan 25, 1982
PVT FORM	12.5MG/5ML	A085287	001	
ROXANE	12.5MG/5ML	A080643	001	

HYDRAMINE

ALPHARMA US PHARMS	12.5MG/5ML	A080763	002	
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INJECTABLE; INJECTION

BENADRYL

MCNEIL CONS	10MG/ML	N006146	001	
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+

	50MG/ML **	N006146	002	
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BENADRYL PRESERVATIVE FREE

+

MCNEIL CONS	50MG/ML **	N009486	001	
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DIPHENHYDRAMINE HYDROCHLORIDE

BEL MAR	10MG/ML	A080822	001	
DR REDDYS	10MG/ML	A080873	001	
	50MG/ML	A080873	002	
EUROHLTH INTL SARL	50MG/ML	A083183	001	
HOSPIRA	50MG/ML	A040140	001	Nov 20, 1998
LYPHOMED	10MG/ML	A087066	001	
WATSON LABS	10MG/ML	A083533	001	
WYETH AYERST	50MG/ML	A080577	001	

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

ABRAXIS PHARM	50MG/ML	A080586	002	
DR REDDYS	50MG/ML	A080873	003	
INTL MEDICATION	50MG/ML	A084094	001	

SYRUP; ORAL

ANTITUSSIVE

PERRIGO	12.5MG/5ML	A071292	001	Apr 10, 1987
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BELDIN

HALSEY	12.5MG/5ML	A089179	001	Jun 05, 1986
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BENYLIN

PARKE DAVIS	12.5MG/5ML	N006514	004	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

SYRUP;ORAL

DIPHEN

MORTON GROVE 12.5MG/5ML

A070118 001 Oct 01, 1985

DIPHENHYDRAMINE HYDROCHLORIDE

AKORN 12.5MG/5ML

A072416 001 Sep 28, 1990

ALPHARMA US PHARMS 12.5MG/5ML

A070497 001 Apr 25, 1989

CUMBERLAND SWAN 12.5MG/5ML

A073611 001 Aug 20, 1992

HYDRAMINE

ALPHARMA US PHARMS 12.5MG/5ML

A070205 001 Jan 28, 1986

SILPHEN

LANNETT CO INC 12.5MG/5ML

A072646 001 Feb 27, 1992

VICKS FORMULA 44

WARNER CHILCOTT 12.5MG/5ML

A070524 001 Jan 14, 1987

DIPHENHYDRAMINE HYDROCHLORIDE; NAPROXEN SODIUM

TABLET;ORAL

NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE

P AND L 25MG;220MG

A207597 001 Jan 25, 2019

DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

BENYLIN

PARKE DAVIS 12.5MG/5ML;30MG/5ML

N019014 001 Jun 11, 1985

DIPHENIDOL HYDROCHLORIDE

TABLET;ORAL

VONTROL

GLAXOSMITHKLINE EQ 25MG BASE

N016033 001

DIPHENYLPYRALINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

HISPRIL

GLAXOSMITHKLINE 5MG

N011945 001

DIPIVEFRIN HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AKPRO

EPIC PHARMA LLC 0.1%

A074382 001 Sep 29, 1995

DIPIVEFRIN HYDROCHLORIDE

BAUSCH AND LOMB 0.1%

A074188 001 May 19, 1995

FALCON PHARMS 0.1%

A073636 001 Jun 30, 1994

PROPINE

ALLERGAN 0.1%

N018239 001

DIPYRIDAMOLE

INJECTABLE;INJECTION

DIPYRIDAMOLE

DR REDDYS 5MG/ML

A074952 001 Nov 26, 1997

EUGIA PHARMA SPECLTS 5MG/ML

A075769 001 Nov 27, 2002

FRESENIUS KABI USA 5MG/ML

A074956 001 Sep 30, 1998

HOSPIRA 5MG/ML

A074601 001 Dec 19, 1997

IV PERSANTINE

+ BOEHRINGER INGELHEIM 5MG/ML **

N019817 001 Dec 13, 1990

TABLET;ORAL

DIPYRIDAMOLE

ANDA REPOSITORY 25MG

A040898 001 Apr 23, 2008

50MG

A040898 002 Apr 23, 2008

75MG

A040898 003 Apr 23, 2008

ANI PHARMS 25MG

A086944 002 Apr 16, 1991

50MG

A086944 001 Feb 25, 1992

75MG

A086944 003 Feb 25, 1992

GLENMARK GENERICS 25MG

A088999 001 Feb 05, 1991

50MG

A089000 001 Feb 05, 1991

75MG

A089001 001 Feb 05, 1991

PUREPAC PHARM 25MG

A089425 001 Jul 12, 1990

50MG

A089426 001 Jul 12, 1990

75MG

A089427 001 Jul 12, 1990

WATSON LABS 50MG

A087160 001 Jun 07, 1996

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIRITHROMYCINTABLET, DELAYED RELEASE;ORAL
DYNABAC

LILLY RES LABS 250MG N050678 001 Jun 19, 1995

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATE

INTERPHARM	EQ 100MG BASE	A071190 001	Jan 15, 1987
	EQ 150MG BASE	A071191 001	Jan 15, 1987
IVAX SUB TEVA PHARMS	EQ 100MG BASE	A070186 001	Nov 18, 1985
	EQ 150MG BASE	A070187 001	Nov 18, 1985
MYLAN	EQ 100MG BASE	A070138 001	Jun 14, 1985
	EQ 150MG BASE	A070139 001	Jun 14, 1985
RISING	EQ 100MG BASE	A070470 001	Dec 10, 1985
	EQ 150MG BASE	A070471 001	Dec 10, 1985
SUN PHARM INDUSTRIES	EQ 100MG BASE	A070351 001	Dec 17, 1985
	EQ 150MG BASE	A070352 001	Dec 17, 1985
SUPERPHARM	EQ 100MG BASE	A070940 001	Feb 09, 1987
	EQ 150MG BASE	A070941 001	Feb 09, 1987
WATSON LABS	EQ 100MG BASE	A070240 001	Feb 02, 1986
	EQ 150MG BASE	A070241 001	Feb 02, 1986

CAPSULE, EXTENDED RELEASE;ORAL

DISOPYRAMIDE PHOSPHATE

NESHER PHARMS EQ 150MG BASE A071200 001 Dec 15, 1987

DISULFIRAM

TABLET;ORAL

ANTABUSE

+	ODYSSEY PHARMS	250MG	A088482 001	Dec 08, 1983
+		500MG	A088483 001	Dec 08, 1983
+	TEVA WOMENS	250MG **	N007883 003	
+		500MG **	N007883 002	

DISULFIRAM

DASH PHARMS	250MG	A203916 001	Mar 04, 2015
	500MG	A203916 002	Mar 04, 2015
HIKMA	250MG	A202652 001	Feb 05, 2014
	500MG	A202652 002	Feb 05, 2014
STRIDES PHARMA	250MG	A088792 001	Aug 14, 1984
	500MG	A088793 001	Aug 14, 1984
WATSON LABS	250MG	A086889 001	
	250MG	A087973 001	Aug 05, 1983
	500MG	A087974 001	Aug 05, 1983
WATSON LABS TEVA	500MG	A086890 001	

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL

DIVALPROEX SODIUM

RISING	EQ 125MG VALPROIC ACID	A090407 001	Mar 28, 2011
TEVA PHARMS USA	EQ 125MG VALPROIC ACID	A211505 001	Nov 17, 2020

TABLET, DELAYED RELEASE;ORAL

DEPAKOTE CP

ABBOTT	EQ 250MG BASE	N019794 001	Jul 11, 1990
	EQ 500MG BASE	N019794 002	Jul 11, 1990

DIVALPROEX SODIUM

ACTAVIS LABS FL INC	EQ 500MG VALPROIC ACID	A079080 001	Feb 25, 2011
ANCHEN PHARMS	EQ 500MG VALPROIC ACID	A078411 001	Nov 03, 2008
MYLAN	EQ 125MG VALPROIC ACID	A077254 001	Jul 29, 2008
	EQ 125MG VALPROIC ACID	A090062 001	Mar 17, 2009
	EQ 250MG VALPROIC ACID	A077254 002	Jul 29, 2008
	EQ 250MG VALPROIC ACID	A090062 002	Mar 17, 2009
	EQ 500MG VALPROIC ACID	A077254 003	Jul 29, 2008
	EQ 500MG VALPROIC ACID	A090062 003	Mar 17, 2009
TEVA	EQ 125MG VALPROIC ACID	A076941 001	Jul 29, 2008
	EQ 250MG VALPROIC ACID	A076941 002	Jul 29, 2008
	EQ 500MG VALPROIC ACID	A076941 003	Jul 29, 2008

TABLET, EXTENDED RELEASE;ORAL

DIVALPROEX SODIUM

AMTA	EQ 250MG VALPROIC ACID	A214462 001	Mar 15, 2021
	EQ 500MG VALPROIC ACID	A214462 002	Mar 15, 2021
ANCHEN PHARMS	EQ 250MG VALPROIC ACID	A078445 001	Feb 26, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIVALPROEX SODIUMTABLET, EXTENDED RELEASE;ORAL
DIVALPROEX SODIUM

	EQ 500MG VALPROIC ACID	A078445	002	Aug 04, 2009
COSETTE	EQ 500MG VALPROIC ACID	A078700	001	Aug 03, 2009
IMPAX LABS	EQ 250MG VALPROIC ACID	A078791	001	May 06, 2009
	EQ 500MG VALPROIC ACID	A078791	002	Aug 04, 2009

DOBUTAMINE HYDROCHLORIDEINJECTABLE; INJECTION
DOBUTAMINE HYDROCHLORIDE

BAXTER HLTHCARE	EQ 12.5MG BASE/ML	A074381	001	Sep 26, 1996
DR REDDYS	EQ 12.5MG BASE/ML	A074995	001	Mar 31, 1998
HOSPIRA	EQ 12.5MG BASE/ML	A074292	001	Feb 16, 1995
	EQ 1.25GM BASE/100ML	A074634	001	Sep 27, 1996
LUITPOLD	EQ 12.5MG BASE/ML	A074545	001	Jun 25, 1998
TELIGENT	EQ 12.5MG BASE/ML	A074098	001	Feb 21, 1995
TEVA PARENTERAL	EQ 12.5MG BASE/ML	A074206	001	Oct 19, 1993
WATSON LABS	EQ 12.5MG BASE/ML	A074114	001	Nov 30, 1993
WATSON LABS INC	EQ 12.5MG BASE/ML	A074279	001	Feb 18, 1998
DOBUTAMINE HYDROCHLORIDE IN	DEXTROSE 5%			
HOSPIRA	EQ 50MG BASE/100ML	N020269	001	Oct 19, 1993
	EQ 100MG BASE/100ML	N020269	002	Oct 19, 1993
	EQ 200MG BASE/100ML	N020269	003	Oct 19, 1993
DOBUTREX				
+ LILLY	EQ 12.5MG BASE/ML **	N017820	002	

DOCETAXEL

INJECTABLE; INJECTION

DOCEFREZ				
+ SUN PHARM	20MG/VIAL	N022534	001	May 03, 2011
+	80MG/VIAL	N022534	002	May 03, 2011
DOCETAXEL				
+ ACCORD HLTHCARE	20MG/0.5ML (40MG/ML)	N201195	001	Jun 08, 2011
+	80MG/2ML (40MG/ML)	N201195	002	Jun 08, 2011
APOTEX INC	20MG/0.5ML (40MG/ML)	N022312	001	Jan 11, 2012
	80MG/2ML (40MG/ML)	N022312	002	Jan 11, 2012
DFB ONCOLOGY LTD	20MG/ML (20MG/ML)	A206177	001	Jan 20, 2017
	80MG/4ML (20MG/ML)	A206177	002	Jan 20, 2017
	200MG/10ML (20MG/ML)	A206177	003	Jan 20, 2017
HENGRUI PHARMA	40MG/ML	A203170	001	Feb 15, 2017
+ HOSPIRA INC	120MG/6ML (20MG/ML)	N022234	006	Jun 23, 2016
MYLAN LABS LTD	20MG/ML (20MG/ML)	A203892	001	May 11, 2023
	80MG/4ML (20MG/ML)	A203892	002	May 11, 2023
NOVAST LABS	20MG/2ML (10MG/ML)	A207563	001	Aug 31, 2017
PFIZER LABS	20MG/2ML (10MG/ML)	N202356	001	Mar 13, 2014
	80MG/8ML (10MG/ML)	N202356	002	Mar 13, 2014
	130MG/13ML (10MG/ML)	N202356	003	Mar 13, 2014
	200MG/20ML (10MG/ML)	N202356	004	Mar 13, 2014
TEVA PHARMS USA	20MG/ML (20MG/ML)	A203877	001	Sep 16, 2015
	80MG/4ML (20MG/ML)	A203877	002	Sep 16, 2015
TAXOTERE				
+ SANOFI AVENTIS US	40MG/ML **	N020449	001	May 14, 1996
SOLUTION; INTRAVENOUS				
DOCETAXEL				
+ INGENUS PHARMS LLC	20MG/2ML (10MG/ML)	N215813	001	Nov 22, 2022
+	80MG/8ML (10MG/ML)	N215813	002	Nov 22, 2022
+	160MG/16ML (10MG/ML)	N215813	003	Nov 22, 2022

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE				
NOVAST LABS	0.125MG	A212410	001	Dec 27, 2019
	0.25MG	A212410	002	Dec 27, 2019
	0.5MG	A212410	003	Dec 27, 2019
PRINSTON INC	0.125MG	A211223	001	Dec 17, 2019
	0.25MG	A211223	002	Dec 17, 2019
	0.5MG	A211223	003	Dec 17, 2019
RK PHARMA	0.125MG	A215323	001	Apr 14, 2022
	0.25MG	A215323	002	Apr 14, 2022
	0.5MG	A215323	003	Apr 14, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

TEVA PHARMS USA	0.125MG	A210018 001	Apr 15, 2022
	0.25MG	A210018 002	Apr 15, 2022
	0.5MG	A210018 003	Apr 15, 2022

DOLASETRON MESYLATE

INJECTABLE; INJECTION

ANZEMET

+ VALIDUS PHARMS	12.5MG/0.625ML (20MG/ML)	N020624 002	Sep 11, 1997
+	100MG/5ML (20MG/ML)	N020624 001	Sep 11, 1997
	500MG/25ML (20MG/ML)	N020624 003	Dec 11, 2001

TABLET; ORAL

ANZEMET

+ VALIDUS PHARMS	100MG	N020623 002	Sep 11, 1997
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DONEPEZIL HYDROCHLORIDE

SOLUTION; ORAL

ARICEPT

+ EISAI INC	5MG/5ML	N021719 001	Oct 18, 2004
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TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

ACCORD HLTHCARE	5MG	A201335 001	Aug 29, 2011
	10MG	A201335 002	Aug 29, 2011
ACTAVIS ELIZABETH	23MG	A202415 001	Dec 17, 2015
APOTEX	5MG	A078841 001	Jun 02, 2011
	10MG	A078841 002	Jun 02, 2011
CHARTWELL RX	23MG	A203419 001	Apr 12, 2016
HERITAGE PHARMA	5MG	A077344 001	May 31, 2011
	10MG	A077344 002	May 31, 2011
HIKMA PHARMS	5MG	A090247 001	May 31, 2011
	10MG	A090247 002	May 31, 2011
HISUN PHARM HANGZHOU	23MG	A202410 001	Mar 24, 2017
MYLAN PHARMS INC	5MG	A090521 001	May 31, 2011
	10MG	A090521 002	May 31, 2011
OSMOTICA PHARM US	23MG	A203114 001	Jan 26, 2016
PAR PHARM	23MG	A202542 001	Jul 24, 2013
RISING	23MG	A202656 001	Oct 22, 2015
SANDOZ	5MG	A090290 001	May 31, 2011
	10MG	A090290 002	May 31, 2011
SUN PHARM	23MG	A204293 001	Jun 05, 2015
SUN PHARM INDS LTD	5MG	A076786 001	Nov 26, 2010
	10MG	A076786 002	Nov 26, 2010
UNICHEM	5MG	A203656 001	Jun 23, 2016
	10MG	A203656 002	Jun 23, 2016
WOCKHARDT	5MG	A091267 001	May 31, 2011
	10MG	A091267 002	May 31, 2011

TABLET, ORALLY DISINTEGRATING; ORAL

ARICEPT ODT

+ EISAI INC	5MG	N021720 001	Oct 18, 2004
+	10MG	N021720 002	Oct 18, 2004

DONEPEZIL HYDROCHLORIDE

CHARTWELL RX	5MG	A078388 002	Nov 26, 2010
	10MG	A078388 001	Nov 26, 2010
SUN PHARM INDUSTRIES	5MG	A077975 002	Dec 11, 2009
	10MG	A077975 001	Dec 11, 2009
UNICHEM	5MG	A204831 001	Nov 10, 2016
	10MG	A204831 002	Nov 10, 2016
ZYDUS PHARMS USA INC	5MG	A090175 001	May 10, 2011
	10MG	A090175 002	May 10, 2011

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE

AMNEAL PHARMS	10MG; 14MG	A208328 001	Jan 27, 2017
	10MG; 28MG	A208328 002	Jan 27, 2017

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

ABBOTT	40MG/ML	A070656	001	Jan 24, 1989
	80MG/ML	A070657	001	Jan 24, 1989
ABRAXIS PHARM	40MG/ML	A070012	001	Jun 12, 1985
	40MG/ML	A070058	001	Mar 20, 1985
	80MG/ML	A070013	001	Jun 12, 1985
	80MG/ML	A070059	001	Mar 20, 1985
	160MG/ML	A070364	001	Dec 04, 1985
AM REGENT	40MG/ML	A070799	001	Feb 11, 1987
	80MG/ML	A070820	001	Feb 11, 1987
	160MG/ML	A070826	001	Feb 11, 1987
BAXTER HLTHCARE	40MG/ML	N018398	001	
	80MG/ML	N018398	002	Mar 22, 1982
HOSPIRA	40MG/ML	A074403	001	May 23, 1996
IGI LABS INC	40MG/ML	A070087	001	Oct 23, 1985
	80MG/ML	A070089	001	Oct 23, 1985
	80MG/ML	A070090	001	Oct 23, 1985
	80MG/ML	A070091	001	Oct 23, 1985
	160MG/ML	A070092	001	Oct 23, 1985
	160MG/ML	A070093	001	Oct 23, 1985
	160MG/ML	A070094	001	Oct 23, 1985
INTL MEDICATION	40MG/ML	N018014	001	
LYPHOMED	40MG/ML	N018549	001	Mar 11, 1983
SMITH AND NEPHEW	40MG/ML	A070011	001	Aug 29, 1985
	40MG/ML	A070046	001	Aug 29, 1985
	80MG/ML	A070047	001	Aug 29, 1985
TELIGENT	40MG/ML	N018656	001	Jun 28, 1983
TEVA PARENTERAL	40MG/ML	A072999	001	Oct 23, 1991
	80MG/ML	A073000	001	Oct 23, 1991
WARNER CHILCOTT	40MG/ML	A070558	001	Sep 20, 1985
	40MG/ML	N018138	001	
	80MG/ML	A070559	001	Sep 20, 1985
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%				
+ B BRAUN	80MG/100ML	N019099	002	Oct 15, 1986
+	320MG/100ML	N019099	004	Oct 15, 1986
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER				
+ B BRAUN	40MG/100ML	N019099	001	Oct 15, 1986
+	160MG/100ML	N019099	003	Oct 15, 1986
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5%				
HOSPIRA	1.6MG/ML	N020542	001	Aug 30, 1995
INTROPIN				
HOSPIRA	40MG/ML	N017395	001	
	80MG/ML	N017395	002	
	160MG/ML	N017395	003	

DORIPENEM

INJECTABLE; INTRAVENOUS

DORIBAX

+ SHIONOGI INC	250MG/VIAL **	N022106	002	Oct 05, 2010
+	500MG/VIAL **	N022106	001	Oct 12, 2007

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

AM REGENT	EQ 2% BASE	A079186	001	Nov 18, 2009
TEVA PHARMS	EQ 2% BASE	A078756	001	Dec 04, 2008
ZAMBON SPA	EQ 2% BASE	A091034	001	Dec 04, 2013
TRUSOPT				
+ MSD SUB MERCK	EQ 2% BASE **	N020408	001	Dec 09, 1994

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

CHARTWELL MOLECULAR	EQ 2% BASE;EQ 0.5% BASE	A201998	001	Dec 17, 2014
RUBICON	EQ 2% BASE;EQ 0.5% BASE	A078201	001	Oct 28, 2008
TEVA PHARMS	EQ 2% BASE;EQ 0.5% BASE	A078704	001	Sep 28, 2009
ZAMBON SPA	EQ 2% BASE;EQ 0.5% BASE	A091180	001	Dec 04, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXACURIUM CHLORIDEINJECTABLE; INJECTION
NUROMAX

ABBVIE	EQ 1MG BASE/ML	N019946 001	Mar 07, 1991
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DOXAPRAM HYDROCHLORIDEINJECTABLE; INJECTION
DOXAPRAM HYDROCHLORIDE
WATSON LABS

20MG/ML	A073529 001	Jan 30, 1992
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DOXAZOSIN MESYLATETABLET; ORAL
DOXAZOSIN MESYLATE

ACTAVIS ELIZABETH	EQ 1MG BASE	A075574 001	Oct 18, 2000
	EQ 2MG BASE	A075574 002	Oct 18, 2000
	EQ 4MG BASE	A075574 003	Oct 18, 2000
	EQ 8MG BASE	A075574 004	Oct 18, 2000
ANI PHARMS	EQ 1MG BASE	A075432 001	Oct 18, 2000
	EQ 2MG BASE	A075432 002	Oct 18, 2000
	EQ 4MG BASE	A075432 003	Oct 18, 2000
	EQ 8MG BASE	A075432 004	Oct 18, 2000
AUROBINDO PHARMA USA	EQ 1MG BASE	A075509 001	Oct 19, 2000
	EQ 2MG BASE	A075509 002	Oct 19, 2000
	EQ 4MG BASE	A075509 003	Oct 19, 2000
	EQ 8MG BASE	A075509 004	Oct 19, 2000
CHARTWELL RX	EQ 1MG BASE	A075646 001	Oct 18, 2000
	EQ 2MG BASE	A075646 002	Oct 18, 2000
	EQ 4MG BASE	A075646 003	Oct 18, 2000
	EQ 8MG BASE	A075646 004	Oct 18, 2000
GENPHARM	EQ 1MG BASE	A075466 001	Oct 18, 2000
	EQ 2MG BASE	A075466 002	Oct 18, 2000
	EQ 4MG BASE	A075466 003	Oct 18, 2000
	EQ 8MG BASE	A075466 004	Oct 18, 2000
IVAX SUB TEVA PHARMS	EQ 1MG BASE	A075453 001	Oct 18, 2000
	EQ 2MG BASE	A075453 002	Oct 18, 2000
	EQ 4MG BASE	A075453 003	Oct 18, 2000
	EQ 8MG BASE	A075453 004	Oct 18, 2000
NESHER PHARMS	EQ 1MG BASE	A075609 001	Oct 18, 2000
	EQ 2MG BASE	A075609 002	Oct 18, 2000
	EQ 4MG BASE	A075609 003	Oct 18, 2000
	EQ 8MG BASE	A075609 004	Oct 18, 2000
PLIVA	EQ 1MG BASE	A075750 001	Jun 08, 2001
	EQ 2MG BASE	A075750 002	Jun 08, 2001
	EQ 4MG BASE	A075750 003	Jun 08, 2001
	EQ 8MG BASE	A075750 004	Jun 08, 2001
STRIDES PHARMA	EQ 1MG BASE	A076161 001	Jun 10, 2004
	EQ 2MG BASE	A076161 002	Jun 10, 2004
	EQ 4MG BASE	A076161 003	Jun 10, 2004
	EQ 8MG BASE	A076161 004	Jun 10, 2004
TEVA	EQ 1MG BASE	A075353 001	Jan 12, 2001
	EQ 2MG BASE	A075353 002	Jan 12, 2001
	EQ 4MG BASE	A075353 003	Jan 12, 2001
	EQ 8MG BASE	A075353 004	Jan 12, 2001
WATSON LABS INC	EQ 1MG BASE	A075426 001	Oct 18, 2000
	EQ 2MG BASE	A075426 002	Oct 18, 2000
	EQ 4MG BASE	A075426 003	Oct 18, 2000
	EQ 8MG BASE	A075426 004	Oct 18, 2000

DOXEPIN HYDROCHLORIDECAPSULE; ORAL
DOXEPIN HYDROCHLORIDE

DAVA PHARMS INC	EQ 10MG BASE	A071685 001	Jan 05, 1988
	EQ 25MG BASE	A071686 001	Jan 05, 1988
	EQ 50MG BASE	A071673 001	Jan 05, 1988
	EQ 75MG BASE	A071674 001	Jan 05, 1988
	EQ 100MG BASE	A071675 001	Jan 05, 1988
	EQ 150MG BASE	A071676 001	Jan 05, 1988
LANNETT CO INC	EQ 10MG BASE	A212997 001	Jul 24, 2020
	EQ 25MG BASE	A212997 002	Jul 24, 2020
	EQ 50MG BASE	A212997 003	Jul 24, 2020
	EQ 75MG BASE	A212997 004	Jul 24, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXEPIN HYDROCHLORIDE

CAPSULE;ORAL

DOXEPIN HYDROCHLORIDE

	EQ 100MG BASE	A212997 005	Jul 24, 2020
NEW RIVER	EQ 10MG BASE	N016987 001	
	EQ 25MG BASE	N016987 002	
	EQ 50MG BASE	N016987 003	
	EQ 75MG BASE	N016987 006	
	EQ 100MG BASE	N016987 004	
	EQ 150MG BASE	N016987 007	Apr 13, 1987
PAR PHARM	EQ 10MG BASE	A071422 002	Nov 09, 1987
	EQ 25MG BASE	A071422 003	Nov 09, 1987
	EQ 50MG BASE	A071422 004	Nov 09, 1987
	EQ 75MG BASE	A071422 005	Nov 09, 1987
	EQ 100MG BASE	A071422 001	Nov 09, 1987
PUREPAC PHARM	EQ 10MG BASE	A073054 001	Dec 28, 1990
	EQ 25MG BASE	A072109 001	Dec 28, 1990
	EQ 50MG BASE	A073055 001	Dec 28, 1990
	EQ 75MG BASE	A072386 001	Sep 08, 1988
	EQ 100MG BASE	A072110 001	Sep 08, 1988
	EQ 150MG BASE	A072387 001	Sep 08, 1988
QUANTUM PHARMICS	EQ 10MG BASE	A070972 001	Sep 29, 1987
	EQ 25MG BASE	A070973 001	Sep 29, 1987
	EQ 50MG BASE	A070931 001	Sep 29, 1987
	EQ 75MG BASE	A070932 001	Sep 29, 1987
	EQ 100MG BASE	A072375 001	Mar 15, 1989
	EQ 150MG BASE	A072376 001	Mar 15, 1989
SANDOZ	EQ 10MG BASE	A071487 001	Mar 02, 1987
	EQ 25MG BASE	A070827 001	May 15, 1986
	EQ 50MG BASE	A070828 001	May 15, 1986
	EQ 75MG BASE	A070825 001	May 15, 1986
	EQ 100MG BASE	A071562 001	Mar 02, 1987
SUN PHARM INDUSTRIES	EQ 25MG BASE	A071502 001	Feb 18, 1988
	EQ 50MG BASE	A071653 001	Feb 18, 1988
	EQ 75MG BASE	A071654 001	Feb 18, 1988
	EQ 100MG BASE	A071521 001	Feb 18, 1988
WATSON LABS	EQ 10MG BASE	A070952 001	Mar 04, 1987
	EQ 10MG BASE	A071485 001	Apr 30, 1987
	EQ 10MG BASE	A072985 001	Mar 29, 1991
	EQ 25MG BASE	A070953 001	May 15, 1986
	EQ 25MG BASE	A071486 001	Apr 30, 1987
	EQ 25MG BASE	A072986 001	Mar 29, 1991
	EQ 50MG BASE	A070954 001	May 15, 1986
	EQ 50MG BASE	A071238 001	Apr 30, 1987
	EQ 75MG BASE	A071326 001	Apr 30, 1987
	EQ 75MG BASE	A071763 001	Feb 09, 1988
	EQ 100MG BASE	A070955 001	May 15, 1986
	EQ 100MG BASE	A071239 001	Apr 30, 1987
	EQ 150MG BASE	A071764 001	Feb 09, 1988
WATSON LABS TEVA	EQ 50MG BASE	A072987 001	Mar 29, 1991
SINEQUAN			
+ PFIZER	EQ 10MG BASE **	N016798 003	
+	EQ 25MG BASE **	N016798 001	
+	EQ 50MG BASE **	N016798 002	
+	EQ 75MG BASE **	N016798 006	
+	EQ 100MG BASE **	N016798 005	
+	EQ 150MG BASE **	N016798 007	
CONCENTRATE;ORAL			
DOXEPIN HYDROCHLORIDE			
PHARM ASSOC	EQ 10MG BASE/ML	A075924 001	Jan 15, 2004
PHARMOBEDIANT CNSLTG	EQ 10MG BASE/ML	A071918 001	Jul 20, 1988
TEVA PHARMS	EQ 10MG BASE/ML	A071609 001	Nov 09, 1987
SINEQUAN			
+ PFIZER	EQ 10MG BASE/ML **	N017516 001	
TABLET;ORAL			
DOXEPIN HYDROCHLORIDE			
ZYDUS PHARMS	EQ 3MG BASE	A202761 001	Aug 16, 2023
	EQ 6MG BASE	A202761 002	Aug 16, 2023

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

HIKMA	0.5MCG	A091433 001	Sep 23, 2011
	1MCG	A091433 002	Jan 14, 2014
	2.5MCG	A091433 003	Jan 14, 2014

INJECTABLE; INJECTION

DOXERCALCIFEROL

AMNEAL	4MCG/2ML (2MCG/ML)	A208975 001	May 24, 2017
EPIC PHARMA LLC	2MCG/ML (2MCG/ML)	A203929 002	Mar 28, 2016
	4MCG/2ML (2MCG/ML)	A203929 001	May 07, 2015
+ HOSPIRA	10MCG/5ML (2MCG/ML)	N208614 002	Jul 24, 2018
SUN PHARM	2MCG/ML (2MCG/ML)	A203875 001	Nov 14, 2019
	4MCG/2ML (2MCG/ML)	A203875 002	Nov 14, 2019

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

PFIZER	2MG/ML	A063165 001	Jan 30, 1991
	200MG/100ML	A063165 002	Jan 30, 1991

DOXORUBICIN HYDROCHLORIDE

ALMAJECT	2MG/ML	A065515 001	Nov 08, 2012
HISUN PHARM HANGZHOU	20MG/VIAL	A206062 001	May 13, 2019
HLTHCARE	2MG/ML	A200146 001	Jul 18, 2012
MYLAN LABS LTD	2MG/ML	A200901 001	Feb 14, 2012
	10MG/VIAL	A200170 001	Oct 28, 2011
PFIZER	10MG/VIAL	N050467 001	
	20MG/VIAL	N050467 003	May 20, 1985
	50MG/VIAL	N050467 002	
	150MG/VIAL	N050467 004	Jul 22, 1987
PHARMACHEMIE BV	2MG/ML	A063336 001	Feb 28, 1995
	10MG/VIAL	A063097 001	May 21, 1990
	20MG/VIAL	A063097 002	May 21, 1990
	50MG/VIAL	A063097 003	May 21, 1990
	200MG/100ML	A063336 004	Feb 28, 1995
TEVA PHARMS USA	2MG/ML	A064140 001	Jul 28, 1995
	200MG/100ML	A064140 002	Jul 28, 1995

RUBEX

BRISTOL MYERS SQUIBB	10MG/VIAL	A062926 001	Apr 13, 1989
	50MG/VIAL	A062926 002	Apr 13, 1989
	100MG/VIAL	A062926 003	Apr 13, 1989

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

IMPAX LABS INC	EQ 150MG BASE	A200065 001	Feb 17, 2011
RISING	EQ 50MG BASE	A208942 001	Jan 21, 2021
	EQ 75MG BASE	A208942 002	Jan 21, 2021
	EQ 100MG BASE	A208942 003	Jan 21, 2021
SANDOZ INC	EQ 50MG BASE	A065032 001	Jun 30, 2000
	EQ 100MG BASE	A065032 002	Jun 30, 2000
STRIDES PHARMA	EQ 75MG BASE	A065055 004	Apr 18, 2005
	EQ 150MG BASE	A202778 001	Jun 08, 2012
WATSON LABS	EQ 50MG BASE	A065041 001	Apr 28, 2000
	EQ 100MG BASE	A065041 002	Apr 28, 2000

FOR SUSPENSION; ORAL

DOXYCHEL

RACHELLE	EQ 25MG BASE/5ML	A061720 001	
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TABLET; ORAL

DOXYCYCLINE

APPCO	EQ 50MG BASE	A065377 001	Nov 07, 2006
	EQ 75MG BASE	A065377 002	Nov 07, 2006
	EQ 100MG BASE	A065377 003	Nov 07, 2006
	EQ 150MG BASE	A065427 001	Jun 07, 2007
SANDOZ INC	EQ 50MG BASE	A065353 001	Nov 27, 2006
	EQ 75MG BASE	A065353 002	Nov 27, 2006
	EQ 100MG BASE	A065353 003	Nov 27, 2006
SUN PHARM INDUSTRIES	EQ 50MG BASE	A065471 001	Apr 17, 2009
	EQ 75MG BASE	A065471 002	Apr 17, 2009
	EQ 100MG BASE	A065471 003	Apr 17, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

ACTICLATE CAP

+ ALMIRALL

EQ 75MG BASE

N208253 001 Apr 26, 2016

DOXY-LEMMON

TEVA

EQ 50MG BASE

A062497 001 Aug 23, 1984

EQ 100MG BASE

A062497 002 Jun 15, 1984

DOXYCYCLINE HYCLATE

AJANTA PHARMA LTD

EQ 50MG BASE

A211012 001 Sep 24, 2018

EQ 100MG BASE

A211012 002 Sep 24, 2018

HALSEY

EQ 50MG BASE

A062119 002 May 24, 1985

EQ 100MG BASE

A062119 001 May 24, 1985

HEATHER

EQ 50MG BASE

A062463 001 Dec 07, 1983

EQ 100MG BASE

A062463 002 Dec 07, 1983

HIKMA INTL PHARMS

EQ 20MG BASE

A065103 001 May 13, 2005

INTERPHARM

EQ 50MG BASE

A062763 001 Sep 02, 1988

EQ 100MG BASE

A062763 002 Sep 02, 1988

MUTUAL PHARM

EQ 50MG BASE

A062418 001 Jan 28, 1983

EQ 100MG BASE

A062418 002 Jan 28, 1983

NESHER PHARMS

EQ 50MG BASE

A208263 001 Nov 22, 2021

EQ 100MG BASE

A208263 002 Nov 22, 2021

NOSTRUM LABS INC

EQ 50MG BASE

A209393 001 Dec 10, 2020

EQ 100MG BASE

A209393 002 Dec 10, 2020

PAR PHARM

EQ 50MG BASE

A062434 001 Oct 19, 1984

EQ 100MG BASE

A062442 001 Dec 22, 1983

PVT FORM

EQ 50MG BASE

A062631 001 Jul 24, 1986

EQ 100MG BASE

A062631 002 Jul 24, 1986

RANBAXY

EQ 50MG BASE

A062479 001 Dec 23, 1983

EQ 100MG BASE

A062479 002 Dec 23, 1983

STRIDES PHARMA

EQ 50MG BASE

A062337 001 Mar 29, 1982

EQ 100MG BASE

A062337 002 Mar 29, 1982

SUPERPHARM

EQ 50MG BASE

A062469 001 Oct 31, 1984

EQ 100MG BASE

A062469 002 Oct 31, 1984

WARNER CHILCOTT

EQ 50MG BASE

A062594 001 Dec 05, 1985

EQ 100MG BASE

A062594 002 Dec 05, 1985

WATSON LABS

EQ 50MG BASE

A061717 001

EQ 50MG BASE

A062142 001

EQ 100MG BASE

A061717 002

EQ 100MG BASE

A062142 002

ZHEJIANG YONGTAI

EQ 50MG BASE

A212610 001 Mar 31, 2020

EQ 100MG BASE

A212610 002 Mar 31, 2020

PERIOSTAT

+ COLLAGENEX

EQ 20MG BASE **

N050744 001 Sep 30, 1998

VIBRAMYCIN

+ PFIZER

EQ 50MG BASE **

N050007 001

CAPSULE, COATED PELLETS; ORAL

DOXYCYCLINE HYCLATE

PLIVA

EQ 100MG BASE

A063187 001 Jun 30, 1992

CAPSULE, DELAYED RELEASE; ORAL

DORYX

+ MAYNE PHARMA

EQ 75MG BASE

N050582 002 Aug 13, 2001

+

EQ 100MG BASE

N050582 001 Jul 22, 1985

WARNER CHILCOTT

EQ 100MG BASE

A062653 001 Oct 30, 1985

DOXYCYCLINE HYCLATE

BAUSCH

EQ 75MG BASE

A065281 001 Dec 21, 2005

EQ 100MG BASE

A065281 002 Dec 21, 2005

INJECTABLE; INJECTION

DOXYCHEL HYCLATE

RACHELLE

EQ 100MG BASE/VIAL

A061953 001

DOXYCYCLINE

HIKMA

EQ 100MG BASE/VIAL

A062450 001 Oct 27, 1983

EQ 100MG BASE/VIAL

A062569 001 Mar 09, 1988

EQ 200MG BASE/VIAL

A062450 002 Oct 27, 1983

EQ 200MG BASE/VIAL

A062569 002 Mar 09, 1988

DOXYCYCLINE HYCLATE

PAR STERILE PRODUCTS

EQ 100MG BASE/VIAL

A216690 001 Dec 07, 2022

VIBRAMYCIN

+ PFIZER

EQ 100MG BASE/VIAL **

N050442 002

+

EQ 200MG BASE/VIAL **

N050442 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE HYCLATE

SYSTEM, EXTENDED RELEASE;PERIODONTAL

ATRIDOX			
+	DEN-MAT	50MG	N050751 001 Sep 03, 1998
TABLET;ORAL			
DOXY-LEMMON			
	TEVA	EQ 100MG BASE	A062581 001 Mar 15, 1985
DOXYCYCLINE HYCLATE			
	ALEMBIC	EQ 100MG BASE	A210536 001 May 14, 2020
	AMNEAL	EQ 100MG BASE	A216599 001 Oct 24, 2022
	AMNEAL PHARMS CO	EQ 75MG BASE	A209372 001 Oct 06, 2017
		EQ 150MG BASE	A209372 002 Oct 06, 2017
	AVET LIFESCIENCES	EQ 100MG BASE	A209969 001 Nov 09, 2018
	HEATHER	EQ 100MG BASE	A062462 001 May 11, 1983
	HERITAGE PHARMA	EQ 20MG BASE	A065163 001 May 13, 2005
	INTERPHARM	EQ 100MG BASE	A062764 001 Sep 02, 1988
	MUTUAL PHARM	EQ 100MG BASE	A062391 001 Sep 30, 1982
	PHARMOBEDIENT	EQ 100MG BASE	A213475 001 Mar 10, 2021
	PRAXGEN	EQ 100MG BASE	A212487 001 Mar 30, 2022
	RISING	EQ 75MG BASE	A209987 001 Oct 05, 2020
		EQ 150MG BASE	A209987 002 Oct 05, 2020
	STRIDES PHARMA	EQ 100MG BASE	A062538 001 Apr 07, 1986
	SUPERPHARM	EQ 100MG BASE	A062494 001 Feb 20, 1985
	WARNER CHILCOTT	EQ 100MG BASE	A062593 001 Aug 28, 1985
	WATSON LABS	EQ 50MG BASE	A062392 001 Mar 31, 1983
		EQ 100MG BASE	A062392 002 Mar 31, 1983
LYMEPAK			
+	CHARTWELL PHARMA	EQ 100MG BASE	N209844 001 Jun 15, 2018
PERIOSTAT			
+	GALDERMA LABS LP	EQ 20MG BASE **	N050783 001 Feb 02, 2001
VIBRA-TABS			
+	PFIZER	EQ 100MG BASE **	N050533 001
TABLET, DELAYED RELEASE;ORAL			
DORYX			
+	MAYNE PHARMA	EQ 50MG BASE	N050795 006 Dec 19, 2014
+		EQ 100MG BASE	N050795 002 May 06, 2005
DORYX MPC			
+	MAYNE PHARMA	EQ 120MG BASE	N050795 008 May 20, 2016
DOXYCYCLINE HYCLATE			
	AUROBINDO PHARMA USA	EQ 150MG BASE	A091052 001 Feb 08, 2012
	IMPAX LABS INC	EQ 75MG BASE	A090505 001 Dec 28, 2010
		EQ 100MG BASE	A090505 002 Dec 28, 2010
	LUPIN	EQ 50MG BASE	A208741 001 Aug 11, 2023
		EQ 60MG BASE	A208741 002 Aug 11, 2023
		EQ 75MG BASE	A208741 003 Aug 11, 2023
		EQ 80MG BASE	A208741 004 Aug 11, 2023
		EQ 100MG BASE	A208741 005 Aug 11, 2023
		EQ 120MG BASE	A208741 006 Aug 11, 2023
		EQ 150MG BASE	A208741 007 Aug 11, 2023
		EQ 200MG BASE	A208741 008 Aug 11, 2023
	RISING	EQ 50MG BASE	A090431 003 May 23, 2016
		EQ 75MG BASE	A090431 001 Dec 28, 2010
		EQ 80MG BASE	A090431 004 Apr 29, 2016
		EQ 100MG BASE	A090431 002 Dec 28, 2010
		EQ 200MG BASE	A090431 005 May 19, 2016
	ZYDUS PHARMS	EQ 75MG BASE	A206772 001 Dec 21, 2018
		EQ 100MG BASE	A206772 002 Dec 21, 2018
		EQ 150MG BASE	A206772 003 Dec 21, 2018
<u>DOXYLAMINE SUCCINATE</u>			
CAPSULE;ORAL			
UNISOM			
	PFIZER	25MG	N019440 001 Feb 05, 1986
TABLET;ORAL			
DECAPRYN			
	SANOFI AVENTIS US	12.5MG	N006412 015
		25MG	N006412 014
DOXY-SLEEP-AID			
	PAR PHARM	25MG	A070156 001 Jul 02, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYLAMINE SUCCINATE

TABLET; ORAL

DOXYLAMINE SUCCINATE

COPLBY PHARM	25MG	A088900	002	Feb 12, 1988
QUANTUM PHARMICS	25MG	A088603	001	Aug 07, 1984

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

BENDECTIN

SANOPI AVENTIS US	10MG;10MG **	N010598	002	
DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE				
ACTAVIS LABS FL INC	20MG;20MG	A212472	001	Mar 01, 2022

DROMOSTANOLONE PROPIONATE

INJECTABLE; INJECTION

DROLBAN

LILLY	50MG/ML	N012936	001	
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DRONABINOL

CAPSULE; ORAL

DRONABINOL

INSYS THERAP	2.5MG	A078501	001	Aug 19, 2011
	5MG	A078501	002	Aug 19, 2011
	10MG	A078501	003	Aug 19, 2011
LANNETT CO INC	2.5MG	A201463	001	May 18, 2018
	5MG	A201463	002	May 18, 2018
	10MG	A201463	003	May 18, 2018

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

ABRAXIS PHARM	2.5MG/ML	A070992	001	Nov 17, 1986
	2.5MG/ML	A070993	001	Nov 17, 1986
ASTRAZENECA	2.5MG/ML	A072018	001	Oct 20, 1988
HIKMA	2.5MG/ML	A208197	001	Dec 14, 2017
HOSPIRA	2.5MG/ML	A071645	001	Apr 07, 1988
	2.5MG/ML	A071981	001	Feb 29, 1988
	2.5MG/ML	A072272	001	Aug 31, 1995
IGI LABS INC	2.5MG/ML	A072019	001	Oct 19, 1988
	2.5MG/ML	A072020	001	Oct 19, 1988
	2.5MG/ML	A072021	001	Oct 19, 1988
LUITPOLD	2.5MG/ML	A072335	001	Oct 24, 1988
SMITH AND NEPHEW	2.5MG/ML	A071750	001	Sep 06, 1988
SOLOPAK	2.5MG/ML	A071754	001	Sep 06, 1988
	2.5MG/ML	A071755	001	Sep 06, 1988
WATSON LABS	2.5MG/ML	A073520	001	Nov 27, 1991
	2.5MG/ML	A073521	001	Nov 27, 1991
	2.5MG/ML	A073523	001	Nov 27, 1991
INAPSINE				
+ RISING	2.5MG/ML	N016796	001	

DROPERIDOL; FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE AND DROPERIDOL

ASTRAZENECA	2.5MG/ML;EQ 0.05MG BASE/ML	A072026	001	Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	A072027	001	Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	A072028	001	Apr 13, 1989
HOSPIRA	2.5MG/ML;EQ 0.05MG BASE/ML	A071982	001	May 04, 1988
INNOVAR				
EPIC PHARMA LLC	2.5MG/ML;EQ 0.05MG BASE/ML	N016049	001	

DROSPIRENONE

TABLET, CHEWABLE; ORAL

DROSPIRENONE

+ EXELTIS USA INC	3.5MG	N216285	001	Jun 29, 2022
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DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

BARR	3MG;0.02MG	A078515	001	Mar 30, 2009
JUBILANT CADISTA	3MG;0.02MG	A209423	001	Dec 22, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

KYRA

SUN PHARM 3MG;0.02MG A202318 001 Jul 23, 2019

TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL

APOTEX 3MG;0.03MG A205876 001 Sep 21, 2016

BARR 3MG;0.03MG A077527 001 May 09, 2008

JUBILANT CADISTA 3MG;0.03MG A210017 001 Sep 10, 2018

KEMEYA

SUN PHARM 3MG;0.03MG A202138 001 Mar 13, 2019

DROXIDOPA

CAPSULE; ORAL

DROXIDOPA

CHARTWELL RX 100MG A214217 001 May 05, 2022

200MG A214217 002 May 05, 2022

300MG A214217 003 May 05, 2022

HIKMA 100MG A212835 001 Feb 18, 2021

200MG A212835 002 Feb 18, 2021

300MG A212835 003 Feb 18, 2021

TEVA PHARMS USA INC 100MG A213162 001 Feb 18, 2021

200MG A213162 002 Feb 18, 2021

300MG A213162 003 Feb 18, 2021

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

DRIZALMA SPRINKLE

+ SUN PHARM EQ 20MG BASE N212516 001 Jul 19, 2019

+ EQ 30MG BASE N212516 002 Jul 19, 2019

+ EQ 40MG BASE N212516 003 Jul 19, 2019

+ EQ 60MG BASE N212516 004 Jul 19, 2019

DULOXETINE HYDROCHLORIDE

APOTEX EQ 20MG BASE A202045 001 Jun 11, 2014

EQ 30MG BASE A202045 002 Jun 11, 2014

EQ 60MG BASE A202045 003 Jun 11, 2014

TEVA PHARMS USA EQ 20MG BASE A090783 001 Dec 11, 2013

EQ 30MG BASE A090783 002 Dec 11, 2013

EQ 60MG BASE A090783 003 Dec 11, 2013

YAOPHARMA CO LTD EQ 20MG BASE A207219 001 Aug 16, 2019

EQ 30MG BASE A207219 002 Aug 16, 2019

EQ 60MG BASE A207219 003 Aug 16, 2019

DUTASTERIDE

CAPSULE; ORAL

DUTASTERIDE

ACTAVIS LABS FL INC 0.5MG A202808 001 Nov 20, 2015

APOTEX 0.5MG A204292 001 Nov 24, 2015

HERITAGE PHARMS INC 0.5MG A207935 001 Oct 13, 2017

HIKMA 0.5MG A202204 001 Nov 23, 2015

MYLAN 0.5MG A203241 001 Jun 14, 2016

NOSTRUM LABS INC 0.5MG A204705 001 Nov 20, 2015

RISING 0.5MG A202530 001 Nov 20, 2015

STRIDES PHARMA 0.5MG A208227 001 Jun 22, 2018

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

ACTAVIS LABS FL INC 0.5MG;0.4MG A202975 001 Nov 20, 2015

DYCLONINE HYDROCHLORIDE

SOLUTION; TOPICAL

DYCLONE

+ ASTRAZENECA 0.5% ** N009925 002

+ 1% ** N009925 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DYDROGESTERONE

TABLET; ORAL

GYNOREST

+	SOLVAY	5MG **	N017388	001
+		10MG **	N017388	002

DYPHYLLINE

ELIXIR; ORAL

NEOTHYLLINE

	TEVA	160MG/15ML	N007794	003
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INJECTABLE; INJECTION

NEOTHYLLINE

	TEVA	250MG/ML	N009088	001
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TABLET; ORAL

DILOR

	SAVAGE LABS	200MG	A084514	001
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DILOR-400

	SAVAGE LABS	400MG	A084751	001
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LUFYLLIN

	MYLAN SPECIALITY LP	200MG	A084566	001
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		400MG	A084566	002
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NEOTHYLLINE

	TEVA	200MG	N007794	001
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		400MG	N007794	002
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ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

	FERA PHARMS LLC	0.03%	N011963	002
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		0.06%	N011963	004
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		0.25%	N011963	003
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ECONAZOLE NITRATE

CREAM; TOPICAL

ECONAZOLE NITRATE

	AUROBINDO PHARMA USA	1%	A210364	001	Apr 18, 2018
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	CHARTWELL RX	1%	A076075	001	Nov 26, 2002
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	VERTICE	1%	A076574	001	Dec 17, 2004
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SPECTAZOLE

+	ALVOGEN	1% **	N018751	001	Dec 23, 1982
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EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

CALCIUM DISODIUM VERSENATE

+	BAUSCH	200MG/ML **	N008922	001
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TABLET; ORAL

CALCIUM DISODIUM VERSENATE

	BAUSCH	500MG	N008922	002
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EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

EDROPHONIUM CHLORIDE

	HOSPIRA	10MG/ML	A040131	001	Feb 24, 1998
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	WATSON LABS	10MG/ML	A040044	001	Mar 20, 1996
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EDROPHONIUM CHLORIDE PRESERVATIVE FREE

	WATSON LABS	10MG/ML	A040043	001	Mar 20, 1996
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ENLON

	MYLAN INSTITUTIONAL	10MG/ML	A088873	001	Aug 06, 1985
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REVERSOL

	ORGANON USA INC	10MG/ML	A089624	001	May 13, 1988
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TENSILON

+	PAI HOLDINGS PHARM	10MG/ML **	N007959	001
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TENSILON PRESERVATIVE FREE

+	PAI HOLDINGS PHARM	10MG/ML **	N007959	002
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EFAVIRENZ

CAPSULE; ORAL

SUSTIVA

+	BRISTOL MYERS SQUIBB	50MG	N020972	001	Sep 17, 1998
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+		100MG **	N020972	002	Sep 17, 1998
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+		200MG	N020972	003	Sep 17, 1998
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EFAVIRENZ

TABLET;ORAL

EFAVIRENZ

AUROBINDO PHARMA	600MG	A205322 001	Aug 30, 2018
MYLAN	600MG	A091471 001	Feb 17, 2016
STRIDES PHARMA	600MG	A078509 001	Jun 16, 2021
SUSTIVA			
+ BRISTOL MYERS SQUIBB	300MG **	N021360 001	Feb 01, 2002
+	600MG	N021360 002	Feb 01, 2002

EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

+ AUROBINDO PHARMA LTD	600MG;300MG;300MG	N022343 001	Aug 15, 2018
MACLEODS PHARMS LTD	400MG;300MG;300MG	N210649 001	Mar 15, 2019

EFINACONAZOLE

SOLUTION;TOPICAL

EFINACONAZOLE

AUROBINDO PHARMA LTD	10%	A212066 001	Mar 29, 2021
LUPIN LTD	10%	A212169 001	Mar 02, 2022
PADAGIS US	10%	A211851 001	Dec 16, 2020
ZYDUS	10%	A212178 001	Jul 15, 2022

EFLORNITHINE HYDROCHLORIDE

CREAM;TOPICAL

VANIQA

+ SKINMEDICA	13.9%	N021145 001	Jul 27, 2000
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INJECTABLE;INJECTION

ORNIDYL

SANOFI AVENTIS US	200MG/ML	N019879 002	Nov 28, 1990
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ELETRIPTAN HYDROBROMIDE

TABLET;ORAL

ELETRIPTAN HYDROBROMIDE

STEVENS J	EQ 20MG BASE	A206787 001	May 25, 2018
	EQ 40MG BASE	A206787 002	May 25, 2018
YUNG SHIN PHARM	EQ 20MG BASE	A209680 001	Jul 13, 2020
	EQ 40MG BASE	A209680 002	Jul 13, 2020

ELIGLUSTAT TARTRATE

CAPSULE;ORAL

ELIGLUSTAT TARTRATE

DR REDDYS	EQ 84MG BASE	A212449 001	Aug 17, 2022
TEVA PHARMS USA INC	EQ 84MG BASE	A212474 001	Dec 27, 2021

ELTROMBOPAG OLAMINE

TABLET;ORAL

PROMACTA

+ NOVARTIS	EQ 100MG ACID **	N022291 005	Nov 16, 2012
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ELVITEGRAVIR

TABLET;ORAL

VITEKTA

+ GILEAD SCIENCES INC	85MG	N203093 001	Sep 24, 2014
+	150MG	N203093 002	Sep 24, 2014

EMEDASTINE DIFUMARATE

SOLUTION/DROPS;OPHTHALMIC

EMADINE

+ NOVARTIS	0.05%	N020706 001	Dec 29, 1997
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EMPAGLIFLOZIN

TABLET;ORAL

EMPAGLIFLOZIN

ZYDUS PHARMS	10MG	A212138 001	Aug 03, 2022
	25MG	A212138 002	Aug 03, 2022

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET;ORAL

EMPAGLIFLOZIN AND METFORMIN HYDROCHLORIDE

ZYDUS PHARMS	5MG;500MG	A212198 001	Jul 07, 2022
	5MG;1GM	A212198 002	Jul 07, 2022
	12.5MG;500MG	A212198 003	Jul 07, 2022
	12.5MG;1GM	A212198 004	Jul 07, 2022

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

APOTEX	200MG;300MG	A208740 001	Jun 16, 2021
CIPLA	200MG;300MG	A090958 001	Apr 02, 2021

ENALAPRIL MALEATE

FOR SOLUTION;ORAL

EPANED KIT

+ SILVERGATE PHARMS	1MG/ML **	N204308 001	Aug 13, 2013
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SOLUTION;ORAL

ENALAPRIL MALEATE

AMNEAL	1MG/ML	A212894 001	Jun 29, 2022
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TABLET;ORAL

ENALAPRIL MALEATE

AIPING PHARM INC	2.5MG	A075178 002	Mar 23, 2001
	5MG	A075178 001	Mar 23, 2001
	10MG	A075178 003	Mar 23, 2001
	20MG	A075178 004	Mar 23, 2001
APOTHECON	2.5MG	A075583 001	Aug 22, 2000
	5MG	A075583 002	Aug 22, 2000
	10MG	A075583 003	Aug 22, 2000
	20MG	A075583 004	Aug 22, 2000
AUROBINDO PHARMA USA	2.5MG	A075480 001	Aug 22, 2000
	5MG	A075480 002	Aug 22, 2000
	10MG	A075480 003	Aug 22, 2000
	20MG	A075480 004	Aug 22, 2000
BEXIMCO PHARMS USA	2.5MG	A075621 001	Aug 22, 2000
	5MG	A075621 002	Aug 22, 2000
	10MG	A075621 003	Aug 22, 2000
	20MG	A075621 004	Aug 22, 2000
CHARTWELL RX	2.5MG	A075048 001	Aug 22, 2000
	5MG	A075048 002	Aug 22, 2000
	10MG	A075048 003	Aug 22, 2000
	20MG	A075048 004	Aug 22, 2000
IVAX SUB TEVA PHARMS	2.5MG	A075482 001	Aug 22, 2000
	5MG	A075482 002	Aug 22, 2000
	10MG	A075482 003	Aug 22, 2000
	20MG	A075482 004	Aug 22, 2000
KRKA DD NOVO MESTO	2.5MG	A075370 001	Aug 22, 2000
	5MG	A075370 002	Aug 22, 2000
	10MG	A075369 001	Aug 22, 2000
	20MG	A075369 002	Aug 22, 2000
MYLAN	2.5MG	A075472 001	Aug 22, 2000
	5MG	A075472 002	Aug 22, 2000
	10MG	A075472 003	Aug 22, 2000
	20MG	A075472 004	Aug 22, 2000
SUN PHARM INDS LTD	2.5MG	A075556 001	Aug 22, 2000
	5MG	A075556 002	Aug 22, 2000
	10MG	A075556 003	Aug 22, 2000
	20MG	A075556 004	Aug 22, 2000
WATSON LABS	2.5MG	A075501 001	Aug 22, 2000
	5MG	A075501 002	Aug 22, 2000
	10MG	A075501 003	Aug 22, 2000
	20MG	A075501 004	Aug 22, 2000

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

LEXXEL

ASTRAZENECA	5MG;2.5MG	N020668 002	Oct 28, 1998
	5MG;5MG	N020668 001	Dec 27, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	5MG;12.5MG	A076116 001	Sep 19, 2001
	10MG;25MG	A076116 002	Sep 19, 2001
IVAX SUB TEVA PHARMS	5MG;12.5MG	A075736 001	Mar 25, 2003
	10MG;25MG	A075736 002	Mar 25, 2003
NOSTRUM LABS INC	5MG;12.5MG	A076486 001	Oct 27, 2004
	10MG;25MG	A076486 002	Oct 27, 2004
RISING	5MG;12.5MG	A075624 001	Sep 18, 2001
	10MG;25MG	A075624 002	Sep 18, 2001

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

HOSPIRA	1.25MG/ML	A075456 001	Aug 22, 2000
	1.25MG/ML	A075571 001	Aug 22, 2000
VASOTEC			
+ BIOVAIL LABS INTL	1.25MG/ML **	N019309 001	Feb 09, 1988

ENCORAFENIB

CAPSULE; ORAL

BRAFTOVI

+ ARRAY BIOPHARMA INC	50MG	N210496 001	Jun 27, 2018
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ENFLURANE

LIQUID; INHALATION

ENFLURANE

ABBOTT	99.9%	A070803 001	Sep 08, 1987
PIRAMAL CRITICAL	99.9%	A074396 001	Jul 29, 1994
ETHRANE			
BAXTER HLTHCARE	99.9%	N017087 001	

ENOXACIN

TABLET; ORAL

PENETREX

SANOFI AVENTIS US	200MG	N019616 004	Dec 31, 1991
	400MG	N019616 005	Dec 31, 1991

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

LOVENOX (PRESERVATIVE FREE)

+ SANOFI AVENTIS US	90MG/0.6ML (150MG/ML) **	N020164 006	Jun 02, 2000
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ENTACAPONE

TABLET; ORAL

ENTACAPONE

MYLAN PHARMS INC	200MG	A202394 001	May 13, 2013
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ENTECAVIR

TABLET; ORAL

ENTECAVIR

ACCORD HLTHCARE	0.5MG	A205824 001	Aug 25, 2017
	1MG	A205824 002	Aug 25, 2017
CASI PHARMS INC	0.5MG	A206672 001	May 11, 2017
	1MG	A206672 002	May 11, 2017
CHARTWELL RX	0.5MG	A206294 001	Nov 23, 2016
	1MG	A206294 002	Nov 23, 2016
RISING	0.5MG	A206226 001	Mar 26, 2019
	1MG	A206226 002	Mar 26, 2019
SUNSHINE	0.5MG	A211978 001	May 20, 2020
	1MG	A211978 002	May 20, 2020
TEVA PHARMS USA	0.5MG	A202122 001	Aug 26, 2014
	1MG	A202122 002	Aug 26, 2014
YAOPHARMA CO LTD	0.5MG	A212201 001	Nov 04, 2019
	1MG	A212201 002	Nov 04, 2019
YUNG SHIN PHARM	0.5MG	A208195 001	Nov 10, 2021
	1MG	A208195 002	Nov 10, 2021

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENZALUTAMIDE

CAPSULE; ORAL

ENZALUTAMIDE

ACTAVIS LABS FL INC 40MG
APOTEX 40MGA209614 001 May 14, 2021
A209645 001 Apr 22, 2022EPHEDRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

REZIPRES

+ DR REDDYS LABS SA 47MG/ML (47MG/ML)
+ 47MG/5ML (9.4MG/ML)N213536 002 Jun 14, 2021
N213536 003 Jun 14, 2021EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

EPHEDRINE SULFATE

RENEW PHARMS 50MG/ML (50MG/ML)
ZYDUS PHARMS 50MG/ML (50MG/ML)N208609 001 Mar 01, 2017
A217276 001 May 16, 2023EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ELESTAT

+ ALLERGAN 0.05%

N021565 001 Oct 16, 2003

EPINASTINE HYDROCHLORIDE

CHARTWELL RX 0.05%
EPIC PHARMA LLC 0.05%
SUN PHARM INDS 0.05%A203384 001 Dec 07, 2016
A204055 001 May 05, 2017
A091626 001 Oct 31, 2011EPINEPHRINE

AEROSOL, METERED; INHALATION

BRONKAID MIST

STERLING 0.25MG/INH

N016803 001

EPINEPHRINE

ARMSTRONG PHARMS 0.2MG/INH

A087907 001 May 23, 1984

PRIMATENE MIST

WYETH CONS 0.2MG/INH

N016126 001

INJECTABLE; INJECTION

SUS-PHRINE SULFITE FREE

FOREST LABS 1.5MG/AMP
5MG/MLN007942 003 Feb 05, 1999
N007942 001

INJECTABLE; INTRAMUSCULAR

EPI E Z PEN JR

MYLAN SPECIALITY LP 0.15MG/DELIVERY

N019430 004 Aug 03, 1995

EPIPEN E Z PEN

MYLAN SPECIALITY LP 0.3MG/DELIVERY

N019430 003 Aug 03, 1995

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

TWINJECT 0.15

IMPAX EQ 0.15MG/DELIVERY

N020800 002 May 28, 2004

TWINJECT 0.3

IMPAX EQ 0.3MG/DELIVERY

N020800 001 May 30, 2003

SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

EPINEPHRINE

AM REGENT EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A207568 001 Jul 06, 2018

SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

SYMJEPI

+ ADAMIS PHARMS CORP 0.15MG/0.3ML (0.15MG/0.3ML)

N207534 002 Sep 27, 2018

EPINEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

BRONITIN MIST

WYETH CONS 0.3MG/INH

N016126 002

MEDIHALER-EPI

3M 0.3MG/INH

N010374 003

EPINEPHRINE BITARTRATE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA 0.005MG/ML; 1% **
+ 0.005MG/ML; 1.5% **
+ DENTSPLY PHARM 0.005MG/ML; 1.5% **N017751 006
N017751 007
N021384 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPINEPHRINE BITARTRATE; PRILUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE

ASTRAZENECA 0.005MG/ML; 4% N014763 008

EPINEPHRINE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA 0.005MG/ML; 0.5% ** N017751 004

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ALPHACAINE HYDROCHLORIDE W/ EPINEPHRINE

CARLISLE 0.01MG/ML; 2% A084720 001

0.02MG/ML; 2% A084732 001

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

BELMORA LLC 0.01MG/ML; 2% A080504 004 Oct 19, 1983

0.02MG/ML; 2% A080504 005 Oct 19, 1983

EASTMAN KODAK 0.01MG/ML; 2% A040057 002 Feb 26, 1993

0.02MG/ML; 2% A040057 001 Feb 26, 1993

HOSPIRA 0.005MG/ML; 1% A089649 001 Jun 21, 1988

0.005MG/ML; 1.5% A089650 001 Jun 21, 1988

0.01MG/ML; 2% A078772 001 May 12, 2008

0.02MG/ML; 2% A078772 002 May 12, 2008

WEST-WARD PHARMS INT 0.01MG/ML; 1% A080406 001

0.01MG/ML; 2% A080406 002

LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE

ABBOTT 0.01MG/ML; 1% A083154 001

BEL MAR 0.01MG/ML; 1% A080820 001

0.01MG/ML; 2% A080757 001

DELL LABS 0.01MG/ML; 1% A083389 001

0.01MG/ML; 2% A083390 001

INTL MEDICATION 0.01MG/ML; 1% A086402 001

WATSON LABS 0.01MG/ML; 1% A080377 003

0.01MG/ML; 1% A085463 001

0.01MG/ML; 2% A080377 004

LIDOCATON

PHARMATON 0.01MG/ML; 2% A084729 001 Aug 17, 1983

0.02MG/ML; 2% A084728 001 Aug 17, 1983

OCTOCAINE

SEPTODONT 0.01MG/ML; 2% A084048 001

0.02MG/ML; 2% A084048 002

XYLOCAINE DENTAL WITH EPINEPHRINE

DENTSPLY PHARM 0.01MG/ML; 2% ** N021381 001

0.02MG/ML; 2% ** N021381 002

XYLOCAINE W/ EPINEPHRINE

ASTRAZENECA 0.005MG/ML; 1% N010418 006

0.005MG/ML; 1.5% N010418 010

0.005MG/ML; 2% N010418 008

FRESENIUS KABI USA 0.01MG/ML; 2% N006488 003

PATCH; IONTOPHORESIS, TOPICAL

LIDOSITE TOPICAL SYSTEM KIT

VYTERIS 1.05MG/PATCH; 100MG/PATCH N021504 001 May 06, 2004

SOLUTION; IONTOPHORESIS

IONTOCAINE

IOMED 0.01MG/ML; 2% N020530 001 Dec 21, 1995

SOLUTION; IONTOPHORESIS, TOPICAL

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

EMPI 0.01MG/ML; 2% N021486 001 Oct 26, 2004

EPINEPHRINE; PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE W/ EPINEPHRINE

BEL MAR 0.02MG/ML; 1% A080758 001

0.02MG/ML; 2% A080759 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

EPIRUBICIN HYDROCHLORIDE

ACTAVIS TOTOWA	10MG/5ML (2MG/ML)	A065445 001	Sep 18, 2008
	50MG/25ML (2MG/ML)	A065445 002	Sep 18, 2008
	200MG/100ML (2MG/ML)	A065445 003	Sep 18, 2008
EBEWE PHARMA	50MG/25ML (2MG/ML)	A065339 001	Dec 22, 2009
	200MG/100ML (2MG/ML)	A065339 002	Dec 22, 2009
FRESENIUS KABI USA	10MG/5ML (2MG/ML)	A065408 001	Oct 15, 2007
	50MG/25ML (2MG/ML)	A065408 002	Oct 15, 2007
	150MG/75ML (2MG/ML)	A065408 003	Oct 15, 2007
	200MG/100ML (2MG/ML)	A065408 004	Oct 15, 2007
	200MG/100ML (2MG/ML)	A065411 001	Aug 20, 2007
	50MG/25ML (2MG/ML)	A065411 002	Aug 20, 2007
HOSPIRA	10MG/5ML (2MG/ML)	A065343 001	Apr 19, 2007
	50MG/25ML (2MG/ML)	A065343 002	Apr 19, 2007
	150MG/75ML (2MG/ML)	A065343 003	Apr 19, 2007
	200MG/100ML (2MG/ML)	A065343 004	Apr 19, 2007
MYLAN INSTITUTIONAL	50MG/25ML (2MG/ML)	A065371 001	Nov 28, 2007
	200MG/100ML (2MG/ML)	A065371 002	Nov 28, 2007
MYLAN LABS LTD	50MG/25ML (2MG/ML)	A091599 001	Mar 12, 2012
	200MG/100ML (2MG/ML)	A091599 002	Mar 12, 2012
ZENNOVA	50MG/25ML (2MG/ML)	A090266 001	Apr 15, 2011
	200MG/100ML (2MG/ML)	A090266 002	Apr 15, 2011

POWDER; INTRAVENOUS

EPIRUBICIN HYDROCHLORIDE

HOSPIRA	50MG/VIAL	N050807 001	Sep 15, 2006
	200MG/VIAL	N050807 002	Sep 15, 2006

EPLERENONE

TABLET; ORAL

INSPRA

UPJOHN	100MG	N021437 003	Sep 27, 2002
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EPROSARTAN MESYLATE

TABLET; ORAL

EPROSARTAN MESYLATE

MYLAN PHARMS INC	EQ 400MG BASE	A202012 001	Nov 16, 2011
	EQ 600MG BASE	A202012 002	Nov 16, 2011
TEVETEN			
ABBVIE	EQ 300MG BASE **	N020738 004	Dec 22, 1997
+	EQ 400MG BASE **	N020738 005	Dec 22, 1997
+	EQ 600MG BASE **	N020738 006	May 27, 1999

EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEVETEN HCT

ABBVIE	600MG; 12.5MG	N021268 001	Nov 01, 2001
	600MG; 25MG	N021268 002	Nov 01, 2001

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

AMNEAL PHARMS	2MG/ML	A205581 001	Dec 08, 2016	
	75MG/100ML	A205581 002	Dec 08, 2016	
BAXTER HLTHCARE CORP	2MG/ML	A208554 001	Nov 23, 2018	
HYBIO	2MG/ML	A207864 001	Mar 20, 2020	
	75MG/100ML	A207864 002	Mar 20, 2020	
RISING	2MG/ML	A204589 001	Apr 18, 2017	
	75MG/100ML	A204589 002	Apr 18, 2017	
TEVA PHARMS USA	75MG/100ML	A091555 001	Jun 05, 2015	
USV	2MG/ML	A204361 001	Mar 14, 2019	
	2MG/ML	A204362 001	Mar 11, 2019	
	75MG/100ML	A204361 002	Mar 14, 2019	
INTEGRILIN				
+	MSD SUB MERCK	2MG/ML **	N020718 001	May 18, 1998
+		75MG/100ML **	N020718 002	May 18, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERGOCALCIFEROL

CAPSULE; ORAL

DELTALIN

LILLY 50,000 IU A080884 001

ERGOCALCIFEROL

SIGMAPHARM LABS LLC 50,000 IU A091004 001 Jul 14, 2010

SUN PHARM INDS INC 50,000 IU A040865 001 Dec 29, 2009

VITAMIN D

CHARTWELL MOLECULAR 50,000 IU A080825 001

CHASE CHEM 50,000 IU A080747 001

EVERYLIFE 50,000 IU A080956 001

IMPAX LABS 50,000 IU A080951 001

VITARINE 50,000 IU A084053 001

WEST WARD 50,000 IU A083102 001

ERGOLOID MESYLATES

CAPSULE; ORAL

HYDERGINE LC

NOVARTIS 1MG N018706 001 Jan 18, 1983

SOLUTION; ORAL

HYDERGINE

NOVARTIS 1MG/ML N018418 001

TABLET; ORAL

ERGOLOID MESYLATES

MUTUAL PHARM 1MG A088891 001 Nov 01, 1985

WATSON LABS 1MG A086433 001 May 27, 1982

1MG A087244 001 Aug 16, 1982

GERIMAL

WATSON LABS 1MG A088207 001 Mar 22, 1984

HYDERGINE

NOVARTIS 0.5MG ** N017993 003

+ 1MG ** N017993 001

TABLET; SUBLINGUAL

ALKERGOT

SANDOZ 0.5MG A085153 001

1MG A087417 001

CIRCANOL

3M 0.5MG A084868 001

1MG A085809 001

DEAPRIL-ST

BRISTOL MYERS SQUIBB 1MG A085020 002

ERGOLOID MESYLATES

KV PHARM 0.5MG A085899 001

0.5MG A086265 001

1MG A085900 001

1MG A086264 001

LEDERLE 0.5MG A086984 001

1MG A086985 001

SUN PHARM INDUSTRIES 0.5MG A087407 001

1MG A087552 001

SUPERPHARM 0.5MG A089233 001 Sep 23, 1986

1MG A089234 001 Sep 23, 1986

VANGARD 0.5MG A088013 001 Sep 20, 1982

1MG A088014 001 Sep 20, 1982

WATSON LABS 0.5MG A084930 001

0.5MG A087233 001

1MG A085177 001

1MG A087183 001

GERIMAL

WATSON LABS 0.5MG A086189 001

1MG A086188 001

HYDERGINE

NOVARTIS 0.5MG N009087 002

1MG N009087 001

HYDROGENATED ERGOT ALKALOIDS

IVAX PHARMS 0.5MG A087186 001

1MG A087185 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERGOTAMINE TARTRATE

AEROSOL, METERED; INHALATION

MEDIHALER ERGOTAMINE

3M

0.36MG/INH

N012102 001

TABLET; SUBLINGUAL

ERGOSTAT

WATSON LABS INC

2MG

A088337 001 Jun 08, 1984

WIGRETTES

ORGANON USA INC

2MG

A086750 001 Jul 29, 1982

ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

ERLOTINIB HYDROCHLORIDE

ACCORD HLTHCARE

EQ 25MG BASE

A211083 001 Jul 02, 2020

EQ 100MG BASE

A211083 002 Jul 02, 2020

EQ 150MG BASE

A211083 003 Jul 02, 2020

EUGIA PHARMA

EQ 25MG BASE

A216342 001 Jun 22, 2022

EQ 100MG BASE

A216342 002 Jun 22, 2022

EQ 150MG BASE

A216342 003 Jun 22, 2022

TARCEVA

+ OSI PHARMS

EQ 25MG BASE

N021743 001 Nov 18, 2004

+

EQ 100MG BASE

N021743 002 Nov 18, 2004

+

EQ 150MG BASE

N021743 003 Nov 18, 2004

ERTUGLIFLOZIN

TABLET; ORAL

ERTUGLIFLOZIN

AUROBINDO PHARMA LTD 5MG

A216947 001 Jul 13, 2023

15MG

A216947 002 Jul 13, 2023

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYC

PARKE DAVIS

250MG

A062546 001 Jul 25, 1985

250MG

A062618 001 Sep 25, 1985

WARNER CHILCOTT LLC

250MG

A062338 001

ERYC 125

PARKE DAVIS

125MG

A062648 001 Oct 24, 1985

ERYC SPRINKLES

HOSPIRA

125MG

N050593 001 Jul 22, 1985

ERYTHROMYCIN

BARR

250MG

A063098 001 May 04, 1989

GEL; TOPICAL

E-GLADES

MYLAN

2%

A065009 001 Mar 18, 2002

EMGEL

ALTANA

2%

A063107 001 Aug 23, 1991

ERYTHROMYCIN

VERTICE

2%

A208154 001 Jul 19, 2017

LOTION; TOPICAL

E-SOLVE 2

SYOSSET

2%

A062467 001 Jul 03, 1985

OINTMENT; OPHTHALMIC

ERYTHROMYCIN

PHARMADERM

5MG/GM

A062446 001 Sep 26, 1983

PHARMAFAIR

5MG/GM

A062481 001 Apr 05, 1984

ILOTYCIN

DISTA

0.5% **

N050368 001

OINTMENT; TOPICAL

AKNE-MYCIN

+ BAUSCH

2%

N050584 001 Jan 10, 1985

POWDER; FOR RX COMPOUNDING

ERYTHROMYCIN

PADDOCK LLC

100%

N050610 001 Nov 07, 1986

SOLUTION; TOPICAL

A/T/S

TARO

2%

A062405 001 Nov 18, 1982

C-SOLVE-2

FOUGERA PHARMS

2%

A062468 001 Jul 03, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN

SOLUTION;TOPICAL

ERYDERM					
ARBOR PHARMS INC	2%		A062290	001	
ERYMAX					
MERZ PHARMS	2%		A062508	002	Jul 11, 1985
ERYTHRO-STATIN					
EPIC PHARMA LLC	2%		A064101	001	Oct 22, 1996
ERYTHROMYCIN					
ALPHARMA US PHARMS	1.5%		A062328	001	Apr 19, 1982
	2%		A062326	001	Apr 19, 1982
	2%		A062327	001	Apr 19, 1982
	2%		A062342	001	Feb 25, 1982
	2%		A062957	001	Jul 21, 1988
BAUSCH	2%		A064039	001	Jan 27, 1994
FOUGERA PHARMS	2%		A064187	001	Sep 30, 1997
LILLY	2%		N050532	001	
PAI HOLDINGS PHARM	2%		A208100	001	Nov 20, 2017
PHARMAFAIR	1.5%		A062485	001	Jul 11, 1984
	2%		A062616	001	Jul 25, 1985
PHARMOBEDIENT CNSLTG	2%		A062825	001	Oct 23, 1987
RENAISSANCE PHARMA	2%		A064127	001	Feb 14, 1997
SANSAC					
DOW PHARM	2%		A062522	001	Jan 24, 1985
STATICIN					
+ WESTWOOD SQUIBB	1.5% **		N050526	001	
T-STAT					
WESTWOOD SQUIBB	2% **		A062436	001	Mar 09, 1983
SWAB;TOPICAL					
C-SOLVE-2					
IVAX SUB TEVA PHARMS	2%		A062751	001	Jul 30, 1993
ERYCETTE					
+ JOHNSON AND JOHNSON	2% **		N050594	001	Feb 15, 1985
ERYTHROMYCIN					
FOUGERA PHARMS	2%		A065320	001	Jul 25, 2006
MYLAN	2%		A064128	001	Jul 03, 1996
T-STAT					
WESTWOOD SQUIBB	2%		A062748	001	Jul 23, 1987
TABLET;ORAL					
ERYTHROMYCIN					
ZYDUS PHARMS	250MG		A215440	001	Aug 31, 2023
	500MG		A215440	002	Aug 31, 2023
TABLET, COATED PARTICLES;ORAL					
PCE					
+ AZURITY	333MG		N050611	001	Sep 09, 1986
+	500MG		N050611	002	Aug 22, 1990
TABLET, DELAYED RELEASE;ORAL					
E-BASE					
BARR	333MG		A063028	001	May 15, 1990
	333MG		A063086	001	May 15, 1990
	500MG		A062999	001	Nov 25, 1988
E-MYCIN					
ARBOR PHARMS INC	250MG		A060272	001	
	333MG		A060272	002	
ILOTYCIN					
DISTA	250MG		A061910	001	
R-P MYCIN					
SOLVAY	250MG		A061659	001	
ROBIMYCIN					
ROBINS AH	250MG		A061633	001	

ERYTHROMYCIN ESTOLATE

CAPSULE;ORAL

ERYTHROMYCIN ESTOLATE

BARR	EQ 125MG BASE		A062162	001	
	EQ 250MG BASE		A062162	002	
IVAX SUB TEVA PHARMS	EQ 250MG BASE		A062237	001	
WATSON LABS	EQ 250MG BASE		A062087	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN ESTOLATE

CAPSULE; ORAL

ILOSONE

LILLY

EQ 125MG BASE

A061897 001

EQ 250MG BASE

A061897 002

FOR SUSPENSION; ORAL

ILOSONE

DISTA

EQ 125MG BASE/5ML

A061893 001

SUSPENSION; ORAL

ERYTHROMYCIN ESTOLATE

ALPHARMA US PHARMS

EQ 125MG BASE/5ML

A062353 001 Nov 18, 1982

EQ 250MG BASE/5ML

A062409 001 Dec 16, 1982

COSETTE

EQ 125MG BASE/5ML

A062169 001 Oct 17, 1990

EQ 250MG BASE/5ML

A062169 002 Oct 17, 1990

LIFE LABS

EQ 250MG BASE/5ML

A062362 001 Dec 17, 1982

ILOSONE

LILLY

EQ 125MG BASE/5ML

A061894 001

EQ 125MG BASE/5ML

N050010 001

EQ 250MG BASE/5ML

A061894 002

EQ 250MG BASE/5ML

N050010 002

SUSPENSION/DROPS; ORAL

ILOSONE

LILLY

EQ 100MG BASE/ML

A061894 003

TABLET; ORAL

ILOSONE

LILLY

EQ 500MG BASE

A061896 001

TABLET, CHEWABLE; ORAL

ILOSONE

DISTA

EQ 125MG BASE

A061895 001

EQ 250MG BASE

A061895 002

ERYTHROMYCIN ESTOLATE; SULFISOXAZOLE ACETYL

SUSPENSION; ORAL

ILOSONE SULFA

LILLY

EQ 125MG BASE/5ML; EQ 600MG BASE/5ML

N050599 001 Sep 29, 1989

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

ERYTHROMYCIN ETHYLSUCCINATE

PAR PHARM INC

EQ 200MG BASE/5ML

A211991 001 Oct 23, 2019

EQ 400MG BASE/5ML

A211991 002 Oct 23, 2019

PEDIAMYCIN

ROSS LABS

EQ 200MG BASE/5ML

A062305 001

SUSPENSION; ORAL

E-MYCIN E

PHARMACIA AND UPJOHN

EQ 200MG BASE/5ML

A062198 001

EQ 400MG BASE/5ML

A062198 002

E.E.S. 200

ARBOR PHARMS LLC

EQ 200MG BASE/5ML **

A061639 001

E.E.S. 400

ARBOR PHARMS LLC

EQ 400MG BASE/5ML **

A061639 002

ERYTHROMYCIN ETHYLSUCCINATE

ALPHARMA US PHARMS

EQ 200MG BASE/5ML

A062200 001

EQ 400MG BASE/5ML

A062200 002

DISTA

EQ 200MG BASE/5ML

A062177 001

EQ 400MG BASE/5ML

A062177 002

NASKA

EQ 400MG BASE/5ML

A062674 001 Mar 10, 1987

PARKE DAVIS

EQ 200MG BASE/5ML

A062231 001

EQ 400MG BASE/5ML

A062231 002

PHARMAFAIR

EQ 200MG BASE/5ML

A062559 001 Mar 15, 1985

EQ 400MG BASE/5ML

A062558 001 Mar 15, 1985

PEDIAMYCIN

ARBOR PHARMS LLC

EQ 200MG BASE/5ML

A062304 001

PEDIAMYCIN 400

ARBOR PHARMS LLC

EQ 400MG BASE/5ML

A062304 002

WYAMYCIN E

WYETH AYERST

EQ 200MG BASE/5ML

A062123 002

EQ 400MG BASE/5ML

A062123 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION/DROPS;ORAL

PEDIAMYCIN

ROSS LABS EQ 100MG BASE/2.5ML A062305 002

TABLET;ORAL

E.E.S. 400

ARBOR PHARMS LLC EQ 400MG BASE A061905 001

ERYTHROMYCIN ETHYLSUCCINATE

AUROBINDO PHARMA USA EQ 400MG BASE A062847 001 Sep 14, 1988

BARR EQ 400MG BASE A062256 001

TABLET, CHEWABLE;ORAL

E.E.S.

AZURITY EQ 200MG BASE N050297 002

ERYPED

AZURITY EQ 200MG BASE N050297 003 Jul 05, 1988

PEDIAMYCIN

ROSS LABS EQ 200MG BASE A062306 001

ERYTHROMYCIN ETHYLSUCCINATE; SULFISOXAZOLE ACETYL

GRANULE;ORAL

ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL

BARR EQ 200MG BASE/5ML;EQ 600MG BASE/5ML A062759 001 May 20, 1988

ERYZOLE

ALRA EQ 200MG BASE/5ML;EQ 600MG BASE/5ML A062758 001 Jun 15, 1988

PEDIAZOLE

ROSS LABS EQ 200MG BASE/5ML;EQ 600MG BASE/5ML ** N050529 001

ERYTHROMYCIN GLUCEPTATE

INJECTABLE; INJECTION

ILOTYCIN GLUCEPTATE

DISTA EQ 250MG BASE/VIAL N050370 001

EQ 500MG BASE/VIAL N050370 002

EQ 1GM BASE/VIAL N050370 003

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

ABBOTT EQ 500MG BASE/VIAL A062586 001 Jan 04, 1988

EQ 1GM BASE/VIAL A062586 002 Jan 04, 1988

HOSPIRA EQ 500MG BASE/VIAL N050182 002

EQ 1GM BASE/VIAL A062638 002 Oct 31, 1986

EQ 1GM BASE/VIAL N050182 003

+

EQ 1GM BASE/VIAL N050609 002 Sep 24, 1986

ERYTHROMYCIN

ELKINS SINN EQ 500MG BASE/VIAL A062563 001 Mar 28, 1985

EQ 1GM BASE/VIAL A062563 002 Mar 28, 1985

ERYTHROMYCIN LACTOBIONATE

ABRAXIS PHARM EQ 500MG BASE/VIAL A062604 001 Nov 24, 1986

EQ 1GM BASE/VIAL A062604 002 Nov 24, 1986

BAXTER HLTHCARE EQ 500MG BASE/VIAL A062993 001 May 09, 1989

EQ 1GM BASE/VIAL A062993 002 May 09, 1989

EXELA PHARMA EQ 500MG BASE/VIAL A211086 001 Sep 17, 2020

TEVA PARENTERAL EQ 500MG BASE/VIAL A063253 001 Jul 30, 1993

EQ 1GM BASE/VIAL A063253 002 Jul 30, 1993

ERYTHROMYCIN STEARATE

TABLET;ORAL

BRISTAMYCIN

BRISTOL EQ 250MG BASE A061304 001

EQ 250MG BASE A061887 001

ERYPAR

PARKE DAVIS EQ 250MG BASE A062032 001

EQ 500MG BASE A062032 002

WARNER CHILCOTT EQ 250MG BASE A062322 001

ERYTHROCIN STEARATE

ARBOR PHARMS LLC EQ 125MG BASE A060359 002

EQ 500MG BASE A060359 003

ERYTHROMYCIN STEARATE

ANI PHARMS EQ 250MG BASE A061461 001

EQ 250MG BASE A061591 001

EQ 500MG BASE A061461 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROMYCIN STEARATE

	EQ 500MG BASE	A063179 001	May 15, 1990
LEDERLE	EQ 250MG BASE	A062089 001	
	EQ 500MG BASE	A062089 002	
MYLAN	EQ 250MG BASE	A061505 001	
	EQ 500MG BASE	A061505 002	
PUREPAC PHARM	EQ 250MG BASE	A061743 001	
WATSON LABS	EQ 250MG BASE	A062121 002	
	EQ 500MG BASE	A062121 001	
ETHRIL 250			
BRISTOL MYERS SQUIBB	EQ 250MG BASE	A061605 001	
ETHRIL 500			
BRISTOL MYERS SQUIBB	EQ 500MG BASE	A061605 002	
PFIZER-E			
PFIZER	EQ 250MG BASE	A061791 001	
	EQ 500MG BASE	A061791 002	
WYAMYCIN S			
WYETH AYERST	EQ 250MG BASE	A061675 001	
	EQ 500MG BASE	A061675 002	

ESCITALOPRAM OXALATE

CAPSULE; ORAL

ESCITALOPRAM OXALATE

MYLAN PHARMS INC	EQ 5MG BASE	A077660 001	Jul 31, 2007
	EQ 10MG BASE	A077660 002	Jul 31, 2007
	EQ 20MG BASE	A077660 003	Jul 31, 2007

SOLUTION; ORAL

ESCITALOPRAM OXALATE

ANTRIM PHARMS LLC	EQ 5MG BASE/5ML	A203967 001	May 26, 2015
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LEXAPRO

+ ABBVIE

	EQ 5MG BASE/5ML **	N021365 001	Nov 27, 2002
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TABLET; ORAL

ESCITALOPRAM OXALATE

DASH PHARMS	EQ 5MG BASE	A077550 001	May 14, 2015
	EQ 10MG BASE	A077550 002	May 14, 2015
	EQ 20MG BASE	A077550 003	May 14, 2015
HIKMA PHARMS	EQ 5MG BASE	A078766 001	Sep 11, 2012
	EQ 10MG BASE	A078766 002	Sep 11, 2012
	EQ 20MG BASE	A078766 003	Sep 11, 2012
TEVA PHARMS USA	EQ 5MG BASE	A076765 001	Mar 14, 2012
	EQ 10MG BASE	A076765 002	Mar 14, 2012
	EQ 20MG BASE	A076765 003	Mar 14, 2012

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

BAXTER HLTHCARE	10MG/ML	N019386 003	Aug 15, 1988
	20MG/ML	N019386 007	May 28, 2003

ESMOLOL HYDROCHLORIDE

AM REGENT	10MG/ML	A201126 001	Feb 20, 2015
FRESENIUS KABI USA	10MG/ML	A076573 001	May 02, 2005
	20MG/ML	A076573 002	Aug 23, 2023

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

ESOMEPRAZOLE MAGNESIUM

AMNEAL PHARMS NY	EQ 20MG BASE	A209647 001	Apr 10, 2019
	EQ 40MG BASE	A209647 002	Apr 10, 2019
HEC PHARM	EQ 20MG BASE	A207265 002	May 18, 2018
	EQ 40MG BASE	A207265 001	May 18, 2018
HETERO LABS LTD III	EQ 20MG BASE	A202784 001	Sep 21, 2015
	EQ 40MG BASE	A202784 002	Sep 21, 2015
TORRENT	EQ 20MG BASE	A203636 001	Oct 19, 2015
	EQ 40MG BASE	A203636 002	Oct 19, 2015

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

HETERO LABS LTD III	EQ 20MG BASE	A208939 001	May 28, 2020
MYLAN	EQ 20MG BASE	A212376 001	Oct 16, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESOMEPRAZOLE MAGNESIUM

TABLET, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

P AND L

EQ 20MG BASE

A209202 001 Mar 05, 2019

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

+ DEXCEL

EQ 20MG BASE

N214278 001 Oct 20, 2020

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE;ORAL

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

ACTAVIS LABS FL INC

EQ 20MG BASE;375MG

A204470 001 Aug 24, 2022

EQ 20MG BASE;500MG

A204470 002 Aug 24, 2022

MYLAN

EQ 20MG BASE;375MG

A204920 001 Jul 20, 2021

EQ 20MG BASE;500MG

A204920 002 Jul 20, 2021

VIMOVO

+ HORIZON

EQ 20MG BASE;375MG **

N022511 002 Apr 30, 2010

+

EQ 20MG BASE;500MG **

N022511 001 Apr 30, 2010

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

EUGIA PHARMA

EQ 20MG BASE/VIAL

A204657 001 Aug 10, 2016

MYLAN

EQ 20MG BASE/VIAL

A202686 001 May 17, 2017

EQ 40MG BASE/VIAL

A202686 002 May 17, 2017

SUN PHARM

EQ 20MG BASE/VIAL

A200882 001 Mar 18, 2013

NEXIUM IV

+ ASTRAZENECA

EQ 20MG BASE/VIAL **

N021689 001 Mar 31, 2005

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE STRONTIUM

+ BELCHER

24.65MG

N202342 001 Aug 06, 2013

+

49.3MG

N202342 002 Aug 06, 2013

ESTAZOLAM

TABLET;ORAL

PROSOM

+ ABBOTT

1MG **

N019080 001 Dec 26, 1990

+

2MG **

N019080 002 Dec 26, 1990

ESTRADIOL

FILM, EXTENDED RELEASE;TRANSDERMAL

ALORA

ABBVIE

0.025MG/24HR

N020655 004 Apr 05, 2002

0.05MG/24HR

N020655 001 Dec 20, 1996

0.075MG/24HR

N020655 002 Dec 20, 1996

0.1MG/24HR

N020655 003 Dec 20, 1996

ESTRADIOL

ZYDUS PHARMS

0.025MG/24HR

A202985 001 Mar 29, 2023

0.0375MG/24HR

A202985 002 Mar 29, 2023

0.05MG/24HR

A202985 003 Mar 29, 2023

0.06MG/24HR

A202985 004 Mar 29, 2023

0.075MG/24HR

A202985 005 Mar 29, 2023

0.1MG/24HR

A202985 006 Mar 29, 2023

FEMPATCH

PARKE DAVIS

0.025MG/24HR

N020417 001 Dec 03, 1996

GEL;TOPICAL

ESTROGEL

ASCEND THERAPS US

0.06%

N021166 001 Feb 09, 2004

SYSTEM;TRANSDERMAL

ESCLIM

WOMEN FIRST HLTHCARE

0.025MG/24HR

N020847 001 Aug 04, 1998

0.0375MG/24HR

N020847 002 Aug 04, 1998

0.05MG/24HR

N020847 003 Aug 04, 1998

0.075MG/24HR

N020847 004 Aug 04, 1998

0.1MG/24HR

N020847 005 Aug 04, 1998

ESTRADERM

+ NOVARTIS

0.05MG/24HR

N019081 002 Sep 10, 1986

+

0.1MG/24HR

N019081 003 Sep 10, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRADIOL

SYSTEM; TRANSDERMAL

ESTRADIOL

ORTHO MCNEIL PHARM	0.05MG/24HR	N021048 001	Sep 20, 1999
	0.075MG/24HR	N021048 002	Sep 20, 1999
	0.1MG/24HR	N021048 003	Sep 20, 1999

VIVELLE

SANDOZ	0.025MG/24HR	N020323 005	Aug 16, 2000
	0.0375MG/24HR	N020323 001	Oct 28, 1994
	0.05MG/24HR	N020323 002	Oct 28, 1994
	0.075MG/24HR	N020323 003	Oct 28, 1994
	0.1MG/24HR	N020323 004	Oct 28, 1994

TABLET; ORAL

ESTRACE

BRISTOL MYERS SQUIBB	0.5MG	A081295 001	Jun 30, 1993
+	1MG	A084499 001	
+	2MG	A084500 001	

ESTRADIOL

DR REDDYS LABS SA	0.5MG	A040114 003	Mar 14, 1996
	1MG	A040114 001	Mar 14, 1996
	2MG	A040114 002	Mar 14, 1996
LANNETT CO INC	0.5MG	A040138 001	Jan 30, 1998
	1MG	A040138 002	Jan 30, 1998
	2MG	A040138 003	Jan 30, 1998
MYLAN	0.5MG	A040326 001	Apr 21, 1999
	1MG	A040326 002	Apr 21, 1999
	2MG	A040326 003	Apr 21, 1999
USL PHARMA	0.5MG	A040297 001	Apr 17, 2002
	1MG	A040297 002	Apr 17, 2002
	2MG	A040297 003	Apr 17, 2002

GYNODIOL

DURAMED PHARMS BARR	0.5MG	A040212 001	Dec 29, 1997
	1MG	A040212 002	Dec 29, 1997
	1.5MG	A040212 003	Dec 29, 1997
	2MG	A040212 004	Dec 29, 1997

INNOFEM

NOVO NORDISK INC	0.5MG	A040312 001	Nov 19, 1999
	1MG	A040312 002	Nov 19, 1999
	2MG	A040312 003	Nov 19, 1999

TABLET; VAGINAL

VAGIFEM

+	NOVO NORDISK INC	25MCG **	N020908 001	Mar 26, 1999
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ESTRADIOL ACETATE

TABLET; ORAL

FEMTRACE

+	APIL	0.45MG	N021633 001	Aug 20, 2004
+		0.9MG	N021633 002	Aug 20, 2004
+		1.8MG	N021633 003	Aug 20, 2004

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

PFIZER	1MG/ML	A085470 001	
	3MG/ML	A085470 002	

ESTRADIOL CYPIONATE

DR REDDYS	5MG/ML	A085620 001	
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ESTRADIOL CYPIONATE; MEDROXYPROGESTERONE ACETATE

INJECTABLE; INTRAMUSCULAR

LUNELLE

PHARMACIA AND UPJOHN	5MG/0.5ML; 25MG/0.5ML	N020874 001	Oct 05, 2000
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ESTRADIOL CYPIONATE; TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTADIOL

PHARMACIA AND UPJOHN	2MG/ML; 50MG/ML	N017968 001	
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TESTOSTERONE CYPIONATE-ESTRADIOL CYPIONATE

WATSON LABS	2MG/ML; 50MG/ML	A085603 001	Mar 13, 1986
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRADIOL HEMIHYDRATE

EMULSION; TOPICAL

ESTRASORB

+ EXELTIS USA INC 0.25% N021371 001 Oct 09, 2003

ESTRADIOL VALERATE

INJECTABLE; INJECTION

ESTRADIOL VALERATE

DR REDDYS	20MG/ML	A083547	001	
	40MG/ML	A083714	001	
FOSUN PHARMA	10MG/ML	A040628	001	Oct 04, 2007
	20MG/ML	A040628	002	Oct 04, 2007
	40MG/ML	A040628	003	Oct 04, 2007
WATSON LABS	10MG/ML	A083546	001	

ESTRADIOL VALERATE; TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DITATE-DS

SAVAGE LABS 8MG/ML; 180MG/ML A086423 001

TESTOSTERONE ENANTHATE AND ESTRADIOL VALERATE

WATSON LABS	4MG/ML; 90MG/ML	A085865	001	
	8MG/ML; 180MG/ML	A085860	001	

ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

ACTIVELLA

+ AMNEAL 0.5MG; 0.1MG ** N020907 002 Dec 28, 2006

ESTRADIOL AND NORETHINDRONE ACETATE

NAARI PTE LTD 1MG; 0.5MG A210233 001 Feb 28, 2018

TEVA PHARMS USA 0.5MG; 0.1MG A200747 001 Mar 08, 2012

ETYQA

AUROBINDO PHARMA LTD	0.5MG; 0.1MG	A214729	001	Jun 30, 2023
	1MG; 0.5MG	A214729	002	Jun 30, 2023

ESTRADIOL; NORGESTIMATE

TABLET; ORAL

PREFEST

+ TEVA WOMENS 1MG, 1MG; N/A, 0.09MG ** N021040 001 Oct 22, 1999

ESTRADIOL; PROGESTERONE

CAPSULE; ORAL

ESTRADIOL AND PROGESTERONE

AMNEAL PHARMS 1MG; 100MG A214293 001 May 16, 2022

ESTROGENS, CONJUGATED

TABLET; ORAL

PREMARIN

WYETH PHARMS 2.5MG N004782 002

ESTROGENS, CONJUGATED SYNTHETIC A

CREAM; VAGINAL

SYNTHETIC CONJUGATED ESTROGENS A

TEVA WOMENS 0.625MG/GM N021788 001 Nov 28, 2008

TABLET; ORAL

CENESTIN

+ ASPEN	0.3MG **	N020992	001	Jun 21, 2002
+	0.45MG **	N020992	005	Feb 05, 2004
+	0.625MG **	N020992	002	Mar 24, 1999
+	0.9MG **	N020992	003	Mar 24, 1999
+	1.25MG **	N020992	004	Mar 13, 2000

ESTROGENS, CONJUGATED SYNTHETIC B

TABLET; ORAL

ENJUVIA

ASPEN	0.3MG	N021443	001	Dec 20, 2004
	0.45MG	N021443	002	Dec 20, 2004
	0.625MG **	N021443	003	May 10, 2004
	0.9MG	N021443	005	Apr 27, 2007
	1.25MG **	N021443	004	May 10, 2004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPHASE (PREMARIN; CYCRIN 14/14)

WYETH PHARMS INC 0.625MG, 0.625MG; N/A, 5MG

N020303 002 Dec 30, 1994

PREMPRO (PREMARIN; CYCRIN)

WYETH PHARMS INC 0.625MG, 0.625MG; 2.5MG, 2.5MG

N020303 001 Dec 30, 1994

ESTROGENS, CONJUGATED; MEPROBAMATE

TABLET; ORAL

MILPREM-200

MEDPOINTE PHARM HLC 0.45MG; 200MG

N011045 002

MILPREM-400

MEDPOINTE PHARM HLC 0.45MG; 400MG

N011045 001

PMB 200

WYETH AYERST 0.45MG; 200MG

N010971 005

PMB 400

WYETH AYERST 0.45MG; 400MG

N010971 003

ESTROGENS, ESTERIFIED

TABLET; ORAL

AMNESTROGEN

BRISTOL MYERS SQUIBB 0.3MG

A083266 001

0.625MG

A083266 002

1.25MG

A083266 003

2.5MG

A083266 004

ESTERIFIED ESTROGENS

PVT FORM 0.625MG

A083414 001

1.25MG

A083765 001

2.5MG

A085907 001

SANDOZ 1.25MG

A085302 001

ESTRATAB

SOLVAY 0.3MG

A086715 001

0.625MG

A083209 001

1.25MG

A083856 001

2.5MG

A083857 001

EVEX

ROCHE PALO 0.625MG

A084215 001

1.25MG

A083376 002

FEMOGEN

PVT FORM 0.625MG

A085076 001

1.25MG

A085008 001

2.5MG

A085007 001

ESTRONE

INJECTABLE; INJECTION

ESTROGENIC SUBSTANCE

WYETH AYERST 2MG/ML

A083488 001

ESTRONE

DR REDDYS 5MG/ML

A085239 001

WATSON LABS 2MG/ML

A083397 001

NATURAL ESTROGENIC SUBSTANCE-ESTRONE

WATSON LABS 2MG/ML

A085237 001 Nov 23, 1982

THEELIN

PARKEDALE 1MG/ML

N003977 001

2MG/ML

N003977 002

5MG/ML

N003977 003

ESTROPIPATE

CREAM; VAGINAL

OGEN

PHARMACIA AND UPJOHN 1.5MG/GM

A084710 001

TABLET; ORAL

ESTROPIPATE

BARR 0.75MG

A040135 001 Nov 27, 1996

1.5MG

A040135 002 Nov 27, 1996

3MG

A040135 003 Nov 27, 1996

DURAMED PHARMS BARR 0.75MG

A040296 001 Nov 01, 1999

1.5MG

A040296 002 Nov 01, 1999

3MG

A040296 003 Nov 01, 1999

MYLAN 0.75MG

A040359 001 Aug 26, 1999

1.5MG

A040359 002 Aug 26, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTROPIPATE

TABLET; ORAL

ESTROPIPATE

	3MG	A040359 003	Aug 26, 1999
WATSON LABS	0.75MG	A081213 001	Sep 23, 1993
	1.5MG	A081214 001	Sep 23, 1993
	6MG	A081216 001	Sep 23, 1993
WATSON LABS TEVA	3MG	A081215 001	Sep 23, 1993
OGEN .625			
+ PFIZER	0.75MG	A083220 001	
OGEN 1.25			
+ PFIZER	1.5MG	A083220 002	
OGEN 2.5			
+ PFIZER	3MG	A083220 003	
ORTHO-EST			
SUN PHARM INDS INC	0.75MG	A089567 001	Feb 27, 1991
	1.5MG	A089582 001	Jul 17, 1991

ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

CHARTWELL RX	1MG	A091165 001	Jul 14, 2011
	2MG	A091165 002	Jul 14, 2011
	3MG	A091165 003	Jul 14, 2011
HIKMA	1MG	A091153 001	Apr 15, 2014
	2MG	A091153 002	Apr 15, 2014
	3MG	A091153 003	Apr 15, 2014
NOSTRUM LABS INC	1MG	A203087 001	May 08, 2019
	2MG	A203087 002	May 08, 2019
	3MG	A203087 003	May 08, 2019

ETHACRYNIC ACID

TABLET; ORAL

EDECIN

BAUSCH	50MG	N016092 002	
ETHACRYNIC ACID			
ALVOGEN	25MG	A205709 001	Jul 24, 2018
HIKMA	25MG	A207262 001	Feb 23, 2017

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

ETHAMBUTOL HYDROCHLORIDE

BARR	400MG	A076057 001	Nov 26, 2001
MYAMBUTOL			
STI PHARMA LLC	200MG	N016320 002	
	500MG	N016320 004	

ETHCHLORVYNOL

CAPSULE; ORAL

ETHCHLORVYNOL

BANNER PHARMACAPS	100MG	A084463 001	
	200MG	A084463 002	
	500MG	A084463 003	
	750MG	A084463 004	
PLACIDYL			
ABBVIE	100MG	N010021 004	
	200MG	N010021 007	
	500MG	N010021 002	
	750MG	N010021 010	

ETHINAMATE

CAPSULE; ORAL

VALMID

DISTA	500MG	N009750 001	
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ETHINYL ESTRADIOL

TABLET; ORAL

ESTINYL

SCHERING	0.02MG	N005292 001	
	0.05MG	N005292 002	
	0.5MG	N005292 003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL

TABLET;ORAL

FEMINONE

PHARMACIA AND UPJOHN 0.05MG N016649 001

LYNORAL

ORGANON USA INC 0.01MG N005490 003

0.05MG N005490 002

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET;ORAL-21

DEMULEN 1/35-21

GD SEARLE LLC 0.035MG;1MG ** N018168 001

DEMULEN 1/50-21

GD SEARLE LLC 0.05MG;1MG N016927 001

ZOVIA 1/35E-21

WATSON PHARMS TEVA 0.035MG;1MG A072720 001 Dec 30, 1991

ZOVIA 1/50E-21

WATSON LABS 0.05MG;1MG A072722 001 Dec 30, 1991

TABLET;ORAL-28

DEMULEN 1/35-28

GD SEARLE LLC 0.035MG;1MG ** N018160 001

DEMULEN 1/50-28

GD SEARLE LLC 0.05MG;1MG ** N016936 001

ZOVIA 1/35E-28

DR REDDYS LABS SA 0.035MG;1MG A072721 001 Dec 30, 1991

ETHINYL ESTRADIOL; ETONOGESTREL

RING;VAGINAL

VERARING

DR REDDYS LABS SA 0.015MG/24HR;0.12MG/24HR A207577 001 Dec 09, 2021

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE

TABLET;ORAL-28

NORQUEST FE

PFIZER 0.035MG;75MG;1MG N018926 001 Jul 18, 1986

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE ACETATE

TABLET;ORAL-28

NORLESTRIN FE 1/50

PARKE DAVIS 0.05MG;75MG;1MG N016766 001

NORLESTRIN FE 2.5/50

PARKE DAVIS 0.05MG;75MG;2.5MG N016854 001

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL

FAYOSIM

LUPIN LTD 0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A A205943 001 Mar 29, 2016

LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL

DR REDDYS LABS SA 0.02MG,0.1MG;0.01MG,N/A A200407 001 Oct 25, 2011

LYBREL

+ WYETH PHARMS INC 0.02MG;0.09MG ** N021864 001 May 22, 2007

PREVEN EMERGENCY CONTRACEPTIVE KIT

TEVA BRANDED PHARM 0.05MG;0.25MG N020946 001 Sep 01, 1998

SYLEVIA

SUN PHARM 0.03MG;0.15MG A202988 001 Feb 06, 2019

TABLET;ORAL-21

ALESSE

+ CADENCE HEALTH 0.02MG;0.1MG ** N020683 001 Mar 27, 1997

AVIANE-21

DURAMED PHARMS BARR 0.02MG;0.1MG A075796 002 Apr 30, 2001

ENPRESSE-21

DURAMED PHARMS BARR 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A075809 001 Jul 16, 2001

LESSINA-21

BARR 0.02MG;0.1MG A075803 001 Mar 20, 2002

LEVLITE

+ BAYER HLTHCARE 0.02MG;0.1MG ** N020860 001 Jul 13, 1998

LEVONORGESTREL AND ETHINYL ESTRADIOL

BARR 0.02MG;0.1MG A075862 001 Apr 29, 2003

LEVORA 0.15/30-21

WATSON LABS 0.03MG;0.15MG A073592 001 Dec 13, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL-21

NORDETTE-21

TEVA BRANDED PHARM 0.03MG;0.15MG N018668 001 May 10, 1982

PORTIA-21

BARR 0.03MG;0.15MG A075866 001 May 23, 2002

TRIPHASIL-21

+ WYETH PHARMS 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG ** N019192 001 Nov 01, 1984

TRIVORA-21

DR REDDYS LABS SA 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A074538 001 Dec 18, 1997

TABLET;ORAL-28

ALESSE

+ CADENCE HEALTH 0.02MG;0.1MG ** N020683 002 Mar 27, 1997

CERINTA

SUN PHARM 0.02MG;0.1MG A202817 001 Jan 07, 2019

ELIFEMME

XIROMED 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A202507 001 Dec 04, 2015

LEVLITE

+ BAYER HLTHCARE 0.02MG;0.1MG ** N020860 002 Jul 13, 1998

LEVONORGESTREL AND ETHINYL ESTRADIOL

AMNEAL PHARMS 0.02MG;0.1MG A201108 001 Feb 05, 2014

0.03MG;0.15MG A201095 001 Dec 08, 2014

BARR 0.02MG;0.1MG A075862 002 Apr 29, 2003

MYLAN LABS LTD 0.02MG;0.1MG A202247 001 Dec 08, 2014

0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A202970 001 Mar 23, 2018

NORDETTE-28

+ TEVA BRANDED PHARM 0.03MG;0.15MG ** N018782 001 Jul 21, 1982

ORSYTHIA

VINTAGE PHARMS LLC 0.02MG;0.1MG A077099 001 May 11, 2011

TRIPHASIL-28

+ WYETH PHARMS INC 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG ** N019190 001 Nov 01, 1984

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE;TRANSDERMAL

ORTHO EVRA

+ JANSSEN PHARMS 0.035MG/24HR;0.15MG/24HR ** N021180 001 Nov 20, 2001

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET;ORAL

FEMCON FE

+ APIL 0.035MG;0.4MG ** N021490 001 Nov 14, 2003

TABLET;ORAL-21

BALZIVA-21

BARR 0.035MG;0.4MG A076198 001 Apr 22, 2004

BREVICON 21-DAY

ALLERGAN 0.035MG;0.5MG N017566 001

GENCEPT 10/11-21

BARR 0.035MG,0.035MG;0.5MG,1MG A072694 001 Feb 28, 1992

MODICON 21

ORTHO MCNEIL PHARM 0.035MG;0.5MG ** N017488 001

N.E.E. 1/35 21

LPI 0.035MG;1MG A071541 001 Dec 14, 1987

NORCEPT-E 1/35 21

ORTHO MCNEIL PHARM 0.035MG;1MG A071545 001 Feb 09, 1989

NORETHIN 1/35E-21

WATSON PHARMS TEVA 0.035MG;1MG A071480 001 Apr 12, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL

WATSON LABS 0.035MG;0.4MG A078379 001 Feb 23, 2010

0.035MG;0.5MG A070684 001 Jan 29, 1987

WATSON PHARMS TEVA 0.035MG;1MG A070685 001 Jan 29, 1987

NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)

WATSON LABS 0.035MG,0.035MG;0.5MG,1MG A071043 001 Apr 01, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)

WATSON LABS TEVA 0.035MG,0.035MG;0.5MG,1MG A071041 001 Sep 24, 1991

NORINYL 1+35 21-DAY

ALLERGAN 0.035MG;1MG N017565 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORTREL 0.5/35-21					
BARR	0.035MG;0.5MG		A072692	001	Feb 28, 1992
ORTHO-NOVUM 1/35-21					
ORTHO MCNEIL PHARM	0.035MG;1MG **		N017489	002	
ORTHO-NOVUM 10/11-21					
+ ORTHO MCNEIL JANSSEN	0.035MG,0.035MG;0.5MG,1MG **		N018354	001	Jan 11, 1982
ORTHO-NOVUM 7/14-21					
ORTHO MCNEIL PHARM	0.035MG,0.035MG;0.5MG,1MG **		N019004	001	Apr 04, 1984
ORTHO-NOVUM 7/7/7-21					
JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG		N018985	001	Apr 04, 1984
OVCON-35					
+ WARNER CHILCOTT	0.035MG;0.4MG **		N018127	001	
OVCON-50					
WARNER CHILCOTT	0.05MG;1MG		N018128	001	
TRI-NORINYL 21-DAY					
DR REDDYS LABS SA	0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG		N018977	001	Apr 13, 1984
TABLET; ORAL-28					
BREVICON 28-DAY					
ALLERGAN	0.035MG;0.5MG		N017743	001	
CYCLAFEM 0.5/35					
VINTAGE PHARMS	0.035MG;0.5MG		A203413	001	Dec 16, 2015
CYCLAFEM 1/35					
VINTAGE PHARMS LLC	0.035MG;1MG		A076337	001	Nov 12, 2010
CYCLAFEM 7/7/7					
VINTAGE PHARMS LLC	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG		A076338	001	Nov 16, 2010
GENCEPT 10/11-28					
BARR	0.035MG,0.035MG;0.5MG,1MG		A072697	001	Feb 28, 1992
MODICON 28					
+ JANSSEN PHARMS	0.035MG;0.5MG		N017735	001	
N.E.E. 1/35 28					
LPI	0.035MG;1MG		A071542	001	Dec 14, 1987
NORCEPT-E 1/35 28					
ORTHO MCNEIL PHARM	0.035MG;1MG		A071546	001	Feb 09, 1989
NORETHIN 1/35E-28					
WATSON LABS	0.035MG;1MG		A071481	001	Apr 12, 1988
NORETHINDRONE AND ETHINYL ESTRADIOL					
DR REDDYS LABS SA	0.035MG;0.5MG		A070686	001	Jan 29, 1987
MYLAN LABS LTD	0.035MG;0.4MG		A200897	001	May 11, 2015
	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG		A200486	001	Dec 28, 2015
	0.035MG;0.5MG		A200488	001	Oct 21, 2015
	0.035MG;1MG		A200489	001	Oct 21, 2015
	0.05MG;1MG		A203006	001	Aug 05, 2013
WATSON LABS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG		A076393	001	Feb 04, 2010
NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)					
WATSON LABS	0.035MG,0.035MG;0.5MG,1MG		A071042	001	Sep 24, 1991
NORINYL 1+35 28-DAY					
ALLERGAN	0.035MG;1MG		N017565	002	
ORTHO-NOVUM 1/35-28					
+ JANSSEN PHARMS	0.035MG;1MG **		N017919	002	
ORTHO-NOVUM 10/11-28					
+ ORTHO MCNEIL JANSSEN	0.035MG,0.035MG;0.5MG,1MG		N018354	002	Jan 11, 1982
ORTHO-NOVUM 7/14-28					
ORTHO MCNEIL PHARM	0.035MG,0.035MG;0.5MG,1MG **		N019004	002	Apr 04, 1984
ORTHO-NOVUM 7/7/7-28					
+ JANSSEN PHARMS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG		N018985	002	Apr 04, 1984
OVCON-35					
+ WARNER CHILCOTT LLC	0.035MG;0.4MG **		N017716	001	
OVCON-50					
WARNER CHILCOTT LLC	0.05MG;1MG **		N017576	001	
PIRMELLA 1/35					
LUPIN LTD	0.035MG;1MG		A201512	001	Apr 24, 2013
PIRMELLA 7/7/7					
LUPIN LTD	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,		A201510	001	Apr 24, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONETABLET;ORAL-28
PIRMELLA 7/7/7

1MG

RHUZDAH

AUROBINDO PHARMA 0.035MG;0.4MG

A207585 001 Oct 11, 2022

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

CAPSULE;ORAL

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE
DR REDDYS LABS SA 0.02MG;1MG

A213901 001 Apr 28, 2022

TABLET;ORAL

FEMHRT

+ APIL 0.0025MG;0.5MG
+ 0.005MG;1MG **N021065 001 Jan 14, 2005
N021065 002 Oct 15, 1999

LOESTRIN 24 FE

+ TEVA BRANDED PHARM 0.02MG;1MG **

N021871 001 Feb 17, 2006

MINASTRIN 24 FE

+ APIL 0.02MG;1MG

N203667 001 May 08, 2013

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

MYLAN LABS LTD 0.0025MG;0.5MG
0.005MG;1MGA207260 001 Feb 02, 2017
A207259 001 Dec 27, 2016

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

MYLAN LABS LTD 0.01MG,0.01MG;1MG,N/A

A205049 001 May 31, 2016

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AMNEAL PHARMS 0.02MG;1MG

A078267 001 Sep 01, 2009

0.02MG;1MG

A207514 001 Sep 11, 2017

APOTEX 0.02MG;1MG

A208639 001 Mar 21, 2018

MYLAN LABS LTD 0.02MG;1MG

A202742 001 Oct 30, 2014

0.02MG;1MG

A206120 001 Sep 12, 2017

TABLET;ORAL-21

ESTROSTEP 21

+ APIL 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG **

N020130 001 Oct 09, 1996

GILDESS 1.5/30

VINTAGE PHARMS LLC 0.03MG;1.5MG

A077075 002 Jul 24, 2012

GILDESS 1/20

VINTAGE PHARMS LLC 0.02MG;1MG

A077077 002 Jul 24, 2012

LOESTRIN 21 1.5/30

+ TEVA BRANDED PHARM 0.03MG;1.5MG

N017875 001

LOESTRIN 21 1/20

+ TEVA BRANDED PHARM 0.02MG;1MG **

N017876 001

LOESTRIN FE 1.5/30

+ TEVA BRANDED PHARM 0.03MG;1.5MG

N017355 001

MICROGESTIN 1.5/30

DR REDDYS LABS SA 0.03MG;1.5MG

A075548 002 Jul 30, 2003

MICROGESTIN 1/20

DR REDDYS LABS SA 0.02MG;1MG

A075647 002 Jul 30, 2003

NORLESTRIN 21 1/50

PARKE DAVIS 0.05MG;1MG

N016749 001

NORLESTRIN 21 2.5/50

PARKE DAVIS 0.05MG;2.5MG

N016852 001

TRI-LEGEST 21

BARR 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG

A076405 001 Oct 26, 2007

TABLET;ORAL-28

ESTROSTEP FE

+ APIL 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG **

N020130 002 Oct 09, 1996

GILDESS FE 1.5/30

VINTAGE PHARMS LLC 0.03MG;1.5MG

A077075 001 Apr 28, 2005

GILDESS FE 1/20

VINTAGE PHARMS LLC 0.02MG;1MG

A077077 001 May 20, 2005

LOESTRIN FE 1/20

+ TEVA BRANDED PHARM 0.02MG;1MG

N017354 001

MICROGESTIN FE 1.5/30

DR REDDYS LABS SA 0.03MG;1.5MG

A075548 001 Feb 05, 2001

MICROGESTIN FE 1/20

DR REDDYS LABS SA 0.02MG;1MG

A075647 001 Feb 05, 2001

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

DR REDDYS LABS SA 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG

A076629 001 Mar 18, 2010

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL-28

NORLESTRIN 28 1/50

PARKE DAVIS

0.05MG;1MG

N016723 001

TABLET, CHEWABLE, TABLET;ORAL

LO MINASTRIN FE

+

APIL

0.01MG,0.01MG,N/A;1MG,N/A,N/A

N204654 001 Jul 24, 2013

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET;ORAL-21

ORTHO CYCLEN-21

JANSSEN PHARMS

0.035MG;0.25MG **

N019653 001 Dec 29, 1989

ORTHO TRI-CYCLEN

JANSSEN PHARMS

0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,
0.25MG **

N019697 002 Jul 03, 1992

TABLET;ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL

AMNEAL PHARMS

0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,
0.25MG

A203870 001 Nov 12, 2015

0.035MG;0.25MG

A203865 001 Oct 27, 2015

MYLAN

0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,
0.25MG

A201897 001 Jan 27, 2016

0.035MG;0.25MG

A201896 001 Jan 27, 2016

MYLAN LABS LTD

0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,
0.25MG

A202132 001 Sep 09, 2015

WATSON LABS

0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,
0.25MG

A090479 001 Mar 09, 2011

0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,
0.25MG

A076626 001 Aug 17, 2006

0.035MG;0.25MG

A076627 001 Aug 17, 2006

ORTHO CYCLEN-28

+

JANSSEN PHARMS

0.035MG;0.25MG **

N019653 002 Dec 29, 1989

ORTHO TRI-CYCLEN

+

JANSSEN PHARMS

0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,
0.25MG **

N019697 001 Jul 03, 1992

ORTHO TRI-CYCLEN LO

+

JANSSEN PHARMS

0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,
0.25MG **

N021241 001 Aug 22, 2002

TRI-PREVIFEM

VINTAGE PHARMS LLC

0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,
0.25MG

A076335 001 Mar 26, 2004

ETHINYL ESTRADIOL; NORGESTREL

TABLET;ORAL-21

LO/OVRAL

+

CADENCE HEALTH

0.03MG;0.3MG

N017612 001

LOW-OGESTREL-21

DR REDDYS LABS SA

0.03MG;0.3MG

A075288 001 Jul 28, 1999

OGESTREL 0.5/50-21

WATSON LABS

0.05MG;0.5MG **

A075406 001 Dec 15, 1999

OVRAL

WYETH PHARMS

0.05MG;0.5MG

N016672 001

TABLET;ORAL-28

LO/OVRAL-28

+

WYETH PHARMS

0.03MG;0.3MG **

N017802 001

NORGESTREL AND ETHINYL ESTRADIOL

MYLAN LABS LTD

0.03MG;0.3MG

A201828 001 Jun 21, 2016

0.05MG;0.5MG

A202875 001 May 08, 2017

OGESTREL 0.5/50-28

WATSON LABS

0.05MG;0.5MG

A075406 002 Dec 15, 1999

OVRAL-28

WYETH PHARMS

0.05MG;0.5MG

N016806 001

ETHOPROPAZINE HYDROCHLORIDE

TABLET;ORAL

PARSIDOL

PARKE DAVIS

10MG

N009078 003

50MG

N009078 006

100MG

N009078 008

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHOSUXIMIDE

SYRUP; ORAL

ETHOSUXIMIDE

TEVA PHARMS

250MG/5ML

A081306 001 Jul 30, 1993

ETHOTOIN

TABLET; ORAL

PEGANONE

+ RECORDATI RARE

250MG

N010841 001

500MG

N010841 003

ETHOXZOLAMIDE

TABLET; ORAL

CARDRASE

PHARMACIA AND UPJOHN

62.5MG

N011047 002

125MG

N011047 001

ETHAMIDE

ALLERGAN

125MG

N016144 001

ETHYLESTRENOL

ELIXIR; ORAL

MAXIBOLIN

ORGANON USA INC

2MG/5ML

N014006 002

TABLET; ORAL

MAXIBOLIN

ORGANON USA INC

2MG

N014005 002

ETHYNODIOL DIACETATE; MESTRANOL

TABLET; ORAL-20

OVULEN

GD SEARLE LLC

1MG; 0.1MG

N016029 002

TABLET; ORAL-21

OVULEN-21

GD SEARLE LLC

1MG; 0.1MG

N016029 003

TABLET; ORAL-28

OVULEN-28

GD SEARLE LLC

1MG; 0.1MG

N016705 001

ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA

0.5% **

N017751 003

+

1% **

N017751 005

ETIDRONATE DISODIUM

INJECTABLE; INJECTION

DIDRONEL

MGI PHARMA INC

50MG/ML

N019545 001 Apr 20, 1987

TABLET; ORAL

DIDRONEL

+ APIL

200MG **

N017831 001

+

400MG **

N017831 002

ETIDRONATE DISODIUM

MYLAN

200MG

A075800 001 Jan 24, 2003

400MG

A075800 002 Jan 24, 2003

ETODOLAC

CAPSULE; ORAL

ETODOLAC

ANI PHARMS

200MG

A074840 001 Aug 29, 1997

200MG

A074844 001 Dec 23, 1997

200MG

A074899 001 Jul 08, 1997

300MG

A074840 002 Aug 29, 1997

300MG

A074844 002 Dec 23, 1997

300MG

A074899 002 Jul 08, 1997

BIOPHARM

300MG

A074929 001 Jan 30, 1998

CHARTWELL MOLECULES

200MG

A074842 001 Jul 17, 1997

300MG

A074842 002 Jul 17, 1997

MYLAN

200MG

A074932 001 May 16, 1997

200MG

A075071 001 Sep 30, 1998

300MG

A074932 002 May 16, 1997

300MG

A075071 002 Sep 30, 1998

SANDOZ

200MG

A074942 001 Sep 30, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETODOLAC

CAPSULE; ORAL

ETODOLAC

300MG

A074942 002 Sep 30, 1997

LODINE

+ WYETH PHARMS INC

200MG **

N018922 002 Jan 31, 1991

+

300MG

N018922 003 Jan 31, 1991

TABLET; ORAL

ETODOLAC

BIOPHARM

400MG

A074927 001 Oct 30, 1997

CHARTWELL MOLECULES

400MG

A074841 001 Jun 27, 1997

EDENBRIDGE PHARMS

400MG

A209888 001 Nov 30, 2018

500MG

A209888 002 Nov 30, 2018

IVAX SUB TEVA PHARMS

400MG

A074883 001 Feb 28, 1997

500MG

A074883 002 Nov 20, 1998

MYLAN

400MG

A075012 001 Sep 30, 1998

400MG

A075104 001 Feb 06, 1998

500MG

A075012 002 Sep 30, 1998

500MG

A075104 002 Nov 20, 1998

OXFORD PHARMS

400MG

A074819 001 Feb 28, 1997

500MG

A074819 002 Apr 28, 1998

RANBAXY LABS LTD

400MG

A075226 001 Nov 24, 1998

500MG

A075226 002 Nov 24, 1998

SHREE HARI INTL

400MG

A074839 001 Jul 11, 1997

400MG

A074846 001 Feb 28, 1997

TEVA

400MG

A074847 001 Apr 23, 1999

400MG

A075009 001 Nov 26, 1997

500MG

A074847 002 Apr 23, 1999

500MG

A075009 002 Dec 28, 1999

WATSON LABS

400MG

A074892 001 Apr 16, 1997

400MG

A075069 001 Apr 16, 1998

500MG

A074892 002 Oct 29, 1998

LODINE

+ WYETH PHARMS INC

400MG **

N018922 004 Jul 29, 1993

+

500MG **

N018922 005 Jun 28, 1996

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

ACTAVIS ELIZABETH

400MG

A075696 001 Jul 31, 2000

ANI PHARMS

400MG

A075943 001 Jul 26, 2002

500MG

A075943 002 Jul 26, 2002

600MG

A075943 003 Jul 26, 2002

WATSON LABS FLORIDA

400MG

A075829 001 Nov 30, 2001

500MG

A075829 002 Nov 30, 2001

LODINE XL

WYETH PHARMS INC

400MG **

N020584 001 Oct 25, 1996

500MG **

N020584 003 Jan 20, 1998

+

600MG **

N020584 002 Oct 25, 1996

ETOMIDATE

INJECTABLE; INJECTION

ETOMIDATE

AVET LIFESCIENCES

2MG/ML

A204618 001 Aug 13, 2014

LUITPOLD

2MG/ML

A078867 001 Dec 22, 2009

PAR STERILE PRODUCTS

2MG/ML

A091297 001 Jun 20, 2012

RISING

2MG/ML

A078289 001 Jan 02, 2009

ETONOGESTREL

IMPLANT; IMPLANTATION

IMPLANON

ORGANON

68MG/IMPLANT

N021529 001 Jul 17, 2006

ETOPOSIDE

CAPSULE; ORAL

VEPESID

+ STRIDES PHARMA

50MG

N019557 001 Dec 30, 1986

+

100MG

N019557 002 Dec 30, 1986

INJECTABLE; INJECTION

ETOPOSIDE

DASH PHARMS

20MG/ML

A204927 001 Oct 31, 2017

DASH PHARMS NATCO

20MG/ML

A203507 001 Nov 20, 2017

HOSPIRA

20MG/ML

A074320 001 Aug 30, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETOPOSIDE

INJECTABLE; INJECTION

ETOPOSIDE

	20MG/ML	A074351 001	Aug 30, 1995
PHARMACHEMIE BV	20MG/ML	A074227 001	Feb 22, 1996
PIERRE FABRE	20MG/ML	A074813 001	Jul 09, 1997
TEVA PARENTERAL	20MG/ML	A074510 001	Jun 29, 1995
TEVA PHARMS USA	20MG/ML	A074284 001	Feb 10, 1994
WATSON LABS	20MG/ML	A074228 001	Oct 15, 1996
WATSON LABS INC	20MG/ML	A074968 001	Jan 09, 1998
TOPOSAR			
TEVA PARENTERAL	20MG/ML	A074166 001	Feb 27, 1995
VEPESID			
+ CORDEN PHARMA	20MG/ML **	N018768 001	Nov 10, 1983

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL	N020906 001	Feb 27, 1998
	EQ 1GM BASE/VIAL	N020906 002	Feb 27, 1998

ETRAVIRINE

TABLET; ORAL

ETRAVIRINE

AMNEAL	25MG	A214196 001	Jun 14, 2021
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ETRETINATE

CAPSULE; ORAL

TEGISON

ROCHE	10MG	N019369 001	Sep 30, 1986
	25MG	N019369 002	Sep 30, 1986

EVANS BLUE

INJECTABLE; INJECTION

EVANS BLUE

PARKE DAVIS	0.5% **	N008041 001	
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EXEMESTANE

TABLET; ORAL

EXEMESTANE

ALVOGEN	25MG	A200898 001	Jul 28, 2014
AMNEAL PHARMS	25MG	A206421 001	Dec 28, 2018
DR REDDYS LABS SA	25MG	A208764 001	Aug 08, 2019

EXENATIDE SYNTHETIC

FOR SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

BYDUREON

+ ASTRAZENECA AB	2MG/VIAL	N022200 001	Jan 27, 2012
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BYDUREON PEN

+ ASTRAZENECA AB	2MG	N022200 002	Feb 28, 2014
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EZETIMIBE

TABLET; ORAL

EZETIMIBE

APOTEX	10MG	A208332 001	Jun 12, 2017
RISING	10MG	A201790 001	Apr 26, 2019
TEVA PHARMS USA	10MG	A078724 001	Jun 12, 2017

EZETIMIBE; ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSZET

+ ALTHERA PHARMS	10MG;EQ 5MG BASE	N213072 001	Mar 23, 2021
	10MG;EQ 10MG BASE	N213072 002	Mar 23, 2021
	10MG;EQ 20MG BASE	N213072 003	Mar 23, 2021
	10MG;EQ 40MG BASE	N213072 004	Mar 23, 2021

EZETIMIBE; SIMVASTATIN

TABLET; ORAL

EZETIMIBE AND SIMVASTATIN

ANI PHARMS	10MG;10MG	A201890 001	Apr 26, 2017
	10MG;20MG	A201890 002	Apr 26, 2017
	10MG;40MG	A201890 003	Apr 26, 2017
	10MG;80MG	A201890 004	Apr 26, 2017
AUROBINDO PHARMA USA	10MG;10MG	A200082 001	Dec 17, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EZETIMIBE; SIMVASTATIN

TABLET; ORAL

EZETIMIBE AND SIMVASTATIN

10MG; 20MG	A200082	002	Dec 17, 2020
10MG; 40MG	A200082	003	Dec 17, 2020
10MG; 80MG	A200082	004	Dec 17, 2020

EZOGABINE

TABLET; ORAL

POTIGA

+ GLAXOSMITHKLINE	50MG	N022345	001	Jun 10, 2011
+	200MG	N022345	002	Jun 10, 2011
+	300MG	N022345	003	Jun 10, 2011
+	400MG	N022345	004	Jun 10, 2011

FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

CIPLA	125MG	A078278	001	Mar 21, 2011
	250MG	A078278	002	Mar 21, 2011
	500MG	A078278	003	Mar 21, 2011
HIKMA	125MG	A090128	001	Mar 21, 2011
	250MG	A090128	002	Mar 21, 2011
	500MG	A090128	003	Mar 21, 2011
MYLAN	125MG	A201333	001	Mar 24, 2011
	250MG	A201333	002	Mar 24, 2011
	500MG	A201333	003	Mar 24, 2011
FAMVIR				
+ NOVARTIS	125MG **	N020363	003	Dec 11, 1995
+	250MG **	N020363	001	Apr 26, 1996
+	500MG **	N020363	002	Jun 29, 1994

FAMOTIDINE

FOR SUSPENSION; ORAL

PEPCID

+ SALIX PHARMS	40MG/5ML **	N019527	001	Feb 02, 1987
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INJECTABLE; INJECTION

FAMOTIDINE

APOTEX INC	10MG/ML	A075942	001	Aug 02, 2002
APOTHECON	10MG/ML	A075707	001	Apr 16, 2001
HIKMA	10MG/ML	A075799	001	Apr 30, 2002
HOSPIRA	10MG/ML	A075705	001	Apr 16, 2001
	10MG/ML	A075870	001	Nov 23, 2001
	10MG/ML	A075905	001	Nov 23, 2001
ZYDUS PHARMS	10MG/ML	A215828	001	Nov 22, 2022
FAMOTIDINE PRESERVATIVE FREE				
AKORN	10MG/ML	A076324	001	Nov 27, 2002
APOTHECON	10MG/ML	A075708	001	Apr 16, 2001
HIKMA	10MG/ML	A075789	001	Apr 30, 2002
HOSPIRA	10MG/ML	A075669	001	Apr 16, 2001
FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK)				
AKORN	10MG/ML	A076322	001	Nov 27, 2002
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER				
ABBVIE	0.4MG/ML	A075729	001	Dec 17, 2001
PEPCID				
+ MERCK	10MG/ML **	N019510	001	Nov 04, 1986
PEPCID PRESERVATIVE FREE				
+ MERCK	10MG/ML **	N019510	004	Nov 04, 1986
PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER				
+ MERCK SHARP DOHME	0.4MG/ML **	N020249	001	Feb 18, 1994

TABLET; ORAL

FAMOTIDINE

ACTAVIS ELIZABETH	20MG	A075650	001	Sep 14, 2001
	40MG	A075650	002	Sep 14, 2001
APOTEX	10MG	A075610	001	Mar 12, 2002
MYLAN	10MG	A075674	001	Dec 21, 2001
MYLAN PHARMS INC	20MG	A075457	001	Apr 18, 2001
	40MG	A075457	002	Apr 18, 2001
PERRIGO R AND D	20MG	A077352	002	Jul 27, 2005
	40MG	A077352	001	Jul 27, 2005
RISING	20MG	A075704	001	Apr 16, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FAMOTIDINETABLET; ORAL
FAMOTIDINE

	40MG	A075704 002	Apr 16, 2001
SANDOZ	10MG	A076101 001	Oct 21, 2002
	20MG	A075607 001	May 10, 2001
	20MG	A075793 001	Apr 16, 2001
	40MG	A075607 002	May 10, 2001
	40MG	A075793 002	Apr 16, 2001
SUN PHARM INDUSTRIES	20MG	A075639 002	Dec 12, 2001
	40MG	A075639 001	Dec 12, 2001
TEVA	10MG	A075312 001	May 31, 2001
	20MG	A075311 001	Apr 16, 2001
	40MG	A075311 002	Apr 16, 2001
WATSON LABS	10MG	A075404 001	Nov 28, 2001
	20MG	A075062 002	Apr 16, 2001
	40MG	A075062 001	Apr 16, 2001
PEPCID			
+ BAUSCH	20MG **	N019462 001	Oct 15, 1986
+	40MG **	N019462 002	Oct 15, 1986

TABLET, CHEWABLE; ORAL

FAMOTIDINE

PERRIGO	10MG	A075715 001	Aug 22, 2003
PEPCID AC			
+ J AND J CONSUMER INC	10MG **	N020801 001	Sep 24, 1998
TABLET, ORALLY DISINTEGRATING; ORAL			
FLUXID			
UCB INC	20MG	N021712 001	Sep 24, 2004
	40MG	N021712 002	Sep 24, 2004
PEPCID RPD			
MERCK	20MG	N020752 001	May 28, 1998
	40MG	N020752 002	May 28, 1998

FEBUXOSTAT

TABLET; ORAL

FEBUXOSTAT

MYLAN	40MG	A205385 001	Jul 01, 2019
	80MG	A205385 002	Jul 01, 2019
TORRENT	40MG	A211837 001	Dec 19, 2023
	80MG	A211837 002	Dec 19, 2023

FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

JUBILANT GENERICS	2.5MG	A203983 001	Aug 19, 2016
	5MG	A203983 002	Aug 19, 2016
	10MG	A203983 003	Aug 19, 2016
MYLAN	2.5MG	A078855 001	Apr 17, 2008
	5MG	A078855 002	Apr 17, 2008
	10MG	A078855 003	Apr 17, 2008
SUN PHARM INDUSTRIES	2.5MG	A075896 001	Nov 02, 2004
	5MG	A075896 002	Nov 02, 2004
	10MG	A075896 003	Nov 02, 2004
WOCKHARDT	2.5MG	A091484 001	Aug 15, 2012
	5MG	A091484 002	Aug 15, 2012
	10MG	A091484 003	Aug 15, 2012
PLENDIL			
+ ASTRAZENECA	2.5MG **	N019834 004	Sep 22, 1994
+	5MG **	N019834 001	Jul 25, 1991
+	10MG **	N019834 002	Jul 25, 1991

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

+ LUPIN	30MG **	N021695 004	Oct 18, 2013
	87MG	N021695 002	Nov 30, 2004
+	90MG	N021695 005	Oct 18, 2013
FENOFIBRATE (MICRONIZED)			
IMPAX LABS	67MG	A075868 001	Oct 27, 2003
	134MG	A075868 002	Oct 27, 2003
	200MG	A075868 003	Oct 27, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENOFIBRATE

CAPSULE; ORAL

FENOFIBRATE (MICRONIZED)

MYLAN PHARMS INC	43MG	A202579 001	Jan 10, 2013
	130MG	A202579 002	Jan 10, 2013
NOVAST LABS	67MG	A207564 001	Apr 19, 2019
	134MG	A207564 002	Apr 19, 2019
	200MG	A207564 003	Apr 19, 2019
RISING	67MG	A202676 001	Oct 23, 2012
	134MG	A202676 002	Oct 23, 2012
	200MG	A202676 003	Oct 23, 2012

LIPIDIL

ABBVIE	100MG	N019304 001	Dec 31, 1993
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LIPOFEN

CIPHER PHARMS INC	100MG	N021612 002	Jan 11, 2006
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TRICOR (MICRONIZED)

+ ABBVIE	67MG **	N019304 002	Feb 09, 1998
+	134MG **	N019304 003	Jun 30, 1999
+	200MG **	N019304 004	Jun 30, 1999

TABLET; ORAL

FENOFIBRATE

ALEMBIC	54MG	A213252 001	Jan 17, 2020
	160MG	A213252 002	Jan 17, 2020
MYLAN	107MG	A076520 002	Dec 29, 2005

TRICOR

+ ABBVIE	54MG **	N021203 001	Sep 04, 2001
+	160MG **	N021203 003	Sep 04, 2001

TRIGLIDE

SKYEPHARMA AG	50MG	N021350 001	May 07, 2005
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FENOFIBRIC ACID

TABLET; ORAL

FIBRICOR

+ ATHENA	35MG	N022418 001	Aug 14, 2009
+	105MG	N022418 002	Aug 14, 2009

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

FENOLDOPAM MESYLATE

LUITPOLD	EQ 10MG BASE/ML	A076656 001	Dec 01, 2003
TEVA PARENTERAL	EQ 10MG BASE/ML	A077826 001	Mar 07, 2007

FENOPROFEN CALCIUM

CAPSULE; ORAL

FENOPROFEN CALCIUM

AM THERAP	EQ 200MG BASE	A072307 001	Aug 22, 1988
	EQ 300MG BASE	A072308 001	Aug 22, 1988
HALSEY	EQ 200MG BASE	A072355 001	Aug 17, 1988
	EQ 300MG BASE	A072356 001	Aug 17, 1988
PAR PHARM	EQ 200MG BASE	A072437 001	Aug 22, 1988
	EQ 300MG BASE	A072438 001	Aug 22, 1988
QUANTUM PHARMICS	EQ 200MG BASE	A072214 001	Aug 17, 1988
	EQ 300MG BASE	A071738 001	Aug 17, 1988
RISING	EQ 200MG BASE	A072394 001	Oct 17, 1988
	EQ 300MG BASE	A072395 001	Oct 17, 1988
WARNER CHILCOTT	EQ 200MG BASE	A072946 001	Apr 30, 1991
	EQ 300MG BASE	A072472 001	Apr 30, 1991
WATSON LABS	EQ 200MG BASE	A072294 001	Aug 17, 1988
	EQ 200MG BASE	A072981 001	Aug 19, 1991
	EQ 300MG BASE	A072293 001	Aug 17, 1988
	EQ 300MG BASE	A072982 001	Aug 19, 1991

NALFON

XSPIRE PHARMA	EQ 300MG BASE	N017604 002	
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TABLET; ORAL

FENOPROFEN CALCIUM

AM THERAP	EQ 600MG BASE	A072309 001	Aug 17, 1988
ANI PHARMS	EQ 600MG BASE	A072274 001	May 02, 1988
DAVA PHARMS INC	EQ 600MG BASE	A072326 001	Aug 17, 1988
HALSEY	EQ 600MG BASE	A072357 001	Aug 17, 1988
IVAX SUB TEVA PHARMS	EQ 600MG BASE	A072557 001	Aug 29, 1988
QUANTUM PHARMICS	EQ 600MG BASE	A072194 001	Aug 17, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENOPROFEN CALCIUM

TABLET;ORAL

FENOPROFEN CALCIUM

RISING	EQ 600MG BASE	A072396 001	Oct 17, 1988
STRIDES PHARMA	EQ 600MG BASE	A072429 001	Aug 17, 1988
SUN PHARM INDUSTRIES	EQ 600MG BASE	A072902 001	Dec 21, 1990
USL PHARMA	EQ 600MG BASE	A072362 001	Aug 17, 1988
WATSON LABS	EQ 600MG BASE	A072165 001	Aug 17, 1988
	EQ 600MG BASE	A072602 001	Oct 11, 1988
WATSON LABS TEVA	EQ 600MG BASE	A072407 001	Aug 17, 1988
NALFON			
+ DISTA	EQ 600MG BASE	N017710 001	

FENTANYL

FILM, EXTENDED RELEASE;TRANSDERMAL

DURAGESIC-100			
+ JANSSEN PHARMS	100MCG/HR **	N019813 001	Aug 07, 1990
DURAGESIC-12			
+ JANSSEN PHARMS	12.5MCG/HR **	N019813 005	Feb 04, 2005
DURAGESIC-25			
+ JANSSEN PHARMS	25MCG/HR **	N019813 004	Aug 07, 1990
DURAGESIC-37			
+ JANSSEN PHARMS	37.5MCG/HR **	N019813 006	Jan 24, 2018
DURAGESIC-50			
+ JANSSEN PHARMS	50MCG/HR **	N019813 003	Aug 07, 1990
DURAGESIC-75			
+ JANSSEN PHARMS	75MCG/HR **	N019813 002	Aug 07, 1990
FENTANYL-100			
ACTAVIS LABS UT INC	100MCG/HR	A076709 004	Aug 20, 2007
LAVIPHARM LABS	100MCG/HR	A077051 004	Aug 04, 2006
MAYNE PHARMA	100MCG/HR	A077062 004	Aug 20, 2007
NOVEN	100MCG/HR	A077775 004	Oct 16, 2009
ZYDUS PHARMS	100MCG/HR	A209655 008	Jan 24, 2023
FENTANYL-12			
ZYDUS PHARMS	12.5MCG/HR	A209655 001	Jan 24, 2023
FENTANYL-25			
ACTAVIS LABS UT INC	25MCG/HR	A076709 001	Aug 20, 2007
LAVIPHARM LABS	25MCG/HR	A077051 001	Aug 04, 2006
MAYNE PHARMA	25MCG/HR	A077062 001	Aug 20, 2007
NOVEN	25MCG/HR	A077775 001	Oct 16, 2009
ZYDUS PHARMS	25MCG/HR	A209655 002	Jan 24, 2023
FENTANYL-37			
ZYDUS PHARMS	37.5MCG/HR	A209655 003	Jan 24, 2023
FENTANYL-50			
ACTAVIS LABS UT INC	50MCG/HR	A076709 002	Aug 20, 2007
LAVIPHARM LABS	50MCG/HR	A077051 002	Aug 04, 2006
MAYNE PHARMA	50MCG/HR	A077062 002	Aug 20, 2007
NOVEN	50MCG/HR	A077775 002	Oct 16, 2009
ZYDUS PHARMS	50MCG/HR	A209655 004	Jan 24, 2023
FENTANYL-62			
ZYDUS PHARMS	62.5MCG/HR	A209655 005	Jan 24, 2023
FENTANYL-75			
ACTAVIS LABS UT INC	75MCG/HR	A076709 003	Aug 20, 2007
LAVIPHARM LABS	75MCG/HR	A077051 003	Aug 04, 2006
MAYNE PHARMA	75MCG/HR	A077062 003	Aug 20, 2007
NOVEN	75MCG/HR	A077775 003	Oct 16, 2009
ZYDUS PHARMS	75MCG/HR	A209655 006	Jan 24, 2023
FENTANYL-87			
ZYDUS PHARMS	87.5MCG/HR	A209655 007	Jan 24, 2023
SPRAY;SUBLINGUAL			
SUBSYS			
+ BTCP PHARMA	0.1MG **	N202788 001	Jan 04, 2012
	0.2MG **	N202788 002	Jan 04, 2012
	0.4MG **	N202788 003	Jan 04, 2012
	0.6MG **	N202788 004	Jan 04, 2012
	0.8MG **	N202788 005	Jan 04, 2012
	1.2MG **	N202788 006	Aug 30, 2012
	1.6MG **	N202788 007	Aug 30, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENTANYL CITRATEFILM;BUCCAL
ONSOLIS

ADALVO	EQ 0.2MG BASE	N022266 001	Jul 16, 2009
	EQ 0.4MG BASE	N022266 002	Jul 16, 2009
	EQ 0.6MG BASE	N022266 003	Jul 16, 2009
	EQ 0.8MG BASE	N022266 004	Jul 16, 2009
	EQ 1.2MG BASE	N022266 005	Jul 16, 2009

INJECTABLE;INJECTION

FENTANYL CITRATE

ABBOTT	EQ 0.05MG BASE/ML	A070636 001	Apr 30, 1990
	EQ 0.05MG BASE/ML	A070637 001	Apr 30, 1990
WATSON LABS	EQ 0.05MG BASE/ML	A073488 001	Jun 30, 1992

FENTANYL CITRATE PRESERVATIVE FREE

DR REDDYS	EQ 0.05MG BASE/ML	A074917 001	Feb 03, 1998
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SOLUTION;INTRAVENOUS

FENTANYL CITRATE

+ EXELA PHARMA	EQ 2.5MG BASE/50ML (EQ 0.05MG BASE/ML)	N215870 001	Feb 08, 2023
+	EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML)	N215870 002	Feb 08, 2023

SPRAY, METERED;NASAL

LAZANDA

+ BTCP PHARMA	EQ 0.1MG BASE **	N022569 001	Jun 30, 2011
+	EQ 0.3MG BASE **	N022569 003	Dec 21, 2015
+	EQ 0.4MG BASE **	N022569 002	Jun 30, 2011

TABLET;BUCCAL, SUBLINGUAL

FENTANYL CITRATE

DR REDDYS LABS SA	EQ 0.1MG BASE	A206329 001	Aug 22, 2022
	EQ 0.2MG BASE	A206329 002	Aug 22, 2022
	EQ 0.3MG BASE	A206329 003	Aug 22, 2022
	EQ 0.4MG BASE	A206329 004	Aug 22, 2022
	EQ 0.6MG BASE	A206329 005	Aug 22, 2022
	EQ 0.8MG BASE	A206329 006	Aug 22, 2022
WATSON LABS	EQ 0.1MG BASE	A079075 001	Jan 07, 2011
	EQ 0.2MG BASE	A079075 002	Jan 07, 2011
	EQ 0.4MG BASE	A079075 003	Jan 07, 2011
	EQ 0.6MG BASE	A079075 004	Jan 07, 2011
	EQ 0.8MG BASE	A079075 005	Jan 07, 2011

FENTORA

+ CEPHALON	EQ 0.3MG BASE **	N021947 006	Mar 02, 2007
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TABLET;SUBLINGUAL

ABSTRAL

+ SENTYNL THERAPS INC	EQ 0.1MG BASE **	N022510 001	Jan 07, 2011
+	EQ 0.2MG BASE **	N022510 002	Jan 07, 2011
+	EQ 0.3MG BASE **	N022510 003	Jan 07, 2011
+	EQ 0.4MG BASE **	N022510 004	Jan 07, 2011
+	EQ 0.6MG BASE **	N022510 005	Jan 07, 2011
+	EQ 0.8MG BASE **	N022510 006	Jan 07, 2011

FENTANYL CITRATE

ACTAVIS LABS FL INC	EQ 0.1MG BASE	A207338 001	Nov 17, 2017
	EQ 0.2MG BASE	A207338 002	Nov 17, 2017
	EQ 0.3MG BASE	A207338 003	Nov 17, 2017
	EQ 0.4MG BASE	A207338 004	Nov 17, 2017
	EQ 0.6MG BASE	A207338 005	Nov 17, 2017
	EQ 0.8MG BASE	A207338 006	Nov 17, 2017

TROCHE/LOZENGE;ORAL

FENTANYL

CEPHALON	EQ 0.1MG BASE	N020195 007	Oct 30, 1995
	EQ 0.2MG BASE	N020195 001	Oct 04, 1993
	EQ 0.3MG BASE	N020195 002	Oct 04, 1993
	EQ 0.4MG BASE	N020195 003	Oct 04, 1993

TROCHE/LOZENGE;TRANSMUCOSAL

FENTANYL CITRATE

PAR PHARM	EQ 0.2MG BASE	A077312 001	Oct 30, 2009
	EQ 0.4MG BASE	A077312 002	Oct 30, 2009
	EQ 0.6MG BASE	A077312 003	Oct 30, 2009
	EQ 0.8MG BASE	A077312 004	Oct 30, 2009
	EQ 1.2MG BASE	A077312 005	Oct 30, 2009
	EQ 1.6MG BASE	A077312 006	Oct 30, 2009
SPECGX LLC	EQ 0.2MG BASE	A078907 001	Oct 30, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENTANYL CITRATETROCHE/LOZENGE; TRANSMUCOSAL
FENTANYL CITRATE

EQ 0.4MG BASE	A078907 002	Oct 30, 2009
EQ 0.6MG BASE	A078907 003	Oct 30, 2009
EQ 0.8MG BASE	A078907 004	Oct 30, 2009
EQ 1.2MG BASE	A078907 005	Oct 30, 2009
EQ 1.6MG BASE	A078907 006	Oct 30, 2009

FENTANYL HYDROCHLORIDESYSTEM; IONTOPHORESIS, TRANSDERMAL
IONSYS

+ THE MEDICINES CO	EQ 40MCG BASE/ACTIVATION	N021338 001	May 22, 2006
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FERRIC AMMONIUM CITRATE

FOR SOLUTION; ORAL

FERRISELTZ

OTSUKA	600MG/PACKET	N020292 001	Oct 14, 1997
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FERRIC DERISOMALTOSE

SOLUTION; INTRAVENOUS

MONOFERRIC

+ PHARMACOSMOS AS	100MG/ML (100MG/ML)	N208171 001	Jan 16, 2020
+	500MG/5ML (100MG/ML)	N208171 002	Jan 16, 2020

FERRIC OXYHYDROXIDE

INJECTABLE; INJECTION

DEXFERRUM

AM REGENT	EQ 50MG IRON/ML	N040024 001	Feb 23, 1996
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IRON DEXTRAN

SANOFI AVENTIS US	EQ 50MG IRON/ML	N010787 002	
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PROFERDEX

NEW RIVER	EQ 50MG IRON/ML	N017807 001	
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INJECTABLE; INTRAVENOUS

VENOFER

AM REGENT	EQ 65MG IRON/3.25ML (EQ 20MG IRON/ML)	N021135 005	Mar 29, 2013
	EQ 75MG IRON/3.75ML (EQ 20MG IRON/ML)	N021135 003	Mar 29, 2005

FERRIC PYROPHOSPHATE CITRATE

POWDER; INTRAVENOUS

TRIFERIC

+ ROCKWELL MEDICAL INC	272MG IRON/PACKET	N208551 001	Apr 25, 2016
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SOLUTION; INTRAVENOUS

TRIFERIC

+ ROCKWELL MEDICAL INC	27.2MG IRON/5ML (5.44MG IRON/ML)	N206317 001	Jan 23, 2015
+	272MG IRON/50ML (5.44MG IRON/ML)	N206317 002	Sep 04, 2015

TRIFERIC AVNU

+ ROCKWELL MEDICAL INC	6.75MG IRON/4.5ML (1.5MG IRON/ML)	N212860 001	Mar 27, 2020
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FERROUS CITRATE, FE-59

INJECTABLE; INJECTION

FERROUS CITRATE FE 59

MALLINCKRODT	25uCi/ML	N016729 001	
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FERROUS SULFATE; FOLIC ACID

CAPSULE; ORAL

FOLVRON

LEDERLE	182MG; 0.33MG	N006012 003	
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FERUMOXIDES

INJECTABLE; INJECTION

FERIDEX I.V.

AMAG PHARMS INC	EQ 11.2MG IRON/ML	N020416 001	Aug 30, 1996
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FERUMOXSIIL

SUSPENSION; ORAL

GASTROMARK

AMAG PHARMS INC	EQ 0.175MG IRON/ML	N020410 001	Dec 06, 1996
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

FESOTERODINE FUMARATE

ACCORD HLTHCARE	4MG	A205012 001	Jan 04, 2023
	8MG	A205012 002	Jan 04, 2023
ACTAVIS LABS FL INC	4MG	A204868 001	Jan 04, 2023
	8MG	A204868 002	Jan 04, 2023
LUPIN LTD	4MG	A204983 001	Jan 05, 2023
	8MG	A204983 002	Jan 05, 2023

FEXOFENADINE HYDROCHLORIDE

CAPSULE;ORAL

ALLEGRA

CHATTEM SANOFI	60MG **	N020625 001	Jul 25, 1996
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FEXOFENADINE HYDROCHLORIDE

BARR	60MG	A076169 001	Jul 13, 2005
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SUSPENSION;ORAL

ALLEGRA

+ CHATTEM SANOFI	30MG/5ML	N021963 001	Oct 16, 2006
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CHILDREN'S ALLEGRA HIVES

+ CHATTEM SANOFI	30MG/5ML **	N201373 002	Jan 24, 2011
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FEXOFENADINE HYDROCHLORIDE

P AND L	30MG/5ML	A201311 001	Jul 25, 2012
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TABLET;ORAL

ALLEGRA HIVES

+ CHATTEM SANOFI	60MG	N020872 008	Jan 24, 2011
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CHILDREN'S ALLEGRA ALLERGY

+ CHATTEM SANOFI	30MG **	N020872 005	Jan 24, 2011
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CHILDREN'S ALLEGRA HIVES

+ CHATTEM SANOFI	30MG	N020872 006	Jan 24, 2011
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CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

RISING	30MG	A077081 004	Jul 21, 2011
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SUN PHARM INDS	30MG	A091567 002	Feb 06, 2012
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CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

RISING	30MG	A077081 005	Jul 21, 2011
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SUN PHARM INDS	30MG	A091567 001	Feb 06, 2012
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FEXOFENADINE HYDROCHLORIDE

BARR	30MG	A076191 001	Aug 31, 2005
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	60MG	A076191 002	Aug 31, 2005
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	180MG	A076191 003	Aug 31, 2005
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RISING	30MG	A077081 002	Apr 11, 2008
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FEXOFENADINE HYDROCHLORIDE ALLERGY

SUN PHARM INDS	60MG	A091567 004	Feb 06, 2012
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FEXOFENADINE HYDROCHLORIDE HIVES

SUN PHARM INDS	60MG	A091567 003	Feb 06, 2012
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TABLET, ORALLY DISINTEGRATING;ORAL

CHILDREN'S ALLEGRA ALLERGY

+ CHATTEM SANOFI	30MG	N021909 002	Jan 24, 2011
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CHILDREN'S ALLEGRA HIVES

+ CHATTEM SANOFI	30MG **	N021909 003	Jan 24, 2011
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CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

DR REDDYS LABS LTD	30MG	A202978 001	Jan 18, 2013
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CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD	30MG	A202978 002	Jan 18, 2013
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FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

BARR	60MG;120MG	A076236 001	Apr 14, 2005
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IMPAX PHARMS	60MG;120MG	A076298 001	Nov 12, 2010
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FINAFLOXACIN

SUSPENSION/DROPS;OTIC

XTORO

+ FONSECA BIOSCIENCES	0.3%	N206307 001	Dec 17, 2014
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FINASTERIDE

TABLET; ORAL

FINASTERIDE

ACTAVIS TOTOWA	1MG	A078371	001	Nov 05, 2013
ACTAVIS TOTOWA TEVA	5MG	A077914	001	Mar 28, 2007
GEDEON RICHTER USA	5MG	A077251	001	Dec 22, 2006
IVAX SUB TEVA PHARMS	5MG	A076340	001	Jun 19, 2006
MYLAN	1MG	A078161	001	Nov 05, 2013
	5MG	A077578	001	Dec 18, 2006
TEVA	1MG	A076905	001	Nov 05, 2013

FINGOLIMOD HYDROCHLORIDE

CAPSULE; ORAL

FINGOLIMOD HYDROCHLORIDE

BIOCON LTD	EQ 0.5MG BASE	A207979	001	Dec 04, 2019
CHARTWELL RX	EQ 0.5MG BASE	A207971	001	Jun 29, 2020
TEVA PHARMS USA	EQ 0.25MG BASE	A212152	001	Nov 12, 2021

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

IMPAX PHARMS	100MG	A076234	001	Aug 28, 2003
URISPAS				
ORTHO MCNEIL JANSSEN	100MG	N016769	001	

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

ANI PHARMS	50MG	A076030	001	Oct 28, 2002
	100MG	A076030	002	Oct 28, 2002
	150MG	A076030	003	Oct 28, 2002
CHARTWELL	50MG	A079164	001	Jul 09, 2009
	100MG	A079164	002	Jul 09, 2009
	150MG	A079164	003	Jul 09, 2009
TAMBOCOR				
+ ALVOGEN	50MG **	N018830	004	Aug 23, 1988
+	100MG **	N018830	001	Oct 31, 1985
+	150MG **	N018830	003	Jun 03, 1988
	200MG **	N018830	002	Oct 31, 1985

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

+ AVID RADIOPHARMS INC	10ML (13.5-51mCi/ML)	N202008	001	Apr 06, 2012
+	10-30ML (13.5-51mCi/ML)	N202008	002	Apr 06, 2012

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

AM REGENT	500MG/VIAL	A203008	001	Nov 22, 2017
FUDR				
+ HOSPIRA	500MG/VIAL **	N016929	001	

FLUCONAZOLE

FOR SUSPENSION; ORAL

FLUCONAZOLE

HIKMA	50MG/5ML	A076246	001	Jul 29, 2004
	200MG/5ML	A076246	002	Jul 29, 2004
IVAX SUB TEVA PHARMS	50MG/5ML	A077523	001	Sep 12, 2007
	200MG/5ML	A077523	002	Sep 12, 2007
SUN PHARM INDS LTD	50MG/5ML	A076332	001	Jul 29, 2004
	200MG/5ML	A076332	002	Jul 29, 2004
TARO PHARM INDS	50MG/5ML	A076918	001	Dec 18, 2006
	200MG/5ML	A076918	002	Dec 18, 2006

INJECTABLE; INJECTION

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER

+ PFIZER	200MG/100ML (2MG/ML)	N019950	003	Sep 29, 1992
+	400MG/200ML (2MG/ML)	N019950	005	Jul 08, 1994

DIFLUCAN IN SODIUM CHLORIDE 0.9%

+ PFIZER	200MG/100ML (2MG/ML)	N019950	001	Jan 29, 1990
+	400MG/200ML (2MG/ML)	N019950	006	Jan 29, 1990

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUCONAZOLE

INJECTABLE; INJECTION

DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+	PFIZER	200MG/100ML (2MG/ML)	N019950	002	Jan 29, 1990
+		400MG/200ML (2MG/ML)	N019950	004	Jan 29, 1990

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

	HIKMA FARMACEUTICA	200MG/100ML (2MG/ML)	A078764	001	Jan 30, 2012
		400MG/200ML (2MG/ML)	A078764	002	Jan 30, 2012
	HOSPIRA	200MG/100ML (2MG/ML)	A076304	001	Jul 29, 2004
		400MG/200ML (2MG/ML)	A076304	002	Jul 29, 2004
	MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076888	001	Mar 25, 2005
		400MG/200ML (2MG/ML)	A076888	002	Mar 25, 2005
	WOODWARD	200MG/100ML (2MG/ML)	A077988	001	May 26, 2010
		400MG/200ML (2MG/ML)	A077988	002	May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

	BAXTER HLTHCARE CORP	200MG/100ML (2MG/ML)	A077947	001	May 26, 2010
		400MG/200ML (2MG/ML)	A077947	002	May 26, 2010
	DR REDDYS	200MG/100ML (2MG/ML)	A076653	001	Jul 29, 2004
		400MG/200ML (2MG/ML)	A076653	002	Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	DR REDDYS	200MG/100ML (2MG/ML)	A076837	001	Jan 13, 2005
		400MG/200ML (2MG/ML)	A076837	002	Jan 13, 2005
	HOSPIRA	200MG/100ML (2MG/ML)	A076617	001	Jul 29, 2004
		400MG/200ML (2MG/ML)	A076617	002	Jul 29, 2004
	MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076889	001	Mar 25, 2005
		400MG/200ML (2MG/ML)	A076889	002	Mar 25, 2005

TABLET; ORAL

FLUCONAZOLE

	ANI PHARMS	50MG	A076086	001	Jul 29, 2004
		100MG	A076086	002	Jul 29, 2004
		150MG	A076086	003	Jul 29, 2004
		200MG	A076086	004	Jul 29, 2004
	GEDEON RICHTER USA	50MG	A076432	001	Jul 29, 2004
		100MG	A076432	002	Jul 29, 2004
		150MG	A076432	003	Jul 29, 2004
		200MG	A076432	004	Jul 29, 2004
	IVAX SUB TEVA PHARMS	50MG	A076077	001	Jul 29, 2004
		100MG	A076077	002	Jul 29, 2004
		150MG	A076077	003	Jul 29, 2004
		200MG	A076077	004	Jul 29, 2004
	LUPIN LTD	50MG	A209146	001	Oct 20, 2023
		100MG	A209146	002	Oct 20, 2023
		150MG	A209146	003	Oct 20, 2023
		200MG	A209146	004	Oct 20, 2023
	MYLAN	50MG	A076351	001	Jul 29, 2004
		100MG	A076351	002	Jul 29, 2004
		150MG	A076351	003	Jul 29, 2004
		200MG	A076351	004	Jul 29, 2004
	MYLAN PHARMS INC	50MG	A076042	001	Jul 29, 2004
		100MG	A076042	002	Jul 29, 2004
		150MG	A076042	003	Jul 29, 2004
		200MG	A076042	004	Jul 29, 2004
	PLIVA	50MG	A076424	001	Jul 29, 2004
		100MG	A076424	002	Jul 29, 2004
		150MG	A076424	003	Jul 29, 2004
		200MG	A076424	004	Jul 29, 2004
	RANBAXY LABS LTD	50MG	A076386	001	Jul 29, 2004
		100MG	A076386	002	Jul 29, 2004
		150MG	A076386	003	Jul 29, 2004
		200MG	A076386	004	Jul 29, 2004
	ROXANE	50MG	A076213	001	Jul 29, 2004
		100MG	A076213	002	Jul 29, 2004
		150MG	A076213	003	Jul 29, 2004
		200MG	A076213	004	Jul 29, 2004
	TEVA	50MG	A074681	001	Jul 29, 2004
		100MG	A074681	002	Jul 29, 2004
		150MG	A074681	003	Jul 29, 2004
		200MG	A074681	004	Jul 29, 2004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUCYDOSINE

CAPSULE; ORAL

FLUCYDOSINE

HIKMA	250MG	A206550 001	Oct 17, 2017
	500MG	A206550 002	Oct 17, 2017
STRIDES PHARMA	250MG	A207536 001	Jun 18, 2018
	500MG	A207536 002	Jun 18, 2018

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA

+ GENZYME CORP	50MG/VIAL **	N020038 001	Apr 18, 1991
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FLUDARABINE PHOSPHATE

ACTAVIS LLC	50MG/2ML (25MG/ML)	A203738 001	Feb 28, 2017
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HOSPIRA	50MG/VIAL	A077790 001	Apr 06, 2007
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MYLAN LABS LTD	50MG/2ML (25MG/ML)	A200647 001	Dec 21, 2011
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	50MG/VIAL	A200648 001	Oct 16, 2012
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+ SANDOZ	50MG/2ML (25MG/ML)	N022137 001	Sep 21, 2007
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TABLET; ORAL

OFORTA

SANOFI AVENTIS US	10MG	N022273 001	Dec 18, 2008
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FLUDEOXYGLUCOSE F-18

INJECTABLE; INJECTION

FLUDEOXYGLUCOSE F18

+ DOWNSTATE CLINCL	4-40mCi/ML **	N020306 001	Aug 19, 1994
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+	4-90mCi/ML **	N020306 002	Sep 25, 2001
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INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

ESSENTIAL ISOTOPES	20-300mCi/ML	A203946 001	Feb 05, 2014
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+ FEINSTEIN	20-200mCi/ML **	N021870 001	Aug 19, 2005
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HOT SHOTS NM LLC	4-500mCi/ML	A203937 001	Oct 30, 2014
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MIDWEST MEDCL	20-200mCi/ML	A203736 001	Nov 19, 2015
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UNIV TX SW MEDCTR	20-200mCi/ML	A210265 001	Feb 06, 2020
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WEILL MEDCL COLL	10-100mCi/ML **	N021768 001	Aug 05, 2004
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FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

+ CASPER PHARMA LLC	0.1MG **	N010060 001	
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FLUDROCORTISONE ACETATE

HIKMA	0.1MG	A216013 001	Oct 27, 2022
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HIKMA PHARMS	0.1MG	A091302 001	Jul 22, 2011
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FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

DR REDDYS	0.5MG/5ML (0.1MG/ML)	A076589 002	Oct 12, 2004
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	1MG/10ML (0.1MG/ML)	A076589 001	Oct 12, 2004
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RISING	1MG/10ML (0.1MG/ML)	A078595 002	May 13, 2008
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ROMAZICON

+ HOFFMANN LA ROCHE	1MG/10ML (0.1MG/ML) **	N020073 001	Dec 20, 1991
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+	0.5MG/5ML (0.1MG/ML) **	N020073 002	Dec 20, 1991
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FLUMETHASONE PIVALATE

CREAM; TOPICAL

LOCORTEN

NOVARTIS	0.03%	N016379 001	
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FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROBID

ROCHE PALO	0.25MG/INH	N018340 001	Aug 17, 1984
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AEROSPAN HFA

+ MYLAN SPECIALITY LP	0.078MG/INH	N021247 001	Jan 27, 2006
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SPRAY, METERED; NASAL

FLUNISOLIDE

APOTEX	0.029MG/SPRAY	A077436 001	Aug 09, 2007
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NASALIDE

+ IVAX RES	0.025MG/SPRAY **	N018148 001	
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NASAREL

TEVA BRANDED PHARM	0.029MG/SPRAY	N020409 001	Mar 08, 1995
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOCINOLONE ACETONIDE

CREAM;TOPICAL

FLUCET

ALPHARMA US PHARMS 0.025% A088360 001 Jan 16, 1984

FLUOCINOLONE ACETONIDE

ALLIED 0.01% A088047 001 Dec 16, 1982

ALPHARMA US PHARMS 0.01% A088361 001 Jan 16, 1984

COSETTE 0.025% A089525 001 Jul 26, 1988

INVATECH 0.025% A088045 001 Dec 16, 1982

PERRIGO NEW YORK 0.01% A086810 001 Mar 04, 1982

0.025% A086811 001 Mar 04, 1982

PHARMAFAIR 0.01% A088499 001 Aug 02, 1984

0.025% A088506 001 Aug 02, 1984

TARO 0.01% A040035 001 Oct 31, 1994

0.01% A087102 001 Apr 27, 1982

0.025% A040042 001 Oct 31, 1994

USL PHARMA 0.01% A088757 001 Feb 11, 1985

0.025% A088756 001 Mar 28, 1985

FLUONID

ALLERGAN HERBERT 0.025% A087156 002 Sep 06, 1984

FLUOTREX

SAVAGE LABS 0.01% A088174 001 May 06, 1983

0.025% A088173 001 Mar 09, 1983

SYNALAR-HP

MEDIMETRIKS PHARMS 0.2% N016161 002

GEL;TOPICAL

FLUONID

ALLERGAN HERBERT 0.025% A087300 001 May 27, 1982

OIL;TOPICAL

FLUOCINOLONE ACETONIDE

AKORN 0.01% A091514 001 Jun 25, 2015

AMNEAL 0.01% A201759 001 Oct 17, 2011

0.01% A201764 001 Oct 17, 2011

OIL/DROPS;OTIC

FLUOCINOLONE ACETONIDE

AKORN 0.01% A202705 001 Sep 09, 2016

AMNEAL 0.01% A091306 001 Oct 17, 2011

OINTMENT;TOPICAL

FLUOCINOLONE ACETONIDE

PHARMADERM 0.025% A088046 001 Dec 16, 1982

PHARMAFAIR 0.025% A088507 001 Feb 27, 1984

USL PHARMA 0.025% A088742 001 Feb 08, 1985

FLUONID

ALLERGAN HERBERT 0.025% A087157 001 Sep 06, 1984

FLUOTREX

SAVAGE LABS 0.025% A088172 001 Mar 09, 1983

SOLUTION;TOPICAL

FLUOCINOLONE ACETONIDE

ACTAVIS LABS UT INC 0.01% A208386 001 Oct 21, 2016

ALLIED 0.01% A088048 001 Dec 16, 1982

ALPHARMA US PHARMS 0.01% A087159 001 Jun 16, 1982

BAUSCH AND LOMB 0.01% A040059 001 Dec 20, 1993

COSETTE 0.01% A207441 001 Sep 28, 2016

PAI HOLDINGS PHARM 0.01% A088312 001 Jan 27, 1984

PHARMAFAIR 0.01% A088449 001 Feb 08, 1984

FLUONID

ALLERGAN HERBERT 0.01% A087158 001 Mar 17, 1983

FLUOTREX

SAVAGE LABS 0.01% A088171 001 Mar 09, 1983

FLUOCINONIDE

CREAM;TOPICAL

FLUOCINONIDE

ENCUBE 0.05% A211410 001 Oct 16, 2018

PAI HOLDINGS PHARM 0.1% A211758 001 Apr 03, 2019

PERRIGO NEW YORK 0.05% A071790 001 Jul 13, 1988

+ TARO 0.05% ** N019117 001 Jun 26, 1984

LIDEX

+ ALVOGEN 0.05% ** N016908 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOCINONIDE

CREAM; TOPICAL

LIDEX-E

+ ALVOGEN

0.05% **

N016908 003

GEL; TOPICAL

FLUOCINONIDE

PAI HOLDINGS PHARM

0.05%

A209030 001 Jun 19, 2018

OINTMENT; TOPICAL

FLUOCINONIDE

PAI HOLDINGS PHARM

0.05%

A207680 001 Sep 28, 2018

SOLUTION; TOPICAL

FLUOCINONIDE

PAI HOLDINGS PHARM

0.05%

A207554 001 Mar 18, 2019

TARO

0.05%

A072857 001 Aug 02, 1989

TEVA

0.05%

A072511 001 Feb 07, 1989

TEVA PHARMS

0.05%

A072522 001 Sep 28, 1990

LIDEX

+ ALVOGEN

0.05%

N018849 001 Apr 06, 1984

FLUORESCEIN SODIUM

INJECTABLE; INJECTION

FUNDUSCEIN-25

+ NOVARTIS

25% **

N017869 001

FLUOROMETHOLONE

CREAM; TOPICAL

OXYLONE

PHARMACIA AND UPJOHN

0.025%

N011748 001

OINTMENT; OPHTHALMIC

FML

+ ALLERGAN

0.1%

N017760 001 Sep 04, 1985

SUSPENSION/DROPS; OPHTHALMIC

FLUOR-OP

NOVARTIS

0.1%

A070185 001 Feb 27, 1986

FLUOROMETHOLONE ACETATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC

TOBRASONE

HARROW EYE

0.1%; 0.3%

N050628 001 Jul 21, 1989

FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS; OPHTHALMIC

FML-S

ALLERGAN

0.1%; 10%

N019525 001 Sep 29, 1989

FLUOROURACIL

CREAM; TOPICAL

FLUOROPLEX

+ CHARTWELL RX

1%

N016988 001

FLUOROURACIL

MYLAN

0.5%

A203122 001 Apr 20, 2015

INJECTABLE; INJECTION

ADRUCIL

PHARMACIA AND UPJOHN

50MG/ML

A081222 001 Jun 28, 1991

+

TEVA PARENTERAL

50MG/ML **

N017959 001

TEVA PARENTERAL

50MG/ML

A040023 001 Oct 18, 1991

TEVA PARENTERAL

50MG/ML

A081225 001 Aug 28, 1991

FLUOROURACIL

ABIC

50MG/ML

A088929 001 Mar 04, 1986

ABRAXIS PHARM

50MG/ML

A089152 001 Mar 21, 1986

50MG/ML

A089428 001 Jan 12, 1987

50MG/ML

A089519 001 Mar 12, 1987

BEDFORD

50MG/ML

A089508 001 Jan 26, 1988

EBEWE PHARMA

500MG/10ML (50MG/ML)

A040772 001 Aug 11, 2008

EUGIA PHARMA SPECLTS

500MG/10ML (50MG/ML)

A202668 001 Jul 17, 2012

1GM/20ML (50MG/ML)

A202668 002 Jul 17, 2012

2.5GM/50ML (50MG/ML)

A202669 001 Jul 17, 2012

5GM/100ML (50MG/ML)

A202669 002 Jul 17, 2012

FRESENIUS KABI USA

50MG/ML

A040291 001 Mar 24, 1999

50MG/ML

A040379 001 Nov 15, 2000

MARCHAR

50MG/ML

A087791 001 Jan 18, 1983

NOVAST LABS

500MG/10ML (50MG/ML)

A209219 001 Dec 12, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOROURACIL

INJECTABLE; INJECTION

FLUOROURACIL

	1GM/20ML (50MG/ML)	A209219	002	Dec 12, 2019
	2.5GM/50ML (50MG/ML)	A209271	002	Dec 11, 2019
	5GM/100ML (50MG/ML)	A209271	001	Dec 11, 2019
SANDOZ	2.5GM/50ML (50MG/ML)	A091299	001	May 02, 2011
	5GM/100ML (50MG/ML)	A091299	002	May 02, 2011
SMITH AND NEPHEW	50MG/ML	A088766	001	Dec 28, 1984
	50MG/ML	A088767	001	Dec 28, 1984
	50MG/ML	A089434	001	Mar 26, 1987
SPECTRUM PHARMS	50MG/ML	A087792	001	Oct 13, 1982
+	500MG/10ML (50MG/ML) **	N012209	001	
+	2.5GM/50ML (50MG/ML) **	N012209	002	Jul 29, 2016
TEVA PHARMS USA	500MG/10ML (50MG/ML)	A040333	001	Jan 27, 2000
	2.5GM/50ML (50MG/ML)	A040334	001	Feb 25, 2000
	5GM/100ML (50MG/ML)	A040334	002	Feb 25, 2000

SOLUTION; TOPICAL

EFUDEX

+ BAUSCH

5% **

N016831 002

FLUOROPLEX

ELORAC

1%

N016765 001

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE

SUN PHARM INDUSTRIES	EQ 10MG BASE	A075787	001	Jan 29, 2002	
	EQ 20MG BASE	A075787	002	Jan 29, 2002	
WATSON LABS	EQ 10MG BASE	A075662	001	Jan 29, 2002	
	EQ 20MG BASE	A075662	002	Jan 29, 2002	
FLUOXETINE HYDROCHLORIDE					
ACCORD HLTHCARE	EQ 10MG BASE	A202729	001	Aug 27, 2020	
	EQ 20MG BASE	A202729	002	Aug 27, 2020	
	EQ 40MG BASE	A202729	003	Aug 27, 2020	
ANI PHARMS	EQ 10MG BASE	A076287	001	May 20, 2008	
	EQ 20MG BASE	A076287	002	May 20, 2008	
BARR	EQ 10MG BASE	A074803	002	Jan 30, 2002	
	EQ 20MG BASE	A074803	001	Aug 02, 2001	
	EQ 40MG BASE	A076251	001	May 18, 2005	
BEXIMCO PHARMS USA	EQ 10MG BASE	A075807	001	Jan 29, 2002	
	EQ 20MG BASE	A075807	002	Jan 29, 2002	
CARLSBAD	EQ 10MG BASE	A076022	001	Jan 30, 2002	
	EQ 20MG BASE	A076022	002	Jan 30, 2002	
CR DOUBLE CRANE	EQ 10MG BASE	A076165	001	Feb 01, 2002	
	EQ 20MG BASE	A076165	002	Feb 01, 2002	
GRANULES	EQ 10MG BASE	A078143	001	Jan 16, 2008	
	EQ 20MG BASE	A078143	002	Jan 16, 2008	
	EQ 40MG BASE	A078143	003	Jan 16, 2008	
MYLAN	EQ 10MG BASE	A075207	001	Jan 30, 2002	
	EQ 20MG BASE	A075207	002	Jan 30, 2002	
	EQ 40MG BASE	A075207	003	May 25, 2007	
MYLAN PHARMS INC	EQ 10MG BASE	A075577	001	Jan 29, 2002	
	EQ 20MG BASE	A075577	002	Jan 29, 2002	
NATCO PHARMA USA	EQ 10MG BASE	A078045	001	Nov 17, 2008	
	EQ 20MG BASE	A078045	002	Nov 17, 2008	
SANDOZ	EQ 10MG BASE	A077469	001	Nov 17, 2008	
	EQ 20MG BASE	A077469	002	Nov 17, 2008	
SPECGX LLC	EQ 10MG BASE	A075658	001	Jan 29, 2002	
	EQ 20MG BASE	A075658	002	Jan 29, 2002	
STRIDES PHARMA	EQ 10MG BASE	A076922	001	Dec 16, 2004	
	EQ 20MG BASE	A076922	002	Dec 16, 2004	
	EQ 40MG BASE	A076922	003	Dec 16, 2004	
PROZAC					
ELI LILLY AND CO	EQ 60MG BASE	N018936	004	Jun 15, 1999	
SARAFEM					
+	ELI LILLY AND CO	EQ 10MG BASE **	N018936	007	Jul 06, 2000
+		EQ 20MG BASE **	N018936	008	Jul 06, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

FLUOXETINE HYDROCHLORIDE

BARR EQ 90MG BASE A076237 001 Mar 24, 2010

PROZAC WEEKLY

+ LILLY EQ 90MG BASE ** N021235 001 Feb 26, 2001

SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC EQ 20MG BASE/5ML A075690 001 Jan 31, 2002

AKORN EQ 20MG BASE/5ML A075525 001 Jun 27, 2002

CHARTWELL MOLECULAR EQ 20MG BASE/5ML A076458 001 May 14, 2004

PHARMOBEDIENT CNSLTG EQ 20MG BASE/5ML A075514 001 Aug 29, 2002

SPECGX LLC EQ 20MG BASE/5ML A075920 001 Jan 29, 2002

PROZAC

+ LILLY EQ 20MG BASE/5ML ** N020101 001 Apr 24, 1991

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

BARR EQ 10MG BASE A075810 001 Feb 01, 2002

FOSUN PHARMA EQ 10MG BASE A076024 001 Jan 29, 2002

G AND W LABS INC EQ 60MG BASE A212191 001 Jul 05, 2019

IVAX SUB TEVA PHARMS EQ 10MG BASE A075865 001 Feb 28, 2002

EQ 40MG BASE A075865 003 Aug 30, 2004

RISING EQ 10MG BASE A075755 001 Aug 02, 2001

EQ 20MG BASE A075755 002 Aug 02, 2001

SLATE RUN PHARMA EQ 10MG BASE A211444 001 Sep 13, 2022

EQ 20MG BASE A211444 002 Sep 13, 2022

PROZAC

+ LILLY EQ 10MG BASE ** N020974 001 Mar 09, 1999

+ EQ 20MG BASE ** N020974 002 Mar 09, 1999

SARAFEM

+ APIL EQ 10MG BASE ** N021860 001 May 19, 2006

+ EQ 15MG BASE ** N021860 002 May 19, 2006

+ EQ 20MG BASE ** N021860 003 May 19, 2006

SELFEMRA

TEVA PHARMS USA EQ 10MG BASE A200151 001 Feb 03, 2014

EQ 15MG BASE A200151 002 Feb 03, 2014

EQ 20MG BASE A200151 003 Feb 03, 2014

FLUOXYMESTERONE

TABLET;ORAL

ANDROID-F

VALEANT PHARM INTL 10MG A087196 001

FLUOXYMESTERONE

UPSHER SMITH LABS 10MG A088342 001 Oct 21, 1983

VALEANT PHARM INTL 10MG A088221 001 May 05, 1983

WATSON LABS 2MG A088260 001 Dec 06, 1983

5MG A088265 001 Dec 06, 1983

10MG A088309 001 Dec 06, 1983

HALOTESTIN

PHARMACIA AND UPJOHN 2MG N010611 002

5MG N010611 006

10MG N010611 010

ORA-TESTRYL

BRISTOL MYERS SQUIBB 2MG N011359 001

5MG N011359 002

FLUPHENAZINE DECANOATE

INJECTABLE;INJECTION

FLUPHENAZINE DECANOATE

HOSPIRA 25MG/ML A074966 001 Apr 16, 1998

TEVA PARENTERAL 25MG/ML A074795 001 Sep 10, 1996

PROLIXIN DECANOATE

+ BRISTOL MYERS SQUIBB 25MG/ML ** N016727 001

FLUPHENAZINE ENANTHATE

INJECTABLE;INJECTION

PROLIXIN ENANTHATE

APOTHECON 25MG/ML ** N016110 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS

5MG/ML

A073058 001 Aug 30, 1991

PERMITIL

SCHERING

5MG/ML **

N016008 001

PROLIXIN

APOTHECON

5MG/ML

A070533 001 Nov 07, 1985

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS

2.5MG/5ML

A081310 001 Apr 29, 1993

PROLIXIN

+ APOTHECON

2.5MG/5ML **

N012145 003

INJECTABLE; INJECTION

PROLIXIN

APOTHECON

2.5MG/ML **

N011751 005

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

PRASCO

1MG

A089804 002 Aug 12, 1988

2.5MG

A089804 003 Aug 12, 1988

5MG

A089804 004 Aug 12, 1988

10MG

A089804 001 Aug 12, 1988

SANDOZ

1MG

A089586 002 Oct 16, 1987

2.5MG

A089586 003 Oct 16, 1987

5MG

A089586 004 Oct 16, 1987

10MG

A089586 001 Oct 16, 1987

WATSON LABS

1MG

A088555 001 Dec 18, 1987

2.5MG

A088544 001 Dec 18, 1987

5MG

A088527 001 Dec 18, 1987

10MG

A088550 001 Dec 18, 1987

PERMITIL

SCHERING

0.25MG

N012034 001

2.5MG

N012034 004

5MG

N012034 005

10MG

N012034 006

PROLIXIN

+ APOTHECON

1MG **

N011751 004

+

2.5MG **

N011751 001

+

5MG **

N011751 003

+

10MG **

N011751 002

TABLET, EXTENDED RELEASE; ORAL

PERMITIL

SCHERING

1MG

N012419 004

FLUPREDNISOLONE

TABLET; ORAL

ALPHADROL

PHARMACIA AND UPJOHN

1.5MG

N012259 002

FLURANDRENOLIDE

CREAM; TOPICAL

CORDRAN SP

+ ALMIRALL

0.025% **

N012806 003

+

0.05%

N012806 002

FLURANDRENOLIDE

CHARTWELL RX

0.05%

A205342 001 Apr 13, 2016

LOTION; TOPICAL

CORDRAN

+ ALMIRALL

0.05%

N013790 001

FLURANDRENOLIDE

ALPHARMA US PHARMS

0.05%

A087203 001 Apr 29, 1982

CHARTWELL RX

0.05%

A205343 001 Dec 22, 2016

OINTMENT; TOPICAL

CORDRAN

+ ALMIRALL

0.025% **

N012806 004

+

0.05%

N012806 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLURANDRENOLIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

CORDRAN N

LILLY

0.05%;EQ 3.5MG BASE/GM

N050346 001

OINTMENT; TOPICAL

CORDRAN N

LILLY

0.05%;EQ 3.5MG BASE/GM

N050345 001

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

DALMANE

VALEANT PHARM INTL

15MG **

N016721 001

+

30MG **

N016721 002

FLURAZEPAM HYDROCHLORIDE

CHARTWELL RX

15MG

A071205 001 Nov 25, 1986

30MG

A071068 001 Nov 25, 1986

HALSEY

15MG

A071808 001 Jan 07, 1988

30MG

A071809 001 Jan 07, 1988

HIKMA

30MG

A071108 001 Dec 08, 1986

HIKMA INTL PHARMS

15MG

A071107 001 Dec 08, 1986

PUREPAC PHARM

15MG

A071927 001 Sep 09, 1987

30MG

A071551 001 Sep 09, 1987

RISING

15MG

A071717 002 Jul 31, 1991

30MG

A071717 001 Jul 31, 1991

STRIDES PHARMA

15MG

A070444 001 Mar 20, 1986

30MG

A070445 001 Mar 20, 1986

SUN PHARM INDUSTRIES

15MG

A070454 001 Aug 04, 1986

30MG

A070455 001 Aug 04, 1986

SUPERPHARM

15MG

A071659 001 Aug 04, 1988

30MG

A071660 001 Aug 04, 1988

USL PHARMA

15MG

A070562 001 Jul 09, 1987

30MG

A070563 001 Jul 09, 1987

WARNER CHILCOTT

15MG

A071767 001 Dec 04, 1987

30MG

A071768 001 Dec 04, 1987

FLURBIPROFEN

TABLET; ORAL

ANSAID

+

PHARMACIA AND UPJOHN

50MG

N018766 002 Oct 31, 1988

+

100MG

N018766 003 Oct 31, 1988

FLURBIPROFEN

IVAX SUB TEVA PHARMS

50MG

A074411 001 May 31, 1995

100MG

A074411 002 May 31, 1995

MYLAN

50MG

A074358 001 Jun 20, 1994

100MG

A074358 002 Jun 20, 1994

PLIVA

50MG

A074647 001 Apr 01, 1997

100MG

A074647 002 Apr 01, 1997

RISING

50MG

A074448 001 Jul 28, 1995

100MG

A074448 002 Jul 28, 1995

SUN PHARM INDS INC

50MG

A075058 001 Apr 27, 2001

100MG

A075058 002 Apr 27, 2001

TEVA

50MG

A074405 002 May 24, 1995

100MG

A074405 001 May 24, 1995

THERAGEN

100MG

A074560 002 May 16, 1997

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

OCUFEN

+

ALLERGAN

0.03% **

N019404 001 Dec 31, 1986

FLUTAMIDE

CAPSULE; ORAL

EULEXIN

+

SCHERING

125MG **

N018554 001 Jan 27, 1989

FLUTAMIDE

ACTAVIS LABS FL INC

125MG

A075820 001 Sep 18, 2001

CHARTWELL RX

125MG

A075818 001 Sep 18, 2001

CIPLA

125MG

A075780 001 Sep 19, 2001

MYLAN

125MG

A076224 001 May 09, 2003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUTEMETAMOL F-18

INJECTABLE; INTRAVENOUS

VIZAMYL

+ GE HEALTHCARE 40.5mCi/10ML (4.05mCi/ML) N203137 001 Oct 25, 2013

FLUTICASONE PROPIONATE

AEROSOL, METERED; INHALATION

FLOVENT

GLAXOSMITHKLINE 0.044MG/INH N020548 001 Mar 27, 1996

0.11MG/INH N020548 002 Mar 27, 1996

0.22MG/INH N020548 003 Mar 27, 1996

CREAM; TOPICAL

CUTIVATE

+ FOUGERA PHARMS 0.05% ** N019958 001 Dec 18, 1990

FLUTICASONE PROPIONATE

ENCUBE 0.05% A076633 001 May 14, 2004

NESHER PHARMS 0.05% A076865 001 Sep 10, 2004

LOTION; TOPICAL

CUTIVATE

+ FOUGERA PHARMS 0.05% ** N021152 001 Mar 31, 2005

OINTMENT; TOPICAL

CUTIVATE

+ FOUGERA PHARMS 0.005% ** N019957 001 Dec 14, 1990

FLUTICASONE PROPIONATE

BRIGHT 0.005% A215343 001 Sep 01, 2023

FOUGERA PHARMS 0.005% A076300 001 May 14, 2004

TARO PHARM INDS 0.005% A077145 001 Jun 14, 2005

POWDER; INHALATION

ARMONAIR DIGIHALER

+ TEVA PHARM 0.03MG/INH N208798 008 Apr 08, 2022

ARMONAIR RESPICLICK

+ TEVA PHARM 0.03MG/INH N208798 007 Jul 09, 2021

+ 0.055MG/INH N208798 001 Jan 27, 2017

+ 0.113MG/INH N208798 002 Jan 27, 2017

+ 0.232MG/INH N208798 003 Jan 27, 2017

FLOVENT

GLAXOSMITHKLINE 0.044MG/INH N020549 001 Nov 07, 1997

0.088MG/INH N020549 002 Nov 07, 1997

0.22MG/INH N020549 003 Nov 07, 1997

SPRAY, METERED; NASAL

FLONASE

+ GLAXOSMITHKLINE 0.05MG/SPRAY ** N020121 001 Oct 19, 1994

FLUVASTATIN SODIUM

CAPSULE; ORAL

LESCOL

+ NOVARTIS EQ 20MG BASE ** N020261 001 Dec 31, 1993

+ EQ 40MG BASE ** N020261 002 Dec 31, 1993

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM

MYLAN EQ 80MG BASE A202458 001 Sep 11, 2015

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE

TORRENT 100MG A203240 001 Oct 31, 2014

150MG A203240 002 Oct 31, 2014

LUVOX CR

+ JAZZ PHARMS 100MG ** N022033 001 Feb 28, 2008

+ 150MG ** N022033 002 Feb 28, 2008

TABLET; ORAL

FLUVOXAMINE MALEATE

ACTAVIS ELIZABETH 25MG A075901 001 Dec 28, 2000

50MG A075901 002 Dec 28, 2000

100MG A075901 003 Dec 28, 2000

ANI PHARMS 25MG A075897 001 Jan 25, 2001

25MG A075898 001 Mar 12, 2001

50MG A075897 002 Jan 25, 2001

50MG A075898 002 Mar 12, 2001

100MG A075897 003 Jan 25, 2001

100MG A075898 003 Mar 12, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUVOXAMINE MALEATE

TABLET; ORAL

FLUVOXAMINE MALEATE

CHARTWELL RX	25MG	A075900 001	Feb 23, 2006
	50MG	A075900 002	Feb 23, 2006
	100MG	A075900 003	Feb 23, 2006
HERITAGE PHARMA	25MG	A075894 001	Apr 18, 2001
	50MG	A075894 002	Apr 18, 2001
	100MG	A075894 003	Apr 18, 2001
MYLAN	25MG	A075889 001	Nov 29, 2000
	50MG	A075889 002	Nov 29, 2000
	50MG	A075950 001	Oct 15, 2001
	100MG	A075889 003	Nov 29, 2000
	100MG	A075950 002	Oct 15, 2001
SUN PHARM INDUSTRIES	25MG	A076125 001	Apr 29, 2002
	50MG	A076125 002	Apr 29, 2002
	100MG	A076125 003	Apr 29, 2002
SYNTHON PHARMS	25MG	A075899 001	Jan 17, 2001
	50MG	A075899 002	Jan 17, 2001
	100MG	A075899 003	Jan 17, 2001
TEVA	25MG	A075893 001	Sep 10, 2002
	50MG	A075893 002	Sep 10, 2002
	100MG	A075893 003	Sep 10, 2002
UPSHER SMITH LABS	25MG	A075887 001	Jan 05, 2001
	50MG	A075887 002	Jan 05, 2001
	100MG	A075887 003	Jan 05, 2001
LUVOX			
+ SOLVAY	25MG **	N020243 001	Dec 05, 1994
+	50MG **	N020243 002	Dec 05, 1994
+	100MG **	N020243 003	Dec 05, 1994
+	150MG **	N020243 004	Dec 05, 1994

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

BEN VENUE	5MG/ML	A081066 001	Dec 29, 1993
FOLVITE			
+ WYETH PHARMS INC	5MG/ML **	N005897 008	

TABLET; ORAL

FOLIC ACID

BARR	1MG	A089177 001	Jan 08, 1986
CONTRACT PHARMACAL	1MG	A085061 001	
EVERYLIFE	1MG	A080755 001	
HALSEY	1MG	A083598 001	
HIKMA PHARMS	1MG	A080600 001	
IMPAX LABS	1MG	A080686 001	
IVAX SUB TEVA PHARMS	1MG	A083000 001	
JUBILANT CADISTA	1MG	A040514 001	Jun 14, 2005
LANNETT	1MG	A080816 001	
LILLY	1MG **	N006135 003	
MK LABS	1MG	A083526 001	
NEXGEN PHARMA INC	1MG	A084915 001	
PHARMERAL	1MG	A084158 001	
PIONEER PHARMS	1MG	A088949 001	Sep 13, 1985
PUREPAC PHARM	1MG	A080784 001	
SANDOZ	1MG	A084472 001	
SUN PHARM INDUSTRIES	1MG	A040582 001	Jul 18, 2005
TABLICAPS	1MG	A083133 002	
UDL	1MG	A088199 001	Mar 29, 1983
USL PHARMA	1MG	A087828 001	May 13, 1982
VALEANT PHARM INTL	1MG	A080903 001	
VANGARD	1MG	A088730 001	Mar 23, 1984
VINTAGE	1MG	A040756 001	Jun 04, 2010
VINTAGE PHARMS	1MG	A086296 001	
WATSON LABS	1MG	A083141 001	
	1MG	A085141 002	
WHITEWORTH TOWN PLSN	1MG	A080691 002	
FOLICET			
MISSION PHARMA	1MG	A087438 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOLIC ACID

TABLET; ORAL

FOLVITE

WYETH PHARMS INC

1MG **

N005897 004

FOMEPIZOLE

INJECTABLE; INJECTION

ANTIZOL

+ PAR PHARM INC

1.5GM/1.5ML (1GM/ML) **

N020696 001 Dec 04, 1997

FOMEPIZOLE

MYLAN INSTITUTIONAL

1.5GM/1.5ML (1GM/ML)

A079033 001 Apr 07, 2009

FOMIVIRSEN SODIUM

INJECTABLE; INJECTION

VITRAVENE PRESERVATIVE FREE

NOVARTIS

6.6MG/ML

N020961 001 Aug 26, 1998

FORMOTEROL FUMARATE

POWDER; INHALATION

FORADIL

+ NOVARTIS

0.012MG/INH **

N020831 001 Feb 16, 2001

FORADIL CERTIHALER

NOVARTIS

0.0085MG/INH

N021592 001 Dec 15, 2006

FOSAMPRENAVIR CALCIUM

SUSPENSION; ORAL

LEXIVA

+ VIIV HLTHCARE

EQ 50MG BASE/ML

N022116 001 Jun 14, 2007

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND

+ MERCK AND CO INC

EQ 115MG BASE/VIAL **

N022023 001 Jan 25, 2008

FOSAPREPITANT DIMEGLUMINE

APOTEX

EQ 150MG BASE/VIAL

A205020 001 Sep 05, 2019

ARTHUR GRP

EQ 150MG BASE/VIAL

A211624 001 Sep 05, 2019

EQ 150MG BASE/VIAL

A213199 001 Oct 04, 2021

MYLAN LABS LTD

EQ 115MG BASE/VIAL

A204015 001 Sep 05, 2019

SANDOZ

EQ 150MG BASE/VIAL

A203939 001 Dec 08, 2020

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

HOSPIRA

2.4GM/100ML

A077174 001 May 31, 2005

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

ACTAVIS LABS FL INC

10MG

A076620 001 Oct 15, 2004

20MG

A076620 002 Oct 15, 2004

40MG

A076620 003 Oct 15, 2004

RANBAXY LABS LTD

10MG

A076580 001 Apr 23, 2004

20MG

A076580 002 Apr 23, 2004

40MG

A076580 003 Apr 23, 2004

UPSHER SMITH LABS

10MG

A076188 001 Oct 08, 2004

20MG

A076188 002 Oct 08, 2004

40MG

A076188 003 Oct 08, 2004

WATSON LABS

10MG

A076987 001 Dec 23, 2004

10MG

A077531 001 Aug 31, 2006

20MG

A076987 002 Dec 23, 2004

20MG

A077531 002 Aug 31, 2006

40MG

A076987 003 Dec 23, 2004

40MG

A077531 003 Aug 31, 2006

MONOPRIL

+ BRISTOL MYERS SQUIBB

10MG **

N019915 002 May 16, 1991

+

20MG **

N019915 003 May 16, 1991

+

40MG **

N019915 004 Mar 28, 1995

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

ANI PHARMS	10MG;12.5MG	A076608	001	Dec 03, 2004
	10MG;12.5MG	A077144	001	Aug 16, 2005
	20MG;12.5MG	A076608	002	Dec 03, 2004
	20MG;12.5MG	A077144	002	Aug 16, 2005
AVET LIFESCIENCES	10MG;12.5MG	A079025	001	Sep 17, 2010
	20MG;12.5MG	A079025	002	Sep 17, 2010
MYLAN	10MG;12.5MG	A077705	001	Aug 14, 2006
	20MG;12.5MG	A077705	002	Aug 14, 2006
SUN PHARM INDS LTD	10MG;12.5MG	A076739	001	Dec 17, 2004
	20MG;12.5MG	A076739	002	Dec 17, 2004
TEVA	10MG;12.5MG	A076945	001	Jul 05, 2006
	20MG;12.5MG	A076945	002	Jul 05, 2006
MONOPRIL-HCT				
+ BRISTOL MYERS SQUIBB	10MG;12.5MG **	N020286	002	Nov 30, 1994
+	20MG;12.5MG **	N020286	001	Nov 30, 1994

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

POWDER; INTRAVENOUS

AKYNZEO

+ HELSINN HLTHCARE	EQ 235MG BASE/VIAL;EQ 0.25MG BASE/VIAL	N210493	001	Apr 19, 2018
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FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

AM REGENT	EQ 50MG PHENYTOIN NA/ML	A078277	001	Aug 06, 2007
	EQ 50MG PHENYTOIN NA/ML	A090099	001	May 13, 2010
AMNEAL	EQ 50MG PHENYTOIN NA/ML	A078476	001	Mar 18, 2008
APOTEX INC	EQ 50MG PHENYTOIN NA/ML	A078126	001	Aug 06, 2007
DR REDDYS	EQ 50MG PHENYTOIN NA/ML	A076886	001	Aug 06, 2007
HOSPIRA	EQ 50MG PHENYTOIN NA/ML	A078158	001	Aug 06, 2007
MYLAN LABS LTD	EQ 50MG PHENYTOIN NA/ML	A078736	001	Jun 08, 2010

SOLUTION; INTRAVENOUS

SESQUIENT

+ LUPIN	EQ 100MG PHENYTOIN NA/2ML (EQ 50MG PHENYTOIN NA/ML)	N210864	001	Nov 05, 2020
+	EQ 500MG PHENYTOIN NA/10ML (EQ 50MG PHENYTOIN NA/ML)	N210864	002	Nov 05, 2020

FOSPROPOFOL DISODIUM

SOLUTION; INTRAVENOUS

LUSEDRA

EISAI INC	1050MG/30ML (35MG/ML)	N022244	001	Dec 12, 2008
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FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVATRIPTAN SUCCINATE

MYLAN	EQ 2.5MG BASE	A202931	001	Aug 28, 2014
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FULVESTRANT

SOLUTION; INTRAMUSCULAR

FASLODEX

+ ASTRAZENECA	125MG/2.5ML (50MG/ML)	N021344	002	Apr 25, 2002
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FULVESTRANT

APOTEX	250MG/5ML (50MG/ML)	A211730	001	Jun 11, 2021
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FURAZOLIDONE

SUSPENSION; ORAL

FUROXONE

SHIRE	50MG/15ML	N011323	002	
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TABLET; ORAL

FUROXONE

SHIRE	100MG	N011270	002	
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FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

ABRAXIS PHARM	10MG/ML	N018507	001	Jul 30, 1982
	10MG/ML	N019036	001	Aug 13, 1984
+ AM REGENT	10MG/ML **	N018579	001	Nov 30, 1983
ASTRAZENECA	10MG/ML	A070014	001	Sep 09, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

HIKMA	10MG/ML	A071439 001	Sep 14, 1990
	10MG/ML	N018267 001	
HOSPIRA	10MG/ML	A070578 001	Jul 08, 1987
	10MG/ML	A072080 001	Aug 13, 1991
	10MG/ML	A074337 001	Oct 31, 1994
IGI LABS INC	10MG/ML	A070095 001	Sep 09, 1985
	10MG/ML	A070096 001	Sep 09, 1985
INTL MEDICATION	10MG/ML	N018025 001	
MARSAM PHARMS LLC	10MG/ML	A074017 001	Jun 30, 1994
SMITH AND NEPHEW	10MG/ML	A070023 001	Feb 05, 1986
	10MG/ML	A070078 001	Feb 05, 1986
WARNER CHILCOTT	10MG/ML	N018420 001	Feb 26, 1982
WATSON LABS	10MG/ML	A070019 001	Sep 22, 1986
	10MG/ML	A070604 001	Jan 02, 1987
WYETH AYERST	10MG/ML	N018670 001	Jul 20, 1982

LASIX

+ SANOFI AVENTIS US 10MG/ML **

N016363 001

SOLUTION; ORAL

FUROSEMIDE

PHARMOBEDIANT CNSLTG	10MG/ML	A070655 001	Oct 02, 1987
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LASIX

SANOFI AVENTIS US 10MG/ML

N017688 001

TABLET; ORAL

FUROSEMIDE

ANI PHARMS	20MG	A071379 001	Jan 02, 1987
	40MG	A070413 001	Feb 26, 1986
	80MG	A071594 001	Feb 09, 1988
CHARTWELL RX	20MG	N018413 001	Nov 30, 1983
	40MG	N018413 002	Nov 30, 1983
EPIC PHARMA LLC	40MG	N018750 002	Jul 30, 1984
INTL MEDICATION	20MG	N018753 001	Feb 28, 1984
	40MG	N018753 002	Feb 28, 1984
KALAPHARM	20MG	N018868 001	Jun 28, 1983
	40MG	N018868 002	Jun 28, 1983
STRIDES PHARMA	20MG	N018415 001	Jul 27, 1982
	40MG	N018415 002	Jul 27, 1982
	80MG	N018415 003	Nov 26, 1984
SUN PHARM INDS INC	20MG	A091258 001	Apr 01, 2014
	40MG	A091258 002	Apr 01, 2014
	40MG	N018790 001	Nov 29, 1983
	80MG	A091258 003	Apr 01, 2014
SUN PHARM INDUSTRIES	20MG	A070043 001	Sep 26, 1985
	80MG	A070100 001	Jan 26, 1988
SUPERPHARM	20MG	N018370 002	Jun 26, 1984
	40MG	N018370 001	Feb 10, 1983
WARNER CHILCOTT	20MG	N018419 001	Jan 31, 1983
	40MG	N018419 002	Jan 31, 1983
	80MG	N018419 003	Nov 13, 1984
WATSON LABS	20MG	A070412 001	Feb 26, 1986
	20MG	N018369 001	May 14, 1982
	40MG	A070450 001	Nov 22, 1985
	40MG	N018369 002	May 14, 1982
WATSON LABS TEVA	20MG	A070449 001	Nov 22, 1985
	80MG	A070528 001	Jan 07, 1986

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

CHARTWELL RX	100MG	A091008 001	Oct 26, 2017
	300MG	A091008 002	Oct 26, 2017
	400MG	A091008 003	Oct 26, 2017
HIKMA	100MG	A078150 001	Sep 25, 2007
	300MG	A078150 002	Sep 25, 2007
	400MG	A078150 003	Sep 25, 2007
MYLAN	100MG	A090158 001	Feb 14, 2011
	300MG	A090158 002	Feb 14, 2011
	400MG	A090158 003	Feb 14, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

SANDOZ	100MG	A075428 001	Jan 24, 2006
	100MG	A075539 001	Apr 06, 2005
	300MG	A075428 002	Jan 24, 2006
	300MG	A075539 002	Apr 06, 2005
	400MG	A075428 003	Jan 24, 2006
	400MG	A075539 003	Apr 06, 2005
SUN PHARM INDS LTD	100MG	A076606 001	Oct 07, 2005
	300MG	A076606 002	Oct 07, 2005
	400MG	A076606 003	Oct 07, 2005
SUN PHARM INDUSTRIES	100MG	A076537 001	Jun 30, 2005
	300MG	A076537 002	Jun 30, 2005
	400MG	A076537 003	Jun 30, 2005
TEVA PHARMS	100MG	A075435 001	Oct 08, 2004
	300MG	A075435 002	Oct 08, 2004
	400MG	A075435 003	Oct 08, 2004
WATSON LABS	100MG	A075485 003	May 11, 2007
	300MG	A075485 002	May 11, 2007
	400MG	A075485 001	May 11, 2007

SOLUTION; ORAL

GABAPENTIN

PAI HOLDINGS PHARM	250MG/5ML	A211330 001	Dec 03, 2019
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TABLET; ORAL

GABAPENTIN

CHARTWELL RX	600MG	A076120 001	Jan 27, 2006
	800MG	A076120 002	Jan 27, 2006
HIKMA PHARMS	600MG	A078782 001	Jul 21, 2011
	800MG	A078782 002	Jul 21, 2011
INVATECH	600MG	A076877 001	Jul 06, 2006
	800MG	A076877 002	Jul 06, 2006
IVAX SUB TEVA PHARMS	600MG	A076017 004	Apr 29, 2005
	800MG	A076017 005	Apr 29, 2005
LUPIN LTD	600MG	A209306 001	Aug 24, 2018
	800MG	A209306 002	Aug 24, 2018
MYLAN PHARMS INC	600MG	A090335 001	Jun 01, 2010
	800MG	A090335 002	Jun 01, 2010
RANBAXY	600MG	A076605 001	Sep 14, 2005
	800MG	A076605 002	Sep 14, 2005
TEVA	600MG	A075827 001	Dec 15, 2004
	800MG	A075827 002	Dec 15, 2004
TEVA PHARMS USA	600MG	A205807 001	Mar 10, 2017
	800MG	A205807 002	Mar 10, 2017

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

GE HEALTHCARE	14.35GM/50ML (287MG/ML)	N022066 001	Sep 05, 2007
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GADOFOSVESET TRISODIUM

SOLUTION; INTRAVENOUS

ABLAVAR

LANTHEUS MEDCL	2440MG/10ML (244MG/ML)	N021711 001	Dec 22, 2008
	3660MG/15ML (244MG/ML)	N021711 002	Dec 22, 2008

GADOPENTETATE DIMEGLUMINE

INJECTABLE; INJECTION

MAGNEVIST

+ BAYER HLTHCARE	469.01MG/ML	N019596 001	Jun 02, 1988
+	469.01MG/ML	N021037 001	Mar 10, 2000

GADOVERSE TAMIDE

INJECTABLE; INJECTION

OPTIMARK

+ LIEBEL-FLARSHEIM	1654.5MG/5ML (330.9MG/ML)	N020937 001	Dec 08, 1999
+	3309MG/10ML (330.9MG/ML)	N020937 002	Dec 08, 1999
+	4963.5MG/15ML (330.9MG/ML)	N020937 003	Dec 08, 1999
+	6618MG/20ML (330.9MG/ML)	N020937 004	Dec 08, 1999
+	16.545GM/50ML (330.9MG/ML)	N020975 001	Dec 08, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GADOVERSETAMIDE

INJECTABLE;INJECTION

OPTIMARK IN PLASTIC CONTAINER

+	LIEBEL-FLARSHEIM	3309MG/10ML (330.9MG/ML)	N020976	002	Dec 08, 1999
+		4963.5MG/15ML (330.9MG/ML)	N020976	003	Dec 08, 1999
+		6618MG/20ML (330.9MG/ML)	N020976	004	Dec 08, 1999
+		9927MG/30ML (330.9MG/ML)	N020976	001	Dec 08, 1999

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE;ORAL

GALANTAMINE HYDROBROMIDE

	IMPAX LABS	EQ 8MG BASE	A078484	001	May 27, 2009
		EQ 16MG BASE	A078484	002	May 27, 2009
		EQ 24MG BASE	A078484	003	May 27, 2009
	MYLAN	EQ 8MG BASE	A090900	001	Jan 24, 2011
		EQ 16MG BASE	A090900	002	Jan 24, 2011
		EQ 24MG BASE	A090900	003	Jan 24, 2011
	RAZADYNE ER				
+	JANSSEN PHARMS	EQ 8MG BASE	N021615	001	Apr 01, 2005
+		EQ 16MG BASE	N021615	002	Apr 01, 2005
+		EQ 24MG BASE	N021615	003	Apr 01, 2005

SOLUTION;ORAL

RAZADYNE

	JANSSEN PHARMS	4MG/ML **	N021224	001	Jun 22, 2001
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TABLET;ORAL

GALANTAMINE HYDROBROMIDE

	APOTEX INC	EQ 4MG BASE	A077781	001	Sep 27, 2011
		EQ 8MG BASE	A077781	002	Sep 27, 2011
		EQ 12MG BASE	A077781	003	Sep 27, 2011
	CHARTWELL RX	EQ 4MG BASE	A077587	001	Jul 09, 2009
		EQ 8MG BASE	A077587	002	Jul 09, 2009
		EQ 12MG BASE	A077587	003	Jul 09, 2009
	HERITAGE PHARMA	EQ 4MG BASE	A077585	001	Sep 15, 2009
		EQ 8MG BASE	A077585	002	Sep 15, 2009
		EQ 12MG BASE	A077585	003	Sep 15, 2009
	HIKMA	EQ 4MG BASE	A077608	001	Feb 11, 2009
		EQ 8MG BASE	A077608	002	Feb 11, 2009
		EQ 12MG BASE	A077608	003	Feb 11, 2009
	MYLAN	EQ 4MG BASE	A077590	001	May 29, 2009
		EQ 4MG BASE	A077603	001	Aug 28, 2008
		EQ 8MG BASE	A077590	002	May 29, 2009
		EQ 8MG BASE	A077603	002	Aug 28, 2008
		EQ 12MG BASE	A077590	003	May 29, 2009
		EQ 12MG BASE	A077603	003	Aug 28, 2008
	RAZADYNE				
+	JANSSEN PHARMS	EQ 4MG BASE **	N021169	001	Feb 28, 2001
+		EQ 8MG BASE **	N021169	002	Feb 28, 2001
+		EQ 12MG BASE **	N021169	003	Feb 28, 2001

GALLAMINE TRIETHIODIDE

INJECTABLE;INJECTION

FLAXEDIL

	DAVIS AND GECK	20MG/ML	N007842	001	
		100MG/ML	N007842	002	

GALLIUM CITRATE GA-67

INJECTABLE;INJECTION

GALLIUM CITRATE GA 67

	GE HEALTHCARE	1mCi/ML	N017700	001	
	LANTHEUS MEDCL	2mCi/ML	N017478	001	
	NEOSCAN				
	GE HEALTHCARE	2mCi/ML	N017655	001	

GALLIUM NITRATE

INJECTABLE;INJECTION

GANITE

	CHAPTER 7 TRUSTEE	25MG/ML **	N019961	002	Jan 17, 1991
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GANCICLOVIR

CAPSULE; ORAL

CYTOVENE

+ ROCHE PALO

250MG **

N020460 001 Dec 22, 1994

+

500MG **

N020460 002 Dec 12, 1997

GANCICLOVIR

RANBAXY LABS LTD

250MG

A076457 001 Jun 27, 2003

500MG

A076457 002 Jun 27, 2003

IMPLANT; IMPLANTATION

VITRASERT

BAUSCH AND LOMB

4.5MG

N020569 001 Mar 04, 1996

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

CYTOVENE

+ CHEPLAPHARM

EQ 500MG BASE/VIAL **

N019661 001 Jun 23, 1989

GANCICLOVIR SODIUM

AM REGENT

EQ 500MG BASE/VIAL

A202624 001 Sep 18, 2013

CUSTOPHARM INC

EQ 500MG BASE/VIAL

A212001 001 Jun 20, 2019

STERISCIENCE

EQ 500MG BASE/VIAL

A204560 001 Nov 17, 2017

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

APOTEX INC

0.3%

A079084 001 Aug 19, 2011

RISING

0.5%

A206446 001 Jun 08, 2018

TORRENT

0.5%

A213542 001 Nov 03, 2021

ZYMAR

+ ALLERGAN

0.3%

N021493 001 Mar 28, 2003

GEFITINIB

TABLET; ORAL

IRESSA

ASTRAZENECA

250MG

N021399 001 May 05, 2003

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

ACTAVIS INC

200MG/5.26ML (38MG/ML)

A204549 001 Apr 11, 2016

1GM/26.3ML (38MG/ML)

A204549 002 Apr 11, 2016

2GM/52.6ML (38MG/ML)

A204549 003 Apr 11, 2016

ACTAVIS TOTOWA

EQ 200MG BASE/VIAL

A079160 001 Jul 25, 2011

EQ 1GM BASE/VIAL

A079160 002 Jul 25, 2011

EQ 2GM BASE/VIAL

A079160 003 Jul 28, 2016

AM REGENT

EQ 200MG BASE/VIAL

A202031 001 May 07, 2013

EQ 1GM BASE/VIAL

A202031 002 May 07, 2013

APOTEX

200MG/5.26ML (38MG/ML)

A206776 001 May 23, 2017

1GM/26.3ML (38MG/ML)

A206776 002 May 23, 2017

2GM/52.6ML (38MG/ML)

A206776 003 May 23, 2017

EMCURE PHARMS LTD

EQ 200MG BASE/VIAL

A202063 001 Sep 11, 2012

EQ 1GM BASE/VIAL

A202063 002 Sep 11, 2012

HAMELN RDS GMBH

EQ 200MG BASE/VIAL

A090663 001 Sep 10, 2012

EQ 1GM BASE/VIAL

A090663 002 Sep 10, 2012

MYLAN LABS LTD

EQ 200MG BASE/VIAL

A200145 001 Jul 25, 2011

EQ 1GM BASE/VIAL

A200145 002 Jul 25, 2011

EQ 2GM BASE/VIAL

A200145 003 Jul 25, 2011

SAGENT PHARMS

EQ 200MG BASE/VIAL

A091597 001 May 07, 2013

EQ 1GM BASE/VIAL

A091597 002 May 07, 2013

TEVA PHARMS

EQ 200MG BASE/VIAL

A077983 002 Jan 25, 2011

EQ 1GM BASE/VIAL

A077983 001 Jan 25, 2011

GEMZAR

+ LILLY

EQ 200MG BASE/VIAL **

N020509 001 May 15, 1996

+

EQ 1GM BASE/VIAL **

N020509 002 May 15, 1996

SOLUTION; INTRAVENOUS

INFUGEM

+ SUN PHARM

EQ 1200MG BASE/120ML (EQ 10MG BASE/ML)

N208313 001 Jul 16, 2018

+

EQ 1300MG BASE/130ML (EQ 10MG BASE/ML)

N208313 002 Jul 16, 2018

+

EQ 1400MG BASE/140ML (EQ 10MG BASE/ML)

N208313 003 Jul 16, 2018

+

EQ 1500MG BASE/150ML (EQ 10MG BASE/ML)

N208313 004 Jul 16, 2018

+

EQ 1600MG BASE/160ML (EQ 10MG BASE/ML)

N208313 005 Jul 16, 2018

+

EQ 1700MG BASE/170ML (EQ 10MG BASE/ML)

N208313 006 Jul 16, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GEMCITABINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

INFUGEM

+	EQ 1800MG BASE/180ML (EQ 10MG BASE/ML)	N208313 007	Jul 16, 2018
+	EQ 1900MG BASE/190ML (EQ 10MG BASE/ML)	N208313 008	Jul 16, 2018
+	EQ 2000MG BASE/200ML (EQ 10MG BASE/ML)	N208313 009	Jul 16, 2018
+	EQ 2200MG BASE/220ML (EQ 10MG BASE/ML)	N208313 010	Jul 16, 2018

GEMFIBROZIL

CAPSULE; ORAL

GEMFIBROZIL

MYLAN

300MG

A073466 001 Jan 25, 1993

PUREPAC PHARM

300MG

A072929 001 Jan 29, 1993

LOPID

PFIZER PHARMS

200MG

N018422 001

300MG

N018422 002

TABLET; ORAL

GEMFIBROZIL

CADILA

600MG

A204189 001 Aug 28, 2018

CHARTWELL RX

600MG

A074615 001 Sep 29, 1995

HIKMA PHARMS

600MG

A078599 001 Aug 16, 2010

MYLAN

600MG

A074452 001 Feb 16, 1995

PUREPAC PHARM

600MG

A074360 001 Aug 31, 1994

SUN PHARM INDS INC

600MG

A079239 001 Dec 29, 2008

TEVA

600MG

A074256 001 Oct 31, 1993

WATSON LABS

600MG

A074156 001 Oct 24, 1994

600MG

A074442 001 Apr 28, 1995

GEMIFLOXACIN MESYLATE

TABLET; ORAL

FACTIVE

+	LG CHEM LTD	EQ 320MG BASE **	N021158 001	Apr 04, 2003
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GEMIFLOXACIN MESYLATE

ORBION PHARMS

EQ 320MG BASE

A090466 001 Jun 15, 2015

GENTAMICIN SULFATE

CREAM; TOPICAL

GARAMYCIN

SCHERING

EQ 0.1% BASE **

A060462 001

GENTAFAIR

PHARMAFAIR

EQ 0.1% BASE

A062458 001 Sep 01, 1983

GENTAMICIN SULFATE

ALPHARMA US PHARMS

EQ 0.1% BASE

A062471 001 Sep 27, 1983

FOUGERA PHARMS INC

EQ 0.1% BASE

A062531 001 Jul 05, 1984

PHARMADERM

EQ 1MG BASE/GM

A062530 001 Jul 05, 1984

TARO

EQ 0.1% BASE

A062427 001 May 26, 1983

INJECTABLE; INJECTION

APOGEN

KING PHARMS

EQ 10MG BASE/ML

A062289 001

EQ 40MG BASE/ML

A062289 002

BRISTAGEN

BRISTOL

EQ 40MG BASE/ML

A062288 001

GARAMYCIN

SCHERING

EQ 1MG BASE/ML **

A061716 002

EQ 10MG BASE/ML **

A061739 001

EQ 40MG BASE/ML **

A061716 001

GENTAFAIR

PHARMAFAIR

EQ 40MG BASE/ML

A062493 001 Aug 28, 1985

GENTAMICIN

INTL MEDICATION

EQ 1MG BASE/ML

A062325 003 Jun 23, 1982

EQ 40MG BASE/ML

A062325 001

EQ 100MG BASE/100ML

A062325 004 Jun 23, 1982

GENTAMICIN SULFATE

ABBOTT

EQ 1.2MG BASE/ML

A062413 001 Aug 11, 1983

EQ 1.4MG BASE/ML

A062413 002 Aug 11, 1983

EQ 1.6MG BASE/ML

A062413 003 Aug 11, 1983

EQ 1.8MG BASE/ML

A062413 004 Aug 11, 1983

EQ 2MG BASE/ML

A062413 005 Aug 11, 1983

EQ 60MG BASE/100ML

A062413 006 Aug 11, 1983

EQ 70MG BASE/100ML

A062413 007 Aug 11, 1983

EQ 80MG BASE/100ML

A062413 008 Aug 11, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GENTAMICIN SULFATE

	EQ 90MG BASE/100ML	A062413 009	Aug 11, 1983
	EQ 100MG BASE/100ML	A062413 010	Aug 11, 1983
FRESENIUS KABI USA	EQ 10MG BASE/ML	A062356 001	Mar 04, 1982
	EQ 40MG BASE/ML	A062356 002	Mar 04, 1982
HOSPIRA	EQ 10MG BASE/ML	A062612 004	Feb 20, 1986
KALAPHARM	EQ 40MG BASE/ML	A062354 001	Apr 05, 1982
PHARM SPEC	EQ 40MG BASE/ML	A062340 001	Mar 28, 1983
SOLOPAK	EQ 10MG BASE/ML	A062507 001	Jun 06, 1985
	EQ 40MG BASE/ML	A062507 002	Jun 06, 1985
TEVA PARENTERAL	EQ 10MG BASE/ML	A063149 001	Nov 21, 1991
	EQ 40MG BASE/ML	A063106 002	Nov 21, 1991
WATSON LABS	EQ 10MG BASE/ML	A062318 002	
	EQ 40MG BASE/ML	A062318 001	
WYETH AYERST	EQ 10MG BASE/ML	A062264 001	
	EQ 40MG BASE/ML	A062264 002	
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	EQ 0.8MG BASE/ML	A062814 001	Aug 28, 1987
	EQ 1.2MG BASE/ML	A062814 002	Aug 28, 1987
	EQ 1.4MG BASE/ML	A062814 003	Aug 28, 1987
	EQ 1.6MG BASE/ML	A062814 004	Aug 28, 1987
	EQ 1.8MG BASE/ML	A062814 005	Aug 28, 1987
	EQ 2MG BASE/ML	A062814 006	Aug 28, 1987
	EQ 2.4MG BASE/ML	A062814 007	Aug 28, 1987
	EQ 40MG BASE/100ML	A062814 008	Aug 28, 1987
	EQ 60MG BASE/100ML	A062814 009	Aug 28, 1987
	EQ 70MG BASE/100ML	A062814 010	Aug 28, 1987
	EQ 80MG BASE/100ML	A062814 011	Aug 28, 1987
	EQ 90MG BASE/100ML	A062814 012	Aug 28, 1987
	EQ 100MG BASE/100ML	A062814 013	Aug 28, 1987
	EQ 120MG BASE/100ML	A062814 014	Aug 28, 1987
BAXTER HLTHCARE	EQ 0.8MG BASE/ML	A062373 001	Sep 07, 1982
	EQ 2.4MG BASE/ML	A062373 010	Sep 07, 1982
	EQ 40MG BASE/100ML	A062373 003	Sep 07, 1982
	EQ 60MG BASE/100ML	A062373 004	Sep 07, 1982
HOSPIRA	EQ 1.2MG BASE/ML	A062588 001	Jan 06, 1986
	EQ 1.4MG BASE/ML	A062414 002	Aug 15, 1983
	EQ 1.4MG BASE/ML	A062588 002	Jan 06, 1986
	EQ 1.6MG BASE/ML	A062588 003	Jan 06, 1986
	EQ 1.8MG BASE/ML	A062414 004	Aug 15, 1983
	EQ 1.8MG BASE/ML	A062588 004	Jan 06, 1986
	EQ 2MG BASE/ML	A062414 005	Aug 15, 1983
	EQ 2MG BASE/ML	A062588 005	Jan 06, 1986
	EQ 60MG BASE/100ML	A062414 006	Aug 15, 1983
	EQ 60MG BASE/100ML	A062588 006	Jan 06, 1986
	EQ 70MG BASE/100ML	A062414 007	Aug 15, 1983
	EQ 70MG BASE/100ML	A062588 007	Jan 06, 1986
	EQ 80MG BASE/100ML	A062588 008	Jan 06, 1986
	EQ 90MG BASE/100ML	A062414 009	Aug 15, 1983
	EQ 90MG BASE/100ML	A062588 009	Jan 06, 1986
	EQ 100MG BASE/100ML	A062588 010	Jan 06, 1986
U-GENCIN			
PHARMACIA AND UPJOHN	EQ 10MG BASE/ML	A062248 001	
	EQ 40MG BASE/ML	A062248 002	
INJECTABLE; INTRATHECAL			
GARAMYCIN			
+ SCHERING	EQ 2MG BASE/ML **	N050505 001	
OINTMENT; OPHTHALMIC			
GARAMYCIN			
SCHERING	EQ 0.3% BASE	N050425 001	
GENTACIDIN			
NOVARTIS	EQ 0.3% BASE	A062501 001	Jul 26, 1984
GENTAFAIR			
PHARMAFAIR	EQ 3MG BASE/GM	A062443 001	May 26, 1983
GENTAMICIN SULFATE			
AKORN	EQ 0.3% BASE	A064093 001	Aug 31, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

OINTMENT;TOPICAL

GARAMYCIN

SCHERING

EQ 0.1% BASE **

A060463 001

GENTAFAIR

PHARMAFAIR

EQ 0.1% BASE

A062444 001 May 26, 1983

GENTAMICIN SULFATE

ALPHARMA US PHARMS

EQ 0.1% BASE

A062496 001 Mar 14, 1984

PHARMADERM

EQ 0.1% BASE

A062534 001 Oct 10, 1984

VERTICE

EQ 0.1% BASE

A209233 001 Dec 31, 2018

SOLUTION/DROPS;OPHTHALMIC

GARAMYCIN

+ SCHERING

EQ 0.3% BASE **

N050039 002

GENOPTIC

ALLERGAN

EQ 0.3% BASE

A062452 001 Oct 10, 1984

GENTACIDIN

NOVARTIS

EQ 0.3% BASE

A062480 001 Mar 30, 1984

GENTAFAIR

PHARMAFAIR

EQ 0.3% BASE

A062440 001 May 03, 1983

GENTAK

AKORN

EQ 0.3% BASE

A064163 001 Oct 12, 2001

GENTAMICIN SULFATE

AKORN

EQ 0.3% BASE

A062635 001 Jan 08, 1987

ALCON PHARMS LTD

EQ 0.3% BASE

A062523 001 Nov 25, 1985

PACO

EQ 3MG BASE/ML

A062932 001 Nov 07, 1988

GENTAMICIN SULFATE; PREDNISOLONE ACETATE

OINTMENT;OPHTHALMIC

PRED-G

+ ALLERGAN

EQ 0.3% BASE;0.6%

N050612 001 Dec 01, 1989

SUSPENSION/DROPS;OPHTHALMIC

PRED-G

+ ALLERGAN

EQ 0.3% BASE;1%

N050586 001 Jun 10, 1988

GENTIAN VIOLET

SUPPOSITORY;VAGINAL

GVS

SAVAGE LABS

0.4%

A083513 001

TAMPON;VAGINAL

GENAPAX

KEY PHARMS

5MG

A085017 001

GLATIRAMER ACETATE

FOR SOLUTION;SUBCUTANEOUS

COPAXONE

+ TEVA PHARMS USA

20MG/VIAL

N020622 001 Dec 20, 1996

GLIMEPIRIDE

TABLET;ORAL

GLIMEPIRIDE

ACTAVIS LABS FL INC

1MG

A076995 001 Apr 27, 2010

2MG

A076995 002 Apr 27, 2010

4MG

A076995 003 Apr 27, 2010

EPIC PHARMA LLC

1MG

A077274 001 Oct 06, 2005

2MG

A077274 002 Oct 06, 2005

4MG

A077274 003 Oct 06, 2005

HIKMA PHARMS

1MG

A078952 001 Aug 01, 2013

2MG

A078952 002 Aug 01, 2013

4MG

A078952 003 Aug 01, 2013

MYLAN

1MG

A077486 001 Feb 10, 2006

1MG

A077624 001 Nov 28, 2005

2MG

A077486 002 Feb 10, 2006

2MG

A077624 002 Nov 28, 2005

4MG

A077486 003 Feb 10, 2006

4MG

A077624 003 Nov 28, 2005

RANBAXY

3MG

A077366 001 Oct 06, 2005

6MG

A077366 002 Oct 06, 2005

RANBAXY LABS LTD

1MG

A076875 001 Oct 06, 2005

2MG

A076875 002 Oct 06, 2005

4MG

A076875 003 Oct 06, 2005

8MG

A076875 004 Oct 06, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

TEVA	1MG	A076802 001	Oct 06, 2005
	2MG	A076802 002	Oct 06, 2005
	4MG	A076802 003	Oct 06, 2005
WATSON LABS	1MG	A077280 001	Feb 03, 2006
	2MG	A077280 002	Feb 03, 2006
	4MG	A077280 003	Feb 03, 2006

GLIMEPIRIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDARYL

+ SB PHARMCO	1MG; 4MG **	N021700 001	Nov 23, 2005
+	2MG; 4MG **	N021700 002	Nov 23, 2005
+	2MG; 8MG **	N021700 004	Mar 30, 2007
+	4MG; 4MG **	N021700 003	Nov 23, 2005
+	4MG; 8MG **	N021700 005	Mar 30, 2007

ROSIGLITAZONE MALEATE AND GLIMEPIRIDE

TEVA PHARMS USA	1MG; 4MG	A078709 001	Apr 01, 2016
	2MG; 4MG	A078709 002	Apr 01, 2016
	2MG; 8MG	A078709 004	Apr 01, 2016
	4MG; 4MG	A078709 003	Apr 01, 2016
	4MG; 8MG	A078709 005	Apr 01, 2016

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

ANI PHARMS	5MG	A074387 001	Mar 04, 1996
	10MG	A074387 002	Mar 04, 1996
BARR LABS INC	5MG	A074619 001	Apr 04, 1997
	10MG	A074619 002	Apr 04, 1997
CHARTWELL RX	5MG	A074542 001	Jun 20, 1995
	10MG	A074542 002	Jun 20, 1995
MYLAN	5MG	A074438 001	Jun 20, 1995
	10MG	A074438 002	Jun 20, 1995
OXFORD PHARMS	5MG	A074378 001	Nov 28, 1994
	10MG	A074378 002	Nov 28, 1994
SUN PHARM INDS INC	5MG	A077820 001	Jul 11, 2006
	10MG	A077820 002	Jul 11, 2006
WATSON LABS	5MG	A074370 001	Nov 22, 1994
	10MG	A074370 002	Nov 22, 1994

GLUCOTROL

+ PFIZER	2.5MG **	N017783 003	May 11, 1993
+	5MG	N017783 001	May 08, 1984
+	10MG	N017783 002	May 08, 1984

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

MYLAN	2.5MG	A202298 001	May 19, 2015
	5MG	A202298 002	May 19, 2015
	10MG	A202298 003	May 19, 2015
PAR PHARM	5MG	A076159 002	Sep 20, 2013
	10MG	A076159 001	Sep 20, 2013

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

DASH PHARMS	2.5MG; 250MG	A078083 001	Apr 12, 2007
	2.5MG; 500MG	A078083 002	Apr 12, 2007
	5MG; 500MG	A078083 003	Apr 12, 2007
SUN PHARM INDS INC	2.5MG; 250MG	A077620 001	Jan 11, 2008
	2.5MG; 500MG	A077620 002	Jan 11, 2008
	5MG; 500MG	A077620 003	Jan 11, 2008

METAGLIP

+ BRISTOL MYERS SQUIBB	2.5MG; 250MG **	N021460 001	Oct 21, 2002
+	2.5MG; 500MG **	N021460 002	Oct 21, 2002
+	5MG; 500MG **	N021460 003	Oct 21, 2002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLUCAGON

INJECTABLE; INJECTION

GLUCAGON

+ LILLY

1MG/VIAL

N020928 001 Sep 11, 1998

SOLUTION; SUBCUTANEOUS

GVOKE PFS

+ XERIS

0.5MG/0.1ML (0.5MG/0.1ML)

N212097 001 Sep 10, 2019

GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGON

+ LILLY

EQ 1MG BASE/VIAL **

N012122 001

+

EQ 10MG BASE/VIAL **

N012122 002

GLUTETHIMIDE

CAPSULE; ORAL

DORIDEN

SANOFI AVENTIS US

500MG

N009519 008

TABLET; ORAL

DORIDEN

SANOFI AVENTIS US

250MG

N009519 002

500MG

N009519 005

GLUTETHIMIDE

HALSEY

250MG

A089458 001 Oct 10, 1986

500MG

A089459 001 Oct 10, 1986

LANNETT

250MG

A083475 001

500MG

A085571 001

UCB INC

500MG

A085171 001

UPSHER SMITH LABS

500MG

A083234 002

VITARINE

500MG

A087297 001

WATSON LABS

500MG

A084362 001

500MG

A085763 001

GLYBURIDE

TABLET; ORAL

GLYBURIDE

ACTAVIS ELIZABETH

1.5MG

A075947 001 Nov 14, 2002

3MG

A075947 002 Nov 14, 2002

6MG

A075947 003 Nov 14, 2002

AUROBINDO PHARMA

1.25MG

A077537 001 Oct 18, 2007

2.5MG

A077537 002 Oct 18, 2007

5MG

A077537 003 Oct 18, 2007

CHARTWELL RX

1.25MG

A203581 001 Apr 14, 2016

2.5MG

A203581 002 Apr 14, 2016

5MG

A203581 003 Apr 14, 2016

GLYBURIDE (MICRONIZED)

CHARTWELL RX

1.5MG

A075174 001 Jun 22, 1998

3MG

A075174 002 Jun 22, 1998

HIKMA

1.5MG

A075890 001 Jul 31, 2003

3MG

A075890 002 Jul 31, 2003

6MG

A075890 003 Jul 31, 2003

MYLAN

1.5MG

A074792 001 Jun 26, 1998

3MG

A074792 002 Jun 26, 1998

6MG

A074792 003 Aug 17, 1999

SANOFI AVENTIS US

1.5MG

N020055 001 Apr 17, 1992

3MG

N020055 002 Apr 17, 1992

6MG

N020055 003 Mar 08, 2000

STRIDES PHARMA

1.5MG

A074591 001 Dec 22, 1997

3MG

A074591 002 Dec 22, 1997

4.5MG

A074591 003 Dec 22, 1997

6MG

A074591 004 Dec 22, 1997

GLYNASE

+ PFIZER

4.5MG **

N020051 003 Sep 24, 1993

MICRONASE

+ PFIZER

1.25MG **

N017498 001 May 01, 1984

2.5MG

N017498 002 May 01, 1984

+

5MG **

N017498 003 May 01, 1984

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOVANCE

+	BRISTOL MYERS SQUIBB	1.25MG;250MG **	N021178	001	Jul 31, 2000
+		2.5MG;500MG **	N021178	002	Jul 31, 2000
+		5MG;500MG **	N021178	003	Jul 31, 2000

GLYBURIDE AND METFORMIN HYDROCHLORIDE

	HERITAGE PHARMS	1.25MG;250MG	A079009	001	Jun 03, 2009
		2.5MG;500MG	A079009	002	Jun 03, 2009
		5MG;500MG	A079009	003	Jun 03, 2009
	IMPAX LABS INC	1.25MG;250MG	A076731	001	Nov 19, 2004
		2.5MG;500MG	A076731	002	Nov 19, 2004
		5MG;500MG	A076731	003	Nov 19, 2004
	TEVA	1.25MG;250MG	A076821	001	Jan 27, 2005
		2.5MG;500MG	A076821	002	Jan 27, 2005
		5MG;500MG	A076821	003	Jan 27, 2005

GLYCEROL PHENYLBUTYRATE

LIQUID; ORAL

GLYCEROL PHENYLBUTYRATE

	PAR PHARM INC	1.1GM/ML	A205742	001	Dec 02, 2021
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GLYCINE

SOLUTION; IRRIGATION

GLYCINE 1.5% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	1.5GM/100ML	N018522	001	Feb 19, 1982
	HOSPIRA	1.5GM/100ML	N017633	001	

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

	ABRAXIS PHARM	0.2MG/ML	A088475	001	Jun 12, 1984
	EUGIA PHARMA	0.2MG/ML	A210244	001	Nov 28, 2018
	HOSPIRA	0.2MG/ML	A089393	001	Jun 15, 1988
	LUPIN LTD	0.2MG/ML	A213655	001	Feb 07, 2023
	MANKIND PHARMA	0.2MG/ML	A217354	001	Feb 27, 2023
	TEVA PARENTERAL	0.2MG/ML	A081169	001	Sep 10, 1991
	WATSON LABS	0.2MG/ML	A086947	001	Jun 24, 1983
	ZYDUS PHARMS	0.2MG/ML	A214213	001	Nov 09, 2021

ROBINUL

+	HIKMA	0.2MG/ML **	N017558	001	
	ROBINS AH	0.2MG/ML	N014764	001	

POWDER; INHALATION

SEEBRI

+	NOVARTIS	15.6MCG/INH	N207923	001	Oct 29, 2015
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SOLUTION; INHALATION

LONHALA MAGNAIR KIT

+	SUMITOMO PHARMA AM	25MCG/ML	N208437	001	Dec 05, 2017
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TABLET; ORAL

GLYCOPYRROLATE

	CHARTWELL RX	1MG	A040568	001	Dec 22, 2004
		2MG	A040568	002	Dec 22, 2004
	HIKMA INTL PHARMS	1MG	A040836	001	Mar 05, 2009
		2MG	A040836	002	Mar 05, 2009
	WATSON LABS	1MG	A085562	001	
		1MG	A086902	001	
		2MG	A085563	001	
		2MG	A086178	001	
		2MG	A086900	001	

TABLET, ORALLY DISINTEGRATING; ORAL

DARTISLA ODT

	EDENBRIDGE PHARMS	0.85MG	N215019	002	Oct 11, 2023
+		1.7MG	N215019	001	Dec 16, 2021

GLYCOPYRROLATE; INDACATEROL MALEATE

POWDER; INHALATION

UTIBRON

+	NOVARTIS	15.6MCG/INH; 27.5MCG/INH	N207930	001	Oct 29, 2015
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GONADORELIN ACETATE

INJECTABLE; INJECTION

LUTREPULSE KIT

FERRING

0.8MG/VIAL

N019687 001 Oct 10, 1989

3.2MG/VIAL

N019687 002 Oct 10, 1989

GONADORELIN HYDROCHLORIDE

INJECTABLE; INJECTION

FACTREL

HIKMA

EQ 0.1MG BASE/VIAL

N018123 001 Sep 30, 1982

EQ 0.2MG BASE/VIAL

N018123 002 Sep 30, 1982

EQ 0.5MG BASE/VIAL

N018123 003 Sep 30, 1982

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPTHALMIC

NEO-POLYCIN

DOW PHARM

0.025MG/ML; EQ 1.75MG BASE/ML; 10,000
UNITS/ML

A060427 001

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

IPHARM

0.025MG/ML; EQ 1.75MG BASE/ML; 10,000
UNITS/ML

A062818 001 Oct 11, 1988

WATSON LABS

0.025MG/ML; EQ 1.75MG BASE/ML; 10,000
UNITS/ML

A062788 001 Jun 11, 1987

NEOMYCIN SULFATE AND POLYMYXIN B SULFATE GRAMICIDIN

PHARMAFAIR

0.025MG/ML; EQ 1.75MG BASE/ML; 10,000
UNITS/ML

A062383 001 Aug 31, 1982

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

AKORN

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A079119 001 Sep 10, 2009

AM REGENT

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A091274 001 Sep 22, 2010

BAXTER HLTHCARE CORP

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A078197 001 Dec 31, 2007

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A078198 001 Jun 30, 2008

EQ 4MG BASE/4ML (EQ 1MG BASE/ML)

A078198 002 Jun 30, 2008

EUGIA PHARMA

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A204238 001 Jul 06, 2016

EQ 4MG BASE/4ML (EQ 1MG BASE/ML)

A204238 002 Jul 06, 2016

MYLAN LABS LTD

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A203454 001 Apr 04, 2017

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A203454 002 Apr 04, 2017

EQ 4MG BASE/4ML (EQ 1MG BASE/ML)

A203453 001 Jan 31, 2017

SANDOZ

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A078808 001 Apr 29, 2008

TEVA PHARMS USA

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A077963 001 Jan 03, 2008

WOCKHARDT USA

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A078566 001 Feb 29, 2008

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A078564 001 Jun 30, 2008

EQ 4MG BASE/4ML (EQ 1MG BASE/ML)

A078565 001 Jun 30, 2008

YUNG SHIN PHARM

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A202647 001 Mar 06, 2020

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A202648 001 Jun 29, 2020

EQ 4MG BASE/4ML (EQ 1MG BASE/ML)

A202648 002 Jun 29, 2020

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

DR REDDYS

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A077165 001 Dec 31, 2007

KYTRIL

+ ROCHE

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **

N020239 003 Sep 17, 2004

+

EQ 1MG BASE/ML (EQ 1MG BASE/ML) **

N020239 004 Mar 11, 1994

+

EQ 3MG BASE/ML **

N020239 001 Dec 29, 1993

+

EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **

N020239 002 Mar 11, 1994

SOLUTION; ORAL

GRANISOL

INTRA SANA LABS

EQ 2MG BASE/10ML

A078334 001 Feb 28, 2008

KYTRIL

+ ROCHE

EQ 2MG BASE/10ML **

N021238 001 Jun 27, 2001

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

AUROBINDO PHARMA USA

EQ 1MG BASE

A078725 001 Jan 30, 2008

BARR

EQ 1MG BASE

A078221 001 Dec 31, 2007

EPIC PHARMA LLC

EQ 1MG BASE

A078260 001 Dec 31, 2007

HIKMA

EQ 1MG BASE

A077842 001 Dec 31, 2007

TEVA PHARMS

EQ 1MG BASE

A078080 001 Dec 31, 2007

KYTRIL

+ ROCHE

EQ 1MG BASE **

N020305 001 Mar 16, 1995

+

EQ 2MG BASE **

N020305 002 Jun 15, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GREPAFLOXACIN HYDROCHLORIDE

TABLET; ORAL

RAXAR

OTSUKA	EQ 200MG BASE	N020695 001	Nov 06, 1997
	EQ 400MG BASE	N020695 002	May 14, 1998
	EQ 600MG BASE	N020695 003	May 14, 1998

GRISEOFULVIN, MICROCRYSTALLINE

CAPSULE; ORAL

GRISACTIN

WYETH AYERST	125MG	N050051 002	
	250MG	N050051 001	

SUSPENSION; ORAL

GRIFULVIN V

+ JOHNSON AND JOHNSON	125MG/5ML **	N050448 001	
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TABLET; ORAL

GRIFULVIN V

J AND J	125MG	A060618 001	
	250MG	A060618 002	
	500MG	A060618 003	
VALEANT LUXEMBOURG	125MG	A062279 001	
	250MG **	A062279 002	

GRISACTIN

WYETH AYERST	500MG	A060212 001	
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GRISEOFULVIN, MICROSIZE

SUSPENSION; ORAL

GRIFULVIN V

VALEANT LUXEMBOURG	125MG/5ML **	A062483 001	Jan 26, 1984
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TABLET; ORAL

GRIFULVIN V

VALEANT LUXEMBOURG	500MG	A062279 003	
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GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL

GRISACTIN ULTRA

WYETH AYERST	125MG	A062178 001	
	165MG	A062438 001	Nov 17, 1983
	250MG	A062178 002	
	330MG	A062438 002	Nov 17, 1983

ULTRAGRIS-165

PLIVA	165MG	A062645 001	Jun 30, 1992
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ULTRAGRIS-330

PLIVA	330MG	A062646 001	Jun 30, 1992
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GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

FLOWTUSS

CHARTWELL RX	200MG/5ML; 2.5MG/5ML	N022424 001	May 14, 2015
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OBREDON

+ SOVEREIGN PHARMS	200MG/5ML; 2.5MG/5ML	N205474 001	Nov 14, 2014
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TABLET; ORAL

XTRELUS

+ ECI PHARMS LLC	400MG; 5MG	N208085 001	Apr 25, 2018
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GUAIFENESIN; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYCOFENIX

+ CHARTWELL RX	200MG/5ML; 2.5MG/5ML; 30MG/5ML	N022279 001	May 14, 2015
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GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE

GRANULES	600MG; 60MG	A216082 001	Aug 22, 2022
	1.2GM; 120MG	A216082 002	Aug 22, 2022

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

ANI PHARMS	EQ 4MG BASE	A074149 001	Apr 07, 1995
	EQ 4MG BASE	A074267 001	Jun 01, 1994
	EQ 8MG BASE	A074149 002	Apr 07, 1995
	EQ 8MG BASE	A074267 002	Jun 01, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

CHARTWELL RX	EQ 4MG BASE	A074517 001	Sep 30, 1998
	EQ 8MG BASE	A074517 002	Sep 30, 1998
WATSON LABS	EQ 4MG BASE	A074025 001	Feb 28, 1994
	EQ 8MG BASE	A074025 002	Feb 28, 1994
WYTENSIN			
WYETH AYERST	EQ 4MG BASE	N018587 001	Sep 07, 1982
	EQ 8MG BASE	N018587 002	Sep 07, 1982
	EQ 16MG BASE	N018587 003	Sep 07, 1982

GUANADREL SULFATE

TABLET; ORAL

HYLOREL

PHARMACIA AND UPJOHN	10MG	N018104 001	Dec 29, 1982
	25MG	N018104 002	Dec 29, 1982

GUANETHIDINE MONOSULFATE

TABLET; ORAL

GUANETHIDINE MONOSULFATE

WATSON LABS	EQ 10MG SULFATE	A086113 001	Mar 26, 1985
	EQ 25MG SULFATE	A086114 001	Mar 26, 1985
ISMELIN			
NOVARTIS	EQ 10MG SULFATE	N012329 001	
	EQ 25MG SULFATE	N012329 002	

GUANETHIDINE MONOSULFATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ESIMIL

NOVARTIS	10MG; 25MG	N013553 001	
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GUANFACINE HYDROCHLORIDE

TABLET; ORAL

GUANFACINE HYDROCHLORIDE

WATSON LABS	EQ 1MG BASE	A074762 001	Jun 25, 1997
	EQ 2MG BASE	A074762 002	Jun 25, 1997
TENEX			
+ PROMIUS PHARMA	EQ 1MG BASE **	N019032 001	Oct 27, 1986
+	EQ 2MG BASE **	N019032 002	Nov 07, 1988
	EQ 3MG BASE **	N019032 003	Nov 07, 1988

TABLET, EXTENDED RELEASE; ORAL

GUANFACINE HYDROCHLORIDE

IMPAX LABS INC	EQ 1MG BASE	A202238 001	Oct 20, 2015
	EQ 2MG BASE	A202238 002	Oct 20, 2015
	EQ 3MG BASE	A202238 003	Oct 20, 2015
	EQ 4MG BASE	A202238 004	Oct 20, 2015
MYLAN	EQ 1MG BASE	A202578 001	Jun 02, 2015
	EQ 2MG BASE	A202578 002	Jun 02, 2015
	EQ 3MG BASE	A202578 003	Jun 02, 2015
	EQ 4MG BASE	A202578 004	Jun 02, 2015

GUANIDINE HYDROCHLORIDE

TABLET; ORAL

GUANIDINE HYDROCHLORIDE

MERCK SHARP DOHME	125MG	N001546 001	
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HALAZEPAM

TABLET; ORAL

PAXIPAM

SCHERING	20MG	N017736 003	
	40MG	N017736 004	

HALCINONIDE

CREAM; TOPICAL

HALOG

WESTWOOD SQUIBB	0.025%	N017818 001	
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HALOG-E

SUN PHARM INDS INC	0.1%	N018234 001	
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OINTMENT; TOPICAL

HALOG

BRISTOL MYERS SQUIBB	0.025%	N018125 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOBETASOL PROPIONATE

CREAM;TOPICAL

ULTRAVATE

+ SUN PHARM INDS INC 0.05% **

N019967 001 Dec 27, 1990

LOTION;TOPICAL

HALOBETASOL PROPIONATE

PADAGIS ISRAEL 0.05%

A211464 001 Jun 03, 2020

OINTMENT;TOPICAL

HALOBETASOL PROPIONATE

COSETTE 0.05%

A077721 001 Sep 07, 2006

FOUGERA PHARMS 0.05%

A076903 001 Dec 16, 2004

VERTICE 0.05%

A209978 001 Mar 20, 2018

ULTRAVATE

+ SUN PHARM INDS INC 0.05% **

N019968 001 Dec 17, 1990

HALOFANTRINE HYDROCHLORIDE

TABLET;ORAL

HALFAN

GLAXOSMITHKLINE 250MG

N020250 001 Jul 24, 1992

HALOPERIDOL

TABLET;ORAL

HALDOL

+ ORTHO MCNEIL 0.5MG **

N015921 001

+ 1MG **

N015921 002

+ 2MG **

N015921 003

+ 5MG **

N015921 004

+ 10MG **

N015921 005

+ 20MG **

N015921 006 Feb 02, 1982

HALDOL SOLUTAB

ORTHO MCNEIL PHARM 1MG

N017079 001

HALOPERIDOL

ACTAVIS GROUP 0.5MG

A200854 001 Jul 01, 2022

1MG

A200854 002 Jul 01, 2022

2MG

A200854 003 Jul 01, 2022

5MG

A200854 004 Jul 01, 2022

10MG

A200854 005 Jul 01, 2022

20MG

A200854 006 Jul 01, 2022

AIPING PHARM INC 0.5MG

A071128 001 Feb 17, 1987

1MG

A071129 001 Feb 17, 1987

20MG

A071133 001 May 12, 1987

DURAMED PHARMS BARR 0.5MG

A071216 001 Dec 04, 1986

1MG

A071217 001 Dec 04, 1986

2MG

A071218 001 Dec 04, 1986

5MG

A071219 001 Dec 04, 1986

10MG

A071220 001 Jul 07, 1987

20MG

A071221 001 Jul 07, 1987

LEDERLE 0.5MG

A072727 001 Sep 19, 1989

1MG

A072728 001 Sep 19, 1989

2MG

A072729 001 Sep 19, 1989

5MG

A072730 001 Sep 19, 1989

10MG

A072731 001 Sep 19, 1989

20MG

A072732 001 Sep 19, 1989

PAR PHARM 20MG

A071328 001 Jul 20, 1987

PUREPAC PHARM 0.5MG

A071071 001 Nov 03, 1986

1MG

A071072 001 Nov 03, 1986

2MG

A071073 001 Nov 03, 1986

5MG

A071074 001 Nov 03, 1986

10MG

A071075 001 Aug 04, 1987

20MG

A071076 001 Aug 04, 1987

QUANTUM PHARMICS 0.5MG

A071255 001 Feb 17, 1987

1MG

A071269 001 Feb 17, 1987

2MG

A071256 001 Feb 17, 1987

5MG

A071257 001 Feb 17, 1987

ROYCE LABS 0.5MG

A071722 001 Dec 24, 1987

1MG

A071723 001 Dec 24, 1987

2MG

A071724 001 Dec 24, 1987

5MG

A071725 001 Dec 24, 1987

10MG

A072121 001 Dec 24, 1987

20MG

A072122 001 Dec 24, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

SANDOZ	2MG	A071209	004	Nov 17, 1986
SCS	0.5MG	A070720	001	Jun 10, 1986
	1MG	A070721	001	Jun 10, 1986
	2MG	A070722	001	Jun 10, 1986
	5MG	A070723	001	Jun 10, 1986
	10MG	A070724	001	Jun 10, 1986
	20MG	A070725	001	Sep 24, 1986
STRIDES PHARMA	0.5MG	A071235	002	Nov 03, 1986
	1MG	A071235	003	Nov 03, 1986
	2MG	A071235	001	Nov 03, 1986
	5MG	A071235	004	Nov 03, 1986
	10MG	A071235	005	Jul 20, 1987
WATSON LABS	0.5MG	A070981	001	Mar 06, 1987
	0.5MG	A071571	001	Jun 03, 1988
	1MG	A070982	001	Mar 06, 1987
	1MG	A071572	001	Jun 03, 1988
	2MG	A070983	001	Mar 06, 1987
	2MG	A071573	001	Jun 03, 1988
	5MG	A070984	001	Mar 06, 1987
	5MG	A071374	001	Jun 03, 1988
	10MG	A071375	001	Jun 03, 1988
	10MG	A072113	001	Aug 27, 1991
	20MG	A071376	001	Jun 03, 1988
	20MG	A072353	001	Aug 27, 1991

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALOPERIDOL DECANOATE

HOSPIRA	EQ 50MG BASE/ML	A075176	001	Feb 09, 2000
	EQ 100MG BASE/ML	A075176	002	Feb 09, 2000
SANDOZ	EQ 50MG BASE/ML	A076463	001	Jun 24, 2005
	EQ 100MG BASE/ML	A076463	002	Jun 24, 2005

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALDOL

+ ORTHO MCNEIL

EQ 2MG BASE/ML **

N015922 001

HALOPERIDOL

ALPHARMA	EQ 2MG BASE/ML	A070318	001	Apr 11, 1986
MORTON GROVE	EQ 2MG BASE/ML	A070710	001	Mar 07, 1986
SCS	EQ 2MG BASE/ML	A070726	001	Jun 10, 1986
TEVA	EQ 2MG BASE/ML	A071015	001	Aug 25, 1987
TEVA PHARMS	EQ 2MG BASE/ML	A071617	001	Dec 01, 1988

HALOPERIDOL INTENSOL

HIKMA

EQ 2MG BASE/ML

A072045 001 Apr 12, 1988

INJECTABLE; INJECTION

HALDOL

+ JANSSEN PHARMS

EQ 5MG BASE/ML **

N015923 001

HALOPERIDOL

ABRAXIS PHARM	EQ 5MG BASE/ML	A071187	001	Jan 20, 1987
BAXTER HLTHCARE CORP	EQ 5MG BASE/ML	A076791	001	Aug 25, 2004
	EQ 5MG BASE/ML	A076828	001	Aug 25, 2004
EPIC PHARMA LLC	EQ 5MG BASE/ML	A204849	001	Sep 06, 2017
FOSUN PHARMA	EQ 5MG BASE/ML	A076464	001	Sep 29, 2004
MARSAM PHARMS LLC	EQ 5MG BASE/ML	A072516	001	Feb 25, 1993
	EQ 5MG BASE/ML	A072517	001	Feb 25, 1993
SMITH AND NEPHEW	EQ 5MG BASE/ML	A070802	001	Dec 14, 1987
SOLOPAK	EQ 5MG BASE/ML	A070800	001	Dec 14, 1987
	EQ 5MG BASE/ML	A070801	001	Dec 14, 1987
	EQ 5MG BASE/ML	A070864	001	Dec 14, 1987
TEVA PHARMS USA	EQ 5MG BASE/ML	A076035	001	Aug 29, 2001
WATSON LABS	EQ 5MG BASE/ML	A070713	001	May 17, 1988
	EQ 5MG BASE/ML	A070714	001	May 17, 1988
	EQ 5MG BASE/ML	A070744	001	May 17, 1988

SOLUTION; ORAL

HALOPERIDOL LACTATE

ACTAVIS MID ATLANTIC EQ 1MG BASE/ML

A074536 001 Nov 02, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOPROGIN

CREAM; TOPICAL

HALOTEX

WESTWOOD SQUIBB 1% N016942 001

SOLUTION; TOPICAL

HALOTEX

WESTWOOD SQUIBB 1% N016943 001

HALOTHANE

LIQUID; INHALATION

FLUOTHANE

WYETH AYERST 99.99% N011338 001

HALOTHANE

BH 99.99% A084977 001

HALOCARBON 99.99% A080810 001

HOSPIRA 99.99% A083254 001

HEPARIN CALCIUM

INJECTABLE; INJECTION

CALCIPARINE

SANOFI AVENTIS US 25,000 UNITS/ML N018237 001

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN LOCK FLUSH

HOSPIRA 100 UNITS/ML N005264 010

INTL MEDICATION 10 UNITS/ML A086357 001

500 UNITS/ML A086357 002

LUITPOLD 10 UNITS/ML A089063 001 Oct 09, 1985

100 UNITS/ML A089064 001 Oct 09, 1985

PARKE DAVIS 10 UNITS/ML N017346 006

SMITH AND NEPHEW 10 UNITS/ML A087904 001 Apr 20, 1983

10 UNITS/ML A087958 001 Apr 20, 1983

10 UNITS/ML A088458 001 Jul 26, 1984

10 UNITS/ML A088580 001 Oct 25, 1984

100 UNITS/ML A087906 001 Apr 20, 1983

100 UNITS/ML A087959 001 Apr 20, 1983

100 UNITS/ML A088460 001 Jul 26, 1984

100 UNITS/ML A088581 001 Oct 25, 1984

SOLOPAK 10 UNITS/ML A087903 001 Apr 20, 1983

10 UNITS/ML A088457 001 Oct 25, 1984

100 UNITS/ML A087905 001 Apr 20, 1983

100 UNITS/ML A088459 001 Jul 26, 1984

HEPARIN SODIUM

ABRAXIS PHARM 1,000 UNITS/ML N017033 001

1,000 UNITS/ML N017979 001

5,000 UNITS/ML N017979 003

10,000 UNITS/ML N017979 002

CASI PHARMS INC 5,000 UNITS/ML A091659 001 Jun 08, 2011

CHAMBERLIN PARENTERL 1,000 UNITS/ML N017130 001

5,000 UNITS/ML N017130 002

10,000 UNITS/ML N017130 003

20,000 UNITS/ML N017130 004

DELL LABS 1,000 UNITS/ML N017540 001

5,000 UNITS/ML N017540 002

10,000 UNITS/ML N017540 003

20,000 UNITS/ML N017540 004

40,000 UNITS/ML N017540 005

DR REDDYS 1,000 UNITS/ML A040007 001 Jun 07, 1996

1,000 UNITS/ML N017064 002

2,500 UNITS/ML N017064 015

3,000 UNITS/ML N017064 016

4,000 UNITS/ML N017064 017

5,000 UNITS/ML N017064 003

6,000 UNITS/ML N017064 018

7,500 UNITS/ML N017064 019

10,000 UNITS/ML N017064 004

20,000 UNITS/ML N017064 005

40,000 UNITS/ML N017064 006

EPIC PHARMA LLC 1,000 UNITS/ML N017486 001

5,000 UNITS/ML N017486 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

		10,000 UNITS/ML	N017486	003	
		20,000 UNITS/ML	N017486	004	
		40,000 UNITS/ML	N017486	005	
	FRESENIUS KABI USA	1,000 UNITS/ML	N017651	005	
		5,000 UNITS/ML	N017029	002	
		10,000 UNITS/ML	N017651	003	
		20,000 UNITS/ML	N017651	008	
+	HIKMA	1,000 UNITS/ML **	N017007	001	
+		2,500 UNITS/ML **	N017007	007	
+		5,000 UNITS/ML **	N017007	002	
+		5,000 UNITS/0.5ML **	N017007	010	
+		7,500 UNITS/ML **	N017007	003	
+		10,000 UNITS/ML **	N017007	004	
+		15,000 UNITS/ML **	N017007	005	
+		20,000 UNITS/ML **	N017007	006	
	HOSPIRA	2,500 UNITS/ML	A088099	001	Apr 28, 1983
		10,000 UNITS/ML	A040095	001	Jul 26, 1996
	LILLY	1,000 UNITS/ML	N005521	001	
		10,000 UNITS/ML	N005521	002	
		20,000 UNITS/ML	N005521	004	
	LUITPOLD	1,000 UNITS/ML	A087452	001	Oct 31, 1983
	ORGANON USA INC	1,000 UNITS/ML	N000552	008	
		5,000 UNITS/ML	N000552	009	
		10,000 UNITS/ML	N000552	010	
	PARKE DAVIS	1,000 UNITS/ML	N017346	001	
		5,000 UNITS/ML	N017346	002	
		7,500 UNITS/ML	N017346	003	
		10,000 UNITS/ML	N017346	004	
		20,000 UNITS/ML	N017346	005	
+	PFIZER	10,000 UNITS/ML	N201370	003	Jul 21, 2011
	PHARM SPEC	1,000 UNITS/ML	N017780	001	
		5,000 UNITS/ML	N017780	002	
		10,000 UNITS/ML	N017780	003	
		20,000 UNITS/ML	N017780	004	
		40,000 UNITS/ML	N017780	005	
	PHARMACIA AND UPJOHN	1,000 UNITS/ML	N004570	001	
		5,000 UNITS/ML	N004570	002	
		10,000 UNITS/ML	N004570	003	
	SMITH AND NEPHEW	1,000 UNITS/ML	A088239	001	Jul 26, 1984
	SOLOPAK	1,000 UNITS/ML	A087043	001	
		5,000 UNITS/ML	A087077	001	
		5,000 UNITS/0.5ML	A087395	001	
		10,000 UNITS/ML	A087107	001	
		10,000 UNITS/0.5ML	A087363	001	
	WATSON LABS INC	1,000 UNITS/ML	A040008	001	Oct 10, 1995
	HEPARIN SODIUM 1,000 UNITS	IN DEXTROSE 5% IN PLASTIC CONTAINER			
	MCGAW	200 UNITS/100ML	N019130	001	Dec 31, 1984
	HEPARIN SODIUM 1,000 UNITS	IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
	B BRAUN	200 UNITS/100ML	N019042	001	Mar 29, 1985
	HEPARIN SODIUM 10,000 UNITS	AND DEXTROSE 5% IN PLASTIC CONTAINER			
	BAXTER HLTHCARE	2,000 UNITS/100ML	N018814	002	Jul 09, 1985
	HEPARIN SODIUM 10,000 UNITS	IN DEXTROSE 5%			
	HOSPIRA	10,000 UNITS/100ML	N018911	006	Jan 30, 1985
	HEPARIN SODIUM 10,000 UNITS	IN SODIUM CHLORIDE 0.45%			
	HOSPIRA	10,000 UNITS/100ML	N018911	001	Jan 30, 1985
		10,000 UNITS/100ML	N018916	005	Jan 31, 1984
	HEPARIN SODIUM 10,000 UNITS	IN SODIUM CHLORIDE 0.9%			
	HOSPIRA	10,000 UNITS/100ML	N018911	003	Jan 30, 1985
		10,000 UNITS/100ML	N018916	002	Jan 31, 1984
	HEPARIN SODIUM 12,500 UNITS	IN DEXTROSE 5%			
	HOSPIRA	5,000 UNITS/100ML	N018911	007	Jan 30, 1985
	HEPARIN SODIUM 12,500 UNITS	IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
	B BRAUN	5,000 UNITS/100ML	N019802	001	Jul 20, 1992
	HEPARIN SODIUM 12,500 UNITS	IN SODIUM CHLORIDE 0.9%			
	HOSPIRA	5,000 UNITS/100ML	N018911	005	Jan 30, 1985
		5,000 UNITS/100ML	N018916	003	Jan 31, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM 2,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
MCGAW	200 UNITS/100ML	N019130	003	Dec 31, 1984
HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	200 UNITS/100ML	N019042	002	Mar 29, 1985
HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4,000 UNITS/100ML	N018814	001	Oct 31, 1983
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
HOSPIRA	4,000 UNITS/100ML	N019805	001	Jan 25, 1989
HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5,000 UNITS/100ML	N018814	003	Jul 09, 1985
	10,000 UNITS/100ML	N018814	004	Jul 02, 1987
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5%				
HOSPIRA	5,000 UNITS/100ML	N018911	009	Jan 30, 1985
	10,000 UNITS/100ML	N018911	008	Jan 30, 1985
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N019134	001	Mar 29, 1985
HOSPIRA	5,000 UNITS/100ML	N019805	002	Jan 25, 1989
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N019802	005	Jul 20, 1992
	10,000 UNITS/100ML	N019802	002	Jul 20, 1992
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9%				
HOSPIRA	5,000 UNITS/100ML	N018911	004	Jan 30, 1985
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N019135	001	Mar 29, 1985
	5,000 UNITS/100ML	N019802	003	Jul 20, 1992
HOSPIRA	5,000 UNITS/100ML	N018916	009	Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	500 UNITS/100ML	N018609	003	Apr 28, 1982
HEPARIN SODIUM 5,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
MCGAW	1,000 UNITS/100ML	N019130	002	Dec 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.45%				
HOSPIRA	100 UNITS/ML	N018911	002	Jan 30, 1985
	100 UNITS/ML	N018916	004	Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9%				
HOSPIRA	1,000 UNITS/100ML	N018916	001	Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	1,000 UNITS/100ML	N019042	004	Mar 29, 1985
HEPARIN SODIUM PRESERVATIVE FREE				
DR REDDYS	1,000 UNITS/ML	A089464	001	Jun 03, 1986
HOSPIRA	2,000 UNITS/ML	N005264	013	Apr 07, 1986
	2,500 UNITS/ML	N005264	014	Apr 07, 1986
PHARMA SERVE NY	1,000 UNITS/ML	A086129	001	
LIPO-HEPIN				
3M	1,000 UNITS/0.5ML	N017027	001	
	1,000 UNITS/ML	N017027	006	
	5,000 UNITS/0.5ML	N017027	002	
	5,000 UNITS/ML	N017027	008	
	7,500 UNITS/0.5ML	N017027	010	
	10,000 UNITS/0.5ML	N017027	003	
	10,000 UNITS/ML	N017027	009	
	15,000 UNITS/0.5ML	N017027	011	
	20,000 UNITS/0.5ML	N017027	004	
	20,000 UNITS/ML	N017027	007	
	40,000 UNITS/ML	N017027	005	
LIQUAEMIN LOCK FLUSH				
ORGANON USA INC	100 UNITS/ML	N000552	007	
LIQUAEMIN SODIUM				
ORGANON USA INC	1,000 UNITS/ML	N000552	004	
	5,000 UNITS/ML	N000552	003	
	10,000 UNITS/ML	N000552	005	
	20,000 UNITS/ML	N000552	001	
	40,000 UNITS/ML	N000552	002	
LIQUAEMIN SODIUM PRESERVATIVE FREE				
ORGANON USA INC	1,000 UNITS/ML	N000552	011	Apr 11, 1986
	5,000 UNITS/ML	N000552	012	Apr 11, 1986
	10,000 UNITS/ML	N000552	013	Apr 11, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

PANHEPRIN

HOSPIRA

1,000 UNITS/ML

N005264 004

5,000 UNITS/ML

N005264 006

10,000 UNITS/ML

N005264 007

20,000 UNITS/ML

N005264 008

40,000 UNITS/ML

N005264 009

SODIUM HEPARIN

ABRAXIS PHARM

5,000 UNITS/ML

N017033 002

10,000 UNITS/ML

N017033 003

20,000 UNITS/ML

N017033 004

BAXTER HLTHCARE

1,000 UNITS/ML

N017036 001 Mar 04, 1988

HETACILLIN

FOR SUSPENSION; ORAL

VERSAPEN

BRISTOL

EQ 112.5MG AMPICIL/ML

A061398 001

EQ 112.5MG AMPICIL/5ML

N050060 001

EQ 112.5MG AMPICIL/ML

N050060 003

EQ 225MG AMPICIL/5ML

A061398 002

HETACILLIN POTASSIUM

CAPSULE; ORAL

VERSAPEN-K

BRISTOL

EQ 225MG AMPICIL

A061396 001

EQ 450MG AMPICIL

A061396 002

HEXACHLOROPHENE

AEROSOL; TOPICAL

SEPTISOL

VESTAL LABS

0.23%

N017424 001

TURGEX

XTTRIUM

3%

N018375 001

EMULSION; TOPICAL

HEXA-GERM

HUNTINGTON LABS

3%

N017411 001

PHISOHEX

SANOFI AVENTIS US

3%

N006882 001

3%

N008402 001

SOY-DOME

BAYER PHARMS

3%

N017405 001

TURGEX

XTTRIUM

3%

N019055 001 Nov 30, 1984

SOAP; TOPICAL

GAMOPHEN

ARBROOK

2%

N006270 003

SOLUTION; TOPICAL

DIAL

DIAL

0.25%

N017421 002

GERMA-MEDICA

HUNTINGTON LABS

1%

N017412 001

GERMA-MEDICA "MG"

HUNTINGTON LABS

0.25%

N017412 002

SEPTI-SOFT

CALGON

0.25%

N017460 001

SEPTISOL

VESTAL LABS

0.25%

N017423 001

SPONGE; TOPICAL

E-Z SCRUB

BECTON DICKINSON

450MG

N017452 001

HEXASCRUB

PROF DSPLS

3%

N018363 001

PHISO-SCRUB

SANOFI AVENTIS US

3%

N017446 001

SCRUBTEAM SURGICAL SPONGEBRUSH

3M

330MG

N017413 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEXAFLUORENIUM BROMIDE

INJECTABLE; INJECTION

MYLAXEN

MEDPOINTE PHARM HLC 20MG/ML N009789 003

HEXOCYCLIUM METHYLSULFATE

TABLET; ORAL

TRAL

ABBVIE 25MG N010599 001

HEXYLCAINE HYDROCHLORIDE

SOLUTION; TOPICAL

CYCLAINE

MERCK 5% N008472 001

HISTAMINE PHOSPHATE

INJECTABLE; INJECTION

HISTAMINE PHOSPHATE

LILLY EQ 0.1MG BASE/ML N000734 003

EQ 0.2MG BASE/ML N000734 002

EQ 1MG BASE/ML N000734 001

HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

VANTAS

+ ENDO PHARM

50MG N021732 001 Oct 12, 2004

INJECTABLE; INJECTION

SUPPRELIN

SHIRE EQ 0.2MG BASE/ML N019836 001 Dec 24, 1991

EQ 0.5MG BASE/ML N019836 002 Dec 24, 1991

EQ 1MG BASE/ML N019836 003 Dec 24, 1991

HOMATROPINE METHYLBROMIDE

TABLET; ORAL

HOMAPIN-10

MISSION PHARMA 10MG A086308 001

HOMAPIN-5

MISSION PHARMA 5MG A086309 001

TABLET, CHEWABLE; ORAL

EQUIPIN

MISSION PHARMA 3MG A086310 001

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

APOZEAL PHARMS 1.5MG/5ML; 5MG/5ML A204765 001 Mar 06, 2017

IVAX SUB TEVA PHARMS 1.5MG/5ML; 5MG/5ML A040285 001 Jul 19, 1999

NOSTRUM LABS INC 1.5MG/5ML; 5MG/5ML A210663 001 Jun 11, 2019

HYDROPANE

HALSEY 1.5MG/5ML; 5MG/5ML A088066 001 Jun 28, 1985

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

ACTAVIS ELIZABETH 1.5MG; 5MG A040295 001 Dec 01, 2000

NOVEL LABS INC 1.5MG; 5MG A091528 001 Apr 20, 2011

TUSSIGON

KING PHARMS 1.5MG; 5MG A088508 001 Jul 30, 1985

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

APRESOLINE

+ NOVARTIS

20MG/ML ** N008303 003

HYDRALAZINE HYDROCHLORIDE

ABRAXIS PHARM 20MG/ML A089532 001 Aug 11, 1987

MYLAN INSTITUTIONAL 20MG/ML A204680 001 Apr 28, 2016

SMITH AND NEPHEW 20MG/ML A088518 001 Apr 20, 1984

SOLOPAK 20MG/ML A088517 001 Aug 22, 1985

TEVA PARENTERAL 20MG/ML A040373 001 Feb 23, 2000

TABLET; ORAL

APRESOLINE

+ NOVARTIS 10MG ** N008303 004

+ 25MG ** N008303 001

+ 50MG ** N008303 002

+ 100MG ** N008303 005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

DRALZINE

TEVA	25MG	A084301	001	
HYDRALAZINE HYDROCHLORIDE				
ACTAVIS ELIZABETH	25MG	A088560	001	Oct 04, 1984
	50MG	A088649	001	Oct 18, 1984
ACTAVIS GRP PTC	10MG	A091679	001	Mar 04, 2013
	25MG	A091679	002	Mar 04, 2013
	50MG	A091679	003	Mar 04, 2013
	100MG	A091679	004	Mar 04, 2013
ANDA REPOSITORY	10MG	A089359	001	Jul 25, 1986
	25MG	A089258	001	May 05, 1986
	50MG	A089259	001	May 05, 1986
	100MG	A088729	001	Apr 11, 1985
ASCOT	25MG	A088310	001	Dec 19, 1984
	50MG	A088311	001	Dec 19, 1984
CHARTWELL RX	10MG	A088846	001	Feb 26, 1985
	25MG	A088847	001	Feb 26, 1985
	50MG	A088848	001	Feb 26, 1985
	100MG	A088849	001	Feb 26, 1985
HALSEY	10MG	A089218	001	Jan 22, 1986
	25MG	A089130	001	Jan 15, 1986
	50MG	A089222	001	Jan 22, 1986
	100MG	A089178	001	Jan 15, 1986
IMPAX LABS	25MG	A084922	001	
	50MG	A084923	001	
IVAX SUB TEVA PHARMS	10MG	A084443	001	
	25MG	A084437	001	
	50MG	A084469	002	
	100MG	A084581	001	
MUTUAL PHARM	10MG	A088728	001	Apr 11, 1985
	25MG	A084106	002	
	50MG	A084107	002	
MYLAN	10MG	A090413	001	Dec 08, 2010
	25MG	A090413	002	Dec 08, 2010
	50MG	A090413	003	Dec 08, 2010
	100MG	A090413	004	Dec 08, 2010
PUREPAC PHARM	25MG	A088177	001	Jul 29, 1983
	50MG	A088178	001	Aug 15, 1983
QUANTUM PHARMICS	10MG	A088671	001	May 01, 1984
	25MG	A088657	001	Jun 15, 1984
	50MG	A088652	001	May 08, 1984
	100MG	A088686	001	May 01, 1984
RISING	10MG	A209251	001	Jul 09, 2018
	25MG	A209251	002	Jul 09, 2018
	50MG	A209251	003	Jul 09, 2018
	100MG	A209251	004	Jul 09, 2018
STRIDES PHARMA	10MG	A200770	004	Jun 25, 2019
SUPERPHARM	10MG	A088787	001	Aug 28, 1984
	25MG	A088788	001	Aug 28, 1984
	50MG	A088789	001	Aug 28, 1984
UPSHER SMITH LABS	10MG	A083241	001	
	25MG	A083560	001	
	50MG	A083561	001	
	50MG	A085088	001	
USL PHARMA	25MG	A087780	001	Mar 29, 1982
	50MG	A087751	001	Mar 29, 1982
VANGARD	25MG	A087712	001	
	50MG	A087908	001	May 07, 1982
VITARINE	25MG	A086088	001	
WATSON LABS	25MG	A084504	001	
	25MG	A085532	002	May 24, 1982
	50MG	A084503	001	
	50MG	A085533	002	May 25, 1982
WEST WARD	25MG	A088240	001	May 27, 1983
	50MG	A088241	001	May 27, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

APRESAZIDE

+	NOVARTIS	25MG;25MG	A084735	001
+		50MG;50MG	A084810	001
		100MG;50MG	A084811	001

HYDRA-ZIDE

	STRIDES PHARMA	100MG;50MG	A088961	001	Oct 21, 1985
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HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

	SOLVAY	25MG;25MG	A087608	001	Feb 08, 1982
		50MG;50MG	A087213	001	Feb 08, 1982
		100MG;50MG	A087609	001	Feb 08, 1982
	SUPERPHARM	25MG;25MG	A089200	001	Feb 09, 1987
		50MG;50MG	A089201	001	Feb 09, 1987
	WATSON LABS	25MG;25MG	A085457	001	Mar 04, 1982
		50MG;50MG	A085446	001	Mar 04, 1982
		100MG;50MG	A085440	001	Mar 04, 1982

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 100/50

	IVAX PHARMS	100MG;50MG	A088358	001	Apr 10, 1984
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HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 25/25

	IVAX PHARMS	25MG;25MG	A088356	001	Apr 10, 1984
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HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 50/50

	IVAX PHARMS	50MG;50MG	A088357	001	Apr 10, 1984
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TABLET; ORAL

APRESOLINE-ESIDRIX

	NOVARTIS	25MG;15MG	N012026	002
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HYDRALAZINE AND HYDROCHLOROTHIAZIDE

	WATSON LABS	25MG;15MG	A085827	001
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HYDROCHLOROTHIAZIDE W/ HYDRALAZINE

	WATSON LABS	25MG;15MG	A085373	001
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HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

	CHARTWELL RX	25MG;15MG;0.1MG	A084897	001
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HYDRALAZINE HYDROCHLORIDE, HYDROCHLOROTHIAZIDE AND RESERPINE

	IVAX SUB TEVA PHARMS	25MG;15MG;0.1MG	A084291	001
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HYDRALAZINE HYDROCHLORIDE-HYDROCHLOROTHIAZIDE-RESERPINE

	MYLAN	25MG;15MG;0.1MG	A087085	001
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HYDRALAZINE, HYDROCHLOROTHIAZIDE W/ RESERPINE

	WATSON LABS	25MG;15MG;0.1MG	A085771	001
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HYDRAP-ES

	SANDOZ	25MG;15MG;0.1MG	A084876	001
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HYDROCHLOROTHIAZIDE W/ RESERPINE AND HYDRALAZINE

	WATSON LABS	25MG;15MG;0.1MG	A083770	001
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HYDROSERPINE PLUS (R-H-H)

	IVAX SUB TEVA PHARMS	25MG;15MG;0.1MG	A083877	001
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RESERPINE, HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

	SOLVAY	25MG;15MG;0.1MG	A088376	001	Oct 28, 1983
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	SUN PHARM INDUSTRIES	25MG;15MG;0.1MG	A088570	001	Apr 10, 1984
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	WATSON LABS	25MG;15MG;0.1MG	A085549	001
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		25MG;15MG;0.1MG	A087556	001
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RESERPINE, HYDROCHLOROTHIAZIDE, AND HYDRALAZINE HYDROCHLORIDE

	LEDERLE	25MG;15MG;0.1MG	A087709	001	May 13, 1982
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SER-A-GEN

	SOLVAY	25MG;15MG;0.1MG	A087210	001
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SER-AP-ES

	NOVARTIS	25MG;15MG;0.1MG	N012193	005
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UNIPRES

	SOLVAY	25MG;15MG;0.1MG	A085893	001
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		25MG;15MG;0.1MG	A086298	001
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HYDRALAZINE HYDROCHLORIDE; RESERPINE

TABLET; ORAL

DRALSERP

	SANDOZ	25MG;0.1MG	A084617	001
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SERPASIL-APRESOLINE

	NOVARTIS	25MG;0.1MG	N009296	004
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		50MG;0.2MG	N009296	002
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

ALEMBIC PHARMS LTD	12.5MG	A200645	001	Nov 30, 2010
APOTEX	12.5MG	A078389	001	May 16, 2008
CHARTWELL MOLECULAR	12.5MG	A091662	001	Jan 27, 2012
HIKMA INTL PHARMS	12.5MG	A077885	001	Nov 26, 2007
IVAX SUB TEVA PHARMS	12.5MG	A077005	001	Jul 13, 2005
MYLAN	12.5MG	A075640	001	Jan 28, 2000
SUN PHARM INDS INC	12.5MG	A090651	001	Apr 07, 2014

SOLUTION; ORAL

HYDROCHLOROTHIAZIDE

MORTON GROVE	50MG/5ML	A089661	001	Jun 20, 1988
+ ROXANE	50MG/5ML **	A088587	001	Jul 02, 1984
HYDROCHLOROTHIAZIDE INTENSOL				
ROXANE	100MG/ML	A088588	001	Jul 02, 1984

TABLET; ORAL

ESIDRIX

NOVARTIS	25MG	N011793	005	
	50MG	N011793	008	
	100MG	N011793	009	

HYDRO-D

HALSEY	25MG	A086504	001	
	50MG	A083891	002	

HYDROCHLOROTHIAZIDE

ABC HOLDING	50MG	A085672	001	
ACTAVIS ELIZABETH	25MG	A085054	002	
	50MG	A085208	001	
ALRA	25MG	A086369	001	
	50MG	A083554	001	
APOTEX	25MG	A040774	001	Oct 03, 2007
	50MG	A040774	002	Oct 03, 2007
ASCOT	25MG	A087539	001	Feb 03, 1982
	50MG	A087540	001	Feb 03, 1982
AUROLIFE PHARMA LLC	25MG	A083899	001	
	50MG	A085219	001	
BARR	50MG	A084771	001	
CHARTWELL RX	25MG	A085683	001	
	25MG	A087565	001	Mar 09, 1982
	50MG	A083965	001	
	50MG	A084912	001	
DAVA PHARMS INC	100MG	A087060	001	
ELKINS SINN	50MG	A085152	002	
HEATHER	50MG	A084135	001	
HIKMA INTL PHARMS	25MG	A084878	002	Jul 12, 2006
	50MG	A084878	001	
IMPAX LABS	25MG	A084029	001	
	50MG	A083607	002	
	100MG	A085098	001	
INWOOD LABS	25MG	A084776	001	
	25MG	A085067	001	
	50MG	A084776	002	
IVAX SUB TEVA PHARMS	50MG	A084658	001	
+	100MG	A085022	001	
JUBILANT CADISTA	25MG	A040809	001	Sep 04, 2007
	50MG	A040809	002	Sep 04, 2007
LANNETT CO INC	25MG	A084325	001	
	50MG	A084324	001	
MAST MM	25MG	A086192	001	
	50MG	A086192	002	
MYLAN	25MG	A084880	001	
	50MG	A085112	001	
MYLAN PHARMS INC	12.5MG	A040770	001	Jan 23, 2007
	25MG	A040735	002	Jan 23, 2007
	50MG	A040735	003	Jan 23, 2007
PVT FORM	50MG	A086597	001	
ROXANE	25MG	A085004	001	
	50MG	A084536	002	
	50MG	A085005	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

SOLVAY	25MG	A085323	001	
SUN PHARM INDS INC	12.5MG	A040857	001	May 30, 2008
	25MG	A040810	001	Mar 27, 2007
	50MG	A040810	002	Mar 27, 2007
SUN PHARM INDUSTRIES	25MG	A083972	001	
	50MG	A083972	002	
	100MG	A083972	003	
SUPERPHARM	25MG	A088827	001	Dec 28, 1984
	50MG	A088828	001	Dec 28, 1984
	100MG	A088829	001	Dec 28, 1984
TEVA	25MG	A088924	001	Feb 07, 1985
	50MG	A088923	001	Feb 07, 1985
USL PHARMA	25MG	A087827	001	Apr 19, 1982
	50MG	A087752	001	Apr 19, 1982
VANGARD	25MG	A087638	001	
	50MG	A087610	001	
WARNER CHILCOTT	25MG	A087586	001	May 03, 1982
	50MG	A087587	001	May 03, 1982
WATSON LABS	25MG	A081189	001	Jan 24, 1992
	25MG	A083458	001	
	25MG	A085232	002	
	50MG	A083456	001	
	50MG	A085233	001	
	50MG	A086087	001	
	50MG	A086594	001	
	100MG	A081190	001	Jan 24, 1992
	100MG	A085099	001	
	100MG	A087002	001	
WATSON LABS TEVA	50MG	A083232	001	
WEST WARD	25MG	A084899	001	
WHITEWORTH TOWN PLSN	25MG	A083809	002	
	50MG	A083809	001	
	100MG	A085347	001	
HYDRODIURIL				
+ MERCK	25MG **	N011835	003	
+	50MG **	N011835	006	
+	100MG **	N011835	007	
ORETIC				
ABBVIE	25MG	N011971	001	
	50MG	N011971	002	
ZIDE				
SOLVAY	50MG	A083925	001	

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

+ SANOFI AVENTIS US	12.5MG; 75MG **	N020758	001	Sep 30, 1997
+	25MG; 300MG **	N020758	004	Mar 15, 2005

IRBESARTAN AND HYDROCHLOROTHIAZIDE

APOTEX INC	12.5MG; 150MG	A201505	001	Oct 15, 2012
	12.5MG; 300MG	A201505	002	Oct 15, 2012
ATLAS PHARMS LLC	12.5MG; 150MG	A203036	001	Jan 15, 2016
	12.5MG; 300MG	A203036	002	Jan 15, 2016
	25MG; 300MG	A203036	003	Jan 15, 2016
MYLAN PHARMS INC	12.5MG; 150MG	A077969	001	Sep 27, 2012
	12.5MG; 300MG	A077969	002	Sep 27, 2012
	25MG; 300MG	A077969	003	Jul 20, 2016
TEVA	25MG; 300MG	A077369	003	Mar 30, 2012
UNICHEM	12.5MG; 150MG	A207018	001	Sep 19, 2017
	12.5MG; 300MG	A207018	002	Sep 19, 2017
WATSON LABS INC	12.5MG; 150MG	A091539	001	Oct 22, 2012
	12.5MG; 300MG	A091539	002	Oct 22, 2012

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; LABETALOL HYDROCHLORIDE

TABLET; ORAL

NORMOZIDE

SCHERING	25MG;100MG	N019046 001	Apr 06, 1987
	25MG;200MG	N019046 002	Apr 06, 1987
	25MG;300MG	N019046 003	Apr 06, 1987
	25MG;400MG	N019046 004	Apr 06, 1987

TRANDATE HCT

GLAXOSMITHKLINE	25MG;100MG	N019174 001	Apr 10, 1987
	25MG;200MG	N019174 002	Apr 10, 1987
	25MG;300MG	N019174 003	Apr 10, 1987
	25MG;400MG	N019174 004	Apr 10, 1987

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	12.5MG;10MG	A075776 001	Jul 01, 2002
	12.5MG;20MG	A075776 002	Jul 01, 2002
	25MG;20MG	A075776 003	Jul 01, 2002
COREPHARMA	12.5MG;10MG	A076674 001	Oct 05, 2004
	12.5MG;20MG	A076674 002	Oct 05, 2004
	25MG;20MG	A076674 003	Oct 05, 2004
EPIC PHARMA LLC	12.5MG;10MG	A075926 001	Jul 01, 2002
	12.5MG;20MG	A075926 002	Jul 01, 2002
	25MG;20MG	A075926 003	Jul 01, 2002
HIKMA INTL PHARMS	12.5MG;10MG	A076265 001	Jul 08, 2002
	12.5MG;20MG	A076265 002	Jul 08, 2002
	25MG;20MG	A076265 003	Jul 08, 2002
MYLAN	12.5MG;10MG	A076113 001	Jul 01, 2002
	12.5MG;20MG	A076113 002	Jul 01, 2002
	25MG;20MG	A076113 003	Jul 01, 2002
TEVA	12.5MG;10MG	A075869 001	Jul 01, 2002
	12.5MG;20MG	A075869 002	Jul 01, 2002
	25MG;20MG	A075869 003	Jul 01, 2002

PRINZIDE

+	MERCK	12.5MG;10MG **	N019778 003	Nov 18, 1993
+		12.5MG;20MG **	N019778 001	Feb 16, 1989
+		25MG;20MG **	N019778 002	Feb 16, 1989

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

APOTEX	12.5MG;50MG	A090150 001	Oct 06, 2010
	12.5MG;100MG	A090150 002	Aug 11, 2010
	25MG;100MG	A090150 003	Oct 06, 2010
HIKMA	12.5MG;50MG	A077732 002	Oct 06, 2010
	12.5MG;100MG	A077732 001	Apr 06, 2010
	25MG;100MG	A077732 003	Oct 06, 2010
MYLAN	12.5MG;50MG	A091652 001	Oct 06, 2010
	12.5MG;100MG	A091652 002	Apr 06, 2010
	25MG;100MG	A091652 003	Oct 06, 2010
TORRENT PHARMS	12.5MG;50MG	A090528 001	Oct 06, 2010
	12.5MG;100MG	A090528 003	Apr 06, 2010
	25MG;100MG	A090528 002	Oct 06, 2010
WATSON LABS	12.5MG;50MG	A200180 001	Jan 12, 2011
	12.5MG;100MG	A200180 002	Jan 12, 2011
	25MG;100MG	A200180 003	Jan 12, 2011

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDORIL 15

MERCK	15MG;250MG	N013402 001	
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ALDORIL 25

MERCK	25MG;250MG	N013402 002	
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ALDORIL D30

MERCK	30MG;500MG	N013402 003	
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ALDORIL D50

MERCK	50MG;500MG	N013402 004	
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METHYLDOPA AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	15MG;250MG	A070182 001	Jan 15, 1986
	25MG;250MG	A070183 001	Jan 15, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

	30MG; 500MG	A070543	001	Jan 15, 1986
	50MG; 500MG	A070544	001	Jan 15, 1986
DAVA PHARMS INC	15MG; 250MG	A072507	001	Jun 02, 1989
	25MG; 250MG	A072508	001	Jun 02, 1989
	30MG; 500MG	A072509	001	Jun 02, 1989
	50MG; 500MG	A072510	001	Jun 02, 1989
IVAX SUB TEVA PHARMS	15MG; 250MG	A071458	001	Mar 08, 1988
	25MG; 250MG	A071459	001	Mar 08, 1988
	30MG; 500MG	A071460	001	Mar 08, 1988
	50MG; 500MG	A071461	001	Mar 08, 1988
PARKE DAVIS	15MG; 250MG	A071897	001	Nov 23, 1987
	25MG; 250MG	A071898	001	Nov 23, 1987
	30MG; 500MG	A071899	001	Nov 23, 1987
	50MG; 500MG	A071900	001	Nov 23, 1987
PUREPAC PHARM	15MG; 250MG	A070853	001	Oct 08, 1986
	25MG; 250MG	A070688	001	Apr 24, 1986
	30MG; 500MG	A070854	001	Oct 08, 1986
	50MG; 500MG	A070689	001	Apr 24, 1986
RISING	15MG; 250MG	A070265	002	Jan 23, 1986
	25MG; 250MG	A070265	001	Jan 23, 1986
SANDOZ	15MG; 250MG	A070829	001	Mar 09, 1987
	25MG; 250MG	A070830	001	Mar 09, 1987
STRIDES PHARMA	15MG; 250MG	A070616	001	Feb 02, 1987
	25MG; 250MG	A070612	001	Feb 02, 1987
	30MG; 500MG	A070613	001	Feb 02, 1987
	50MG; 500MG	A070614	001	Feb 02, 1987
TEVA	15MG; 250MG	A071819	001	Apr 08, 1988
	25MG; 250MG	A071820	001	Apr 08, 1988
	30MG; 500MG	A071821	001	Apr 08, 1988
	50MG; 500MG	A071822	001	Apr 08, 1988
WATSON LABS	15MG; 250MG	A070365	001	Mar 19, 1986
	15MG; 250MG	A070958	001	Feb 06, 1989
	15MG; 250MG	A071920	001	Aug 29, 1988
	25MG; 250MG	A070366	001	Apr 16, 1986
	25MG; 250MG	A070959	001	Jan 19, 1989
	25MG; 250MG	A071921	001	Aug 29, 1988
	30MG; 500MG	A070367	001	Mar 19, 1986
	30MG; 500MG	A071069	001	Jan 19, 1989
	30MG; 500MG	A071922	001	Aug 29, 1988
	50MG; 500MG	A070368	001	Apr 16, 1986
	50MG; 500MG	A070960	001	Feb 06, 1989
	50MG; 500MG	A071923	001	Aug 29, 1988

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DUTOPROL

+	CONCORDIA	12.5MG; EQ 25MG TARTRATE **	N021956	001	Aug 28, 2006
+		12.5MG; EQ 50MG TARTRATE **	N021956	002	Aug 28, 2006
+		12.5MG; EQ 100MG TARTRATE **	N021956	003	Aug 28, 2006

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

+	VALIDUS PHARMS	25MG; 100MG **	N018303	002	Dec 31, 1984
+		50MG; 100MG **	N018303	003	Dec 31, 1984

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	12.5MG; 7.5MG	A090096	001	Sep 25, 2008
	12.5MG; 15MG	A090096	002	Sep 25, 2008
	25MG; 15MG	A090096	003	Sep 25, 2008
HERITAGE PHARMS	12.5MG; 7.5MG	A202150	001	Mar 07, 2014
	12.5MG; 15MG	A202150	002	Mar 07, 2014
	25MG; 15MG	A202150	003	Mar 07, 2014
UNIRETIC				
UCB INC	12.5MG; 7.5MG **	N020729	001	Jun 27, 1997
	12.5MG; 15MG **	N020729	003	Feb 14, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDETABLET; ORAL
UNIRETIC

25MG; 15MG **

N020729 002 Jun 27, 1997

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

NATCO PHARMA USA 12.5MG; 20MG
12.5MG; 40MG
25MG; 40MGA078827 001 Oct 26, 2016
A078827 002 Oct 26, 2016
A078827 003 Oct 26, 2016TEVA PHARMS USA 12.5MG; 20MG
25MG; 40MGA200532 001 Apr 24, 2017
A200532 003 Apr 24, 2017TORRENT 12.5MG; 20MG
12.5MG; 40MG
25MG; 40MGA206515 001 May 03, 2017
A206515 002 May 03, 2017
A206515 003 May 03, 2017ZYDUS PHARMS 12.5MG; 20MG
12.5MG; 40MG
25MG; 40MGA206377 001 Feb 24, 2023
A206377 002 Feb 24, 2023
A206377 003 Feb 24, 2023HYDROCHLOROTHIAZIDE; PINDOLOL

TABLET; ORAL

VISKAZIDE

NOVARTIS 25MG; 5MG
25MG; 10MGN018872 001 Jul 22, 1987
N018872 002 Jul 22, 1987HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

INDERIDE LA 120/50
WYETH AYERST 50MG; 120MG

N019059 002 Jul 03, 1985

INDERIDE LA 160/50
WYETH AYERST 50MG; 160MG

N019059 003 Jul 03, 1985

INDERIDE LA 80/50
WYETH AYERST 50MG; 80MG

N019059 001 Jul 03, 1985

TABLET; ORAL

INDERIDE-40/25
+ WYETH PHARMS INC 25MG; 40MG **

N018031 001

INDERIDE-80/25
+ WYETH PHARMS INC 25MG; 80MG **

N018031 002

PROPRANOLOL HYDROCHLORIDE & HYDROCHLOROTHIAZIDE
DURAMED PHARMS BARR 25MG; 40MG
25MG; 80MGA071126 001 Mar 02, 1987
A071127 001 Mar 02, 1987

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ACTAVIS ELIZABETH 25MG; 40MG
25MG; 80MGA070851 001 May 15, 1986
A070852 001 May 15, 1986ANI PHARMS 25MG; 40MG
25MG; 40MGA070705 002 Oct 01, 1986
A072043 002 Mar 14, 198825MG; 80MG
25MG; 80MGA070705 001 Oct 01, 1986
A072043 001 Mar 14, 1988CHARTWELL RX 25MG; 40MG
25MG; 80MGA071060 001 Aug 26, 1987
A071061 001 Aug 26, 1987IVAX SUB TEVA PHARMS 25MG; 40MG
25MG; 80MGA071552 001 Dec 01, 1988
A071553 001 Dec 01, 1988RISING 25MG; 40MG
25MG; 80MGA070947 002 Mar 04, 1987
A070947 001 Apr 01, 1987WARNER CHILCOTT 25MG; 40MG
25MG; 80MGA071771 001 Jan 26, 1988
A071772 001 Jan 26, 1988WATSON LABS 25MG; 40MG
25MG; 40MGA070301 001 Apr 18, 1986
A071498 001 Dec 18, 199125MG; 80MG
25MG; 80MGA070305 001 Apr 18, 1986
A071501 001 Dec 18, 1991HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCURETIC

+ PFIZER PHARMS 12.5MG; EQ 10MG BASE
+ 12.5MG; EQ 20MG BASE
+ 25MG; EQ 20MG BASEN020125 001 Dec 28, 1999
N020125 002 Dec 28, 1999
N020125 003 Dec 28, 1999

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX 12.5MG; EQ 10MG BASE
12.5MG; EQ 20MG BASEA076374 001 Mar 31, 2004
A076374 002 Mar 31, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

	25MG;EQ 20MG BASE	A076374	003	Mar 31, 2004
MYLAN	12.5MG;EQ 10MG BASE	A077093	001	Mar 28, 2005
	12.5MG;EQ 20MG BASE	A077093	002	Mar 28, 2005
	25MG;EQ 20MG BASE	A077093	003	Mar 28, 2005
SUN PHARM INDS LTD	12.5MG;EQ 10MG BASE	A078211	001	Mar 04, 2009
	12.5MG;EQ 20MG BASE	A078211	002	Mar 04, 2009
	25MG;EQ 20MG BASE	A078211	003	Mar 04, 2009

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

H.R.-50

WHITEWORTH TOWN PLSN	50MG;0.125MG	A085338	001	
HYDRO-RESERP				
ABC HOLDING	50MG;0.125MG	A084714	002	Jun 29, 1982
HYDRO-SERP "25"				
SANDOZ	25MG;0.125MG	A084827	001	
HYDRO-SERP "50"				
SANDOZ	50MG;0.125MG	A085213	001	
HYDROCHLOROTHIAZIDE W/ RESERPINE				
IVAX SUB TEVA PHARMS	25MG;0.1MG	A083572	001	
	25MG;0.125MG	A083571	001	
	50MG;0.1MG	A083568	001	
	50MG;0.125MG	A083573	001	
PHARMERAL	25MG;0.125MG	A085421	001	
	50MG;0.125MG	A085420	001	
ROXANE	50MG;0.125MG	A084603	001	
WATSON LABS	25MG;0.125MG	A084466	001	
	25MG;0.125MG	A085317	001	
	25MG;0.125MG	A086330	002	
	50MG;0.125MG	A083666	001	
	50MG;0.125MG	A084467	001	
	50MG;0.125MG	A086331	001	
HYDROPRES 25				
MERCK	25MG;0.125MG	N011958	002	
HYDROPRES 50				
MERCK	50MG;0.125MG	N011958	003	
RESERPINE AND HYDROCHLOROTHIAZIDE				
BARR	25MG;0.125MG	A084580	001	
	50MG;0.125MG	A084579	001	
SANDOZ	50MG;0.125MG	A088200	001	Jan 31, 1984
RESERPINE AND HYDROCHLOROTHIAZIDE-50				
WEST WARD	50MG;0.125MG	A088189	001	May 10, 1984
SERPASIL-ESIDRIX #1				
NOVARTIS	25MG;0.1MG	N011878	003	
SERPASIL-ESIDRIX #2				
NOVARTIS	50MG;0.1MG	N011878	005	

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

ALDACTAZIDE

+ PFIZER	50MG;50MG	N012616	005	Dec 30, 1982
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE				
ASCOT	25MG;25MG	A088025	001	Nov 23, 1984
CHARTWELL RX	25MG;25MG	A086881	001	
MUTUAL PHARM	25MG;25MG	A087267	001	
PUREPAC PHARM	25MG;25MG	A087999	001	Nov 06, 1985
SUPERPHARM	25MG;25MG	A089137	001	Aug 26, 1985
WATSON LABS	25MG;25MG	A087398	001	
SPIRONOLACTONE W/ HYDROCHLOROTHIAZIDE				
IVAX PHARMS	25MG;25MG	A087004	002	May 24, 1982
LEDERLE	25MG;25MG	A087511	001	
PARKE DAVIS	25MG;25MG	A087948	001	Feb 22, 1983
PUREPAC PHARM	25MG;25MG	A088054	001	Aug 18, 1983
UPSHER SMITH	25MG;25MG	A087553	001	
USL PHARMA	25MG;25MG	A087651	001	
VANGARD	25MG;25MG	A087655	001	
WATSON LABS	25MG;25MG	A085974	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE W/ HYDROCHLOROTHIAZIDE
25MG; 25MG

A086026 001

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND HYDROCHLOROTHIAZIDE

MACLEODS PHARMS LTD	12.5MG; 40MG
	12.5MG; 80MG
	25MG; 80MG
MYLAN	12.5MG; 40MG
	12.5MG; 80MG
	25MG; 80MG
TORRENT	12.5MG; 40MG
	12.5MG; 80MG
	25MG; 80MG

A204169	001	Nov 02, 2015
A204169	002	Nov 02, 2015
A204169	003	Nov 02, 2015
A091648	001	Feb 25, 2014
A091648	002	Feb 25, 2014
A091648	003	Feb 25, 2014
A201192	001	Feb 25, 2014
A201192	002	Feb 25, 2014
A201192	003	Feb 25, 2014

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

TABLET; ORAL

TIMOLIDE 10-25

MERCK 25MG; 10MG

N018061 001

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

+ GLAXOSMITHKLINE LLC 25MG; 37.5MG **
25MG; 50MG **N016042 003 Mar 03, 1994
N016042 002

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

ANI PHARMS	25MG; 37.5MG
	25MG; 50MG
CHARTWELL RX	25MG; 50MG
DURAMED PHARMS BARR	25MG; 37.5MG
MYLAN	25MG; 37.5MG
NOVARTIS	25MG; 37.5MG
VITARINE	25MG; 50MG

A074970	001	Jan 06, 1998
A074259	001	Mar 30, 1995
A073191	001	Jul 31, 1991
A075052	001	Jun 18, 1999
A074701	001	Jun 07, 1996
A074857	001	Sep 09, 1997
A071737	001	Feb 12, 1988

TABLET; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AM THERAP	50MG; 75MG
ANI PHARMS	50MG; 75MG
PLIVA	25MG; 37.5MG
QUANTUM PHARMICS	50MG; 75MG
WATSON LABS	50MG; 75MG

A072022	001	Apr 17, 1988
A073467	001	Jan 31, 1996
A074026	001	Apr 26, 1996
A071980	001	Apr 17, 1988
A071969	001	Apr 17, 1988

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

APOTEX INC	12.5MG; 80MG
	12.5MG; 160MG
	12.5MG; 320MG
	25MG; 160MG
	25MG; 320MG
CADILA	12.5MG; 80MG
	12.5MG; 160MG
	12.5MG; 320MG
	25MG; 160MG
	25MG; 320MG
WATSON LABS TEVA	12.5MG; 80MG
	12.5MG; 160MG
	12.5MG; 320MG
	25MG; 160MG
	25MG; 320MG

A203026	001	Mar 21, 2013
A203026	002	Mar 21, 2013
A203026	003	Mar 21, 2013
A203026	004	Mar 21, 2013
A203026	005	Mar 21, 2013
A203000	001	Mar 15, 2019
A203000	002	Mar 15, 2019
A203000	003	Mar 15, 2019
A203000	004	Mar 15, 2019
A203000	005	Mar 15, 2019
A091519	001	Mar 21, 2013
A091519	002	Mar 21, 2013
A091519	003	Mar 21, 2013
A091519	004	Mar 21, 2013
A091519	005	Mar 21, 2013

HYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

ZOHYDRO ER

+ RECRO GAINESVILLE	10MG
+	15MG
+	20MG
+	30MG
+	40MG
+	50MG

N202880	001	Oct 25, 2013
N202880	002	Oct 25, 2013
N202880	003	Oct 25, 2013
N202880	004	Oct 25, 2013
N202880	005	Oct 25, 2013
N202880	006	Oct 25, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCODONE BITARTRATE

TABLET, EXTENDED RELEASE;ORAL

VANTRELA ER

+	TEVA BRANDED PHARM	15MG	N207975	001	Jan 17, 2017
+		30MG	N207975	002	Jan 17, 2017
+		45MG	N207975	003	Jan 17, 2017
+		60MG	N207975	004	Jan 17, 2017
+		90MG	N207975	005	Jan 17, 2017

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET;ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

	ANI PHARMS	5MG;200MG	A077454	001	Jun 23, 2010
	SUN PHARM INDS INC	2.5MG;200MG	A091633	001	May 28, 2013
		5MG;200MG	A091633	002	May 28, 2013
		7.5MG;200MG	A091633	003	May 28, 2013
		10MG;200MG	A091633	004	May 28, 2013
	TEVA	7.5MG;200MG	A076023	001	Apr 11, 2003
	REPREXAIN				
	AMNEAL PHARMS NY	2.5MG;200MG	A076642	003	Oct 19, 2007
		10MG;200MG	A076642	004	Oct 19, 2007
	VICOPROFEN				
+	ABBVIE	7.5MG;200MG **	N020716	001	Sep 23, 1997

HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP;ORAL

CODAMINE

	ALPHARMA US PHARMS	5MG/5ML;25MG/5ML	A075103	001	Sep 29, 2000
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HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

	APOZEAL PHARMS	5MG/5ML;60MG/5ML	A206661	001	Jan 23, 2019
	MAYNE PHARMA INC	5MG/5ML;60MG/5ML	A205658	001	Nov 17, 2015
	PADAGIS US	5MG/5ML;60MG/5ML	A204658	001	Apr 29, 2014
	TRIS PHARMA INC	5MG/5ML;60MG/5ML	A203839	001	Oct 28, 2014
	REZIRA				
+	PERSION	5MG/5ML;60MG/5ML **	N022442	001	Jun 08, 2011

HYDROCORTAMATE HYDROCHLORIDE

OINTMENT;TOPICAL

MAGNACORT

	PFIZER	0.5%	N010554	001	
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HYDROCORTISONE

AEROSOL;TOPICAL

AEROSEB-HC

	ALLERGAN HERBERT	0.5%	A085805	001	
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CREAM;TOPICAL

CORT-DOME

	BAYER PHARMS	0.5%	N009585	003	
		1%	N009585	001	

DERMACORT

	MONARCH PHARMS	1%	A083011	002	
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ELDECORT

	VALEANT PHARM INTL	1%	A080459	001	
		2.5%	A084055	001	

FLEXICORT

	WESTWOOD SQUIBB	0.5%	A087136	003	Apr 08, 1982
		1%	A087136	002	Apr 08, 1982
		2.5%	A087136	001	Apr 08, 1982

H-CORT

	PHARM ASSOC	0.5%	A086823	001	
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HC #1

	BAYER PHARMS	0.5%	A080438	001	
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HC #4

	BAYER PHARMS	1%	A080438	002	
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HC (HYDROCORTISONE)

	C AND M PHARMA	0.5%	A080482	003	
		1%	A080482	004	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

CREAM;TOPICAL

HI-COR

C AND M PHARMA 2.5% A080483 001

HYDROCORTISONE

ALPHARMA US PHARMS 2.5% A089754 001 Feb 01, 1989

ALTANA 0.5% A080848 002

1% A080848 003

AMBIX 1% A086080 001

2.5% A086271 001

CHARTWELL MOLECULAR 2.5% A040503 001 Mar 12, 2004

ENCUBE 2.5% A203810 001 Jul 23, 2018

EVERYLIFE 0.5% A080452 001

1% A080452 002

G AND W LABS 1% A084059 001

INGRAM PHARM 0.5% A080456 002

1% A080456 003

IVAX PHARMS 1% A085733 001

NASKA 1% A089706 001 Mar 10, 1988

PERRIGO NEW YORK 0.5% A084970 002

1% A085026 001

PHARMADERM 1% A088845 001 Feb 27, 1986

2.5% A089413 001 Dec 16, 1986

PHARMAFAIR 1% A087838 001 Jul 28, 1982

STIEFEL 1% A086170 001

SYOSSET 0.5% A085527 001

TARO 0.5% A086154 001

1% A086155 001

TEVA 0.5% A080400 002

1% A080400 003

1% A085191 001

2.5% A080400 004

TOPIDERM 1% A089273 001 Feb 17, 1989

USL PHARMA 1% A088027 001 Sep 27, 1983

2.5% A088029 001 Sep 27, 1983

WHITEWORTH TOWN PLSN 1% A080496 002

HYTONE

+ VALEANT INTL 1% ** A080472 003

+ 2.5% ** A080472 004

NOGENIC HC

IVAX PHARMS 1% A087427 001 Apr 04, 1988

NUTRACORT

BAUSCH 0.5% A080442 002

1% A080442 003

PENECORT

ALLERGAN HERBERT 1% A088216 001 Jun 06, 1984

PROTOCORT

MONARCH PHARMS 1% A083011 001

SYNACORT

BAUSCH 0.5% A087459 001

+ 1% A087458 001

+ 2.5% A087457 001

ENEMA;RECTAL

HYDROCORTISONE

TEVA PHARMS 100MG/60ML A074171 001 May 27, 1994

GEL;TOPICAL

NUTRACORT

HEALTHPOINT 1% A084698 001

PENECORT

ALLERGAN HERBERT 1% A088215 001 Jun 06, 1984

INJECTABLE;INJECTION

CORTEF

PHARMACIA AND UPJOHN 50MG/ML N009864 001

LOTION;TOPICAL

ACTICORT

BAKER NORTON 1% A086535 001

ALA-CORT

CROWN LABS 1% A083201 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

LOTION; TOPICAL			
BALNEOL-HC			
SOLVAY	1%	A088041 001	Dec 03, 1982
BETA-HC			
BETA DERMAC	1%	A089495 001	Jan 25, 1988
CETACORT			
BAUSCH	0.5%	A080426 002	
	1%	A080426 001	
CORT-DOME			
BAYER PHARMS	0.5%	N009895 003	
	1%	N009895 001	
DERMACORT			
SOLVAY	0.5%	A084573 002	
	1%	A086462 001	
EPICORT			
BLULINE	0.5%	A083219 002	
GLYCORT			
HERAN	1%	A087489 001	Oct 03, 1983
H-CORT			
PHARM ASSOC	0.5%	A086824 001	
HYDROCORTISONE			
ALPHARMA US PHARMS	0.5%	A087317 001	Jun 07, 1982
	1%	A087315 001	Jun 07, 1982
CHARTWELL MOLECULAR	2.5%	A040417 001	Jul 30, 2003
FOUGERA PHARMS	2.5%	A040351 001	Jul 25, 2000
MERICON	0.5%	A085282 001	
	1%	A085282 002	Feb 26, 1987
NASKA	1%	A089705 001	Apr 25, 1988
PERRIGO NEW YORK	0.5%	A085662 001	
	1%	A085663 001	
TARO	1%	A089024 001	Feb 12, 1986
VERTICE	2.5%	A203804 001	Jul 27, 2018
HYTONE			
+ VALEANT INTL	1% **	A080473 003	
+	2.5% **	A080473 004	Nov 30, 1982
NUTRACORT			
DOW PHARM	0.5%	A080443 002	
	1%	A080443 003	
	2.5%	A087644 001	Aug 24, 1982
STIE-CORT			
PADAGIS US	1%	A089066 001	Nov 25, 1985
OINTMENT; TOPICAL			
CORTRIL			
PFIZER GLOBAL	1%	N009176 001	
	2.5%	N009176 002	
HC (HYDROCORTISONE)			
C AND M PHARMA	0.5%	A080481 001	
	1%	A080481 002	
HYDROCORTISONE			
ACTAVIS MID ATLANTIC	1%	A087796 001	Oct 13, 1982
ALTANA	0.5%	A080489 002	
	1%	A080489 003	
AMBIX	1%	A086079 001	
	2.5%	A086272 001	
NASKA	1%	A089704 001	Mar 10, 1988
PERRIGO NEW YORK	0.5%	A084969 003	
	1%	A085028 001	
PHARMADERM	1%	A088842 001	Feb 09, 1987
TARO	0.5%	A086256 001	
	2.5%	A040310 001	Dec 29, 2000
USL PHARMA	1%	A088061 001	Sep 27, 1983
	2.5%	A088039 001	Sep 27, 1983
HYTONE			
+ DERMIK LABS	1% **	A080474 003	
+	2.5% **	A080474 004	
PENECORT			
ALLERGAN HERBERT	2.5%	A088217 001	Jun 06, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

POWDER;FOR RX COMPOUNDING

H-CORT

TORCH 100% A087834 001 Mar 29, 1982

HYDRO-RX

X GEN PHARMS 100% A085982 001

HYDROCORTISONE

PADDOCK LLC 100% A088082 001 Apr 08, 1983

SOLUTION;TOPICAL

PENECORT

+ ALLERGAN HERBERT 1% A088214 001 Jun 06, 1984

TEXACORT

MISSION PHARMA 1% A080425 001

TABLET;ORAL

CORTRIL

PFIZER 10MG N009127 005

20MG N009127 003

HYDROCORTISONE

BARR 20MG A083999 001

CHARTWELL MOLECULAR 20MG A085070 001

ELKINS SINN 20MG A080624 001

FERRANTE 10MG A080568 001

20MG A080568 002

HIKMA INTL PHARMS 5MG A083365 002 Feb 23, 2015

10MG A083365 003 Feb 23, 2015

20MG A083365 001

IMPAX LABS 20MG A080781 001

INWOOD LABS 20MG A080732 001

NEXGEN PHARMA INC 20MG A083140 001

PANRAY 10MG N009659 001

20MG N009659 002

PARKE DAVIS 20MG A084243 001

PUREPAC PHARM 10MG A084247 003 Aug 31, 1982

20MG A080395 001

20MG A084247 002

ROXANE 10MG A088539 001 Mar 21, 1984

SANDOZ 20MG A080642 002

STRIDES PHARMA 5MG A040761 001 Jul 16, 2007

10MG A040761 002 Jul 16, 2007

20MG A040761 003 Jul 16, 2007

WATSON LABS 20MG A080355 001

WHITEWORTH TOWN PLSN 10MG A080344 001

20MG A080344 002

HYDROCORTONE

MERCK 10MG N008506 007

20MG N008506 011

TABLET;VAGINAL

CORTRIL

PFIPHARMECS 10MG N009796 001

HYDROCORTISONE ACETATE

CREAM;TOPICAL

CARMOL HC

+ FOUGERA PHARMS 1% A080505 001

HEMSOL-HC

ABLE 1% A081274 001 Jun 19, 1992

HYDROCORTISONE ACETATE

CENCI 1% A080419 001 Jan 25, 1982

IMPERIUM 2.5% A040259 001 Jul 29, 1999

PARKE DAVIS 1% A089914 001 Jan 03, 1989

PUREPAC PHARM 0.5% A086050 001

+ 1% A086052 001

MICORT-HC

SEBELA IRELAND LTD 2% A040398 001 Mar 29, 2002

U-CORT

TARO 1% A089472 001 Jun 13, 1988

INJECTABLE;INJECTION

CORTEF ACETATE

PHARMACIA AND UPJOHN 50MG/ML N009378 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE ACETATE

INJECTABLE; INJECTION

CORTRIL

PFIZER 25MG/ML N009164 001

HYDROCORTISONE ACETATE

BEL MAR 25MG/ML A083739 001

50MG/ML A083739 002

EPIC PHARMA LLC 25MG/ML N009637 001

50MG/ML N009637 002

WATSON LABS 25MG/ML A083128 001

25MG/ML A083759 001

50MG/ML A083759 002

50MG/ML A085214 001

HYDROCORTONE

MERCK 25MG/ML N008228 001

50MG/ML N008228 004

LOTION; TOPICAL

DRICORT

INGRAM PHARM 0.5% A086207 001

OINTMENT; OPHTHALMIC

HYDROCORTISONE ACETATE

FERA PHARMS 0.5% A080828 001

OINTMENT; OPHTHALMIC, OTIC

HYDROCORTONE

MERCK 1.5% N009018 003

OINTMENT; TOPICAL

CORTEF ACETATE

PHARMACIA AND UPJOHN 1% N008917 002

+ 2.5% ** N008917 001

PASTE; TOPICAL

ORABASE HCA

COLGATE 0.5% A083205 001

POWDER; FOR RX COMPOUNDING

HYDROCORTISONE ACETATE

X GEN PHARMS 100% A085981 001

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-CORTEF

PHARMACIA AND UPJOHN 1%;EQ 3.5MG BASE/GM A061049 001

2.5%;EQ 3.5MG BASE/GM A061049 002

OINTMENT; OPHTHALMIC

NEO-CORTEF

PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/GM A060610 001

1.5%;EQ 3.5MG BASE/GM A060610 002

OINTMENT; TOPICAL

NEO-CORTEF

PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/GM A060751 001

1%;EQ 3.5MG BASE/GM A060751 002

2.5%;EQ 3.5MG BASE/GM A060751 003

SUSPENSION/DROPS; OPHTHALMIC

COR-OTICIN

EPIC PHARMA LLC 1.5%;EQ 3.5MG BASE/ML A060188 001

NEO-CORTEF

PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/ML A060612 002

1.5%;EQ 3.5MG BASE/ML A060612 001

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL

CORTISPORIN

+ MONARCH PHARMS 0.5%;EQ 3.5MG BASE/GM;10,000 UNITS/GM N050218 001 Aug 09, 1985

HYDROCORTISONE ACETATE; OXYTETRACYCLINE HYDROCHLORIDE

SUSPENSION; OPHTHALMIC

TERRA-CORTRIL

PFIZER 1.5%;EQ 5MG BASE/ML A061016 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED; TOPICAL

HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%

GENUS 1%;1%

A089440 001 May 17, 1988

LOTION; TOPICAL

PRAMOSONE

FERNDALE LABS 0.5%;1%

A083213 002

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

LOCOID

YAMANOUCHI 0.1%

N018795 001 Jan 07, 1983

OINTMENT; TOPICAL

LOCOID

YAMANOUCHI 0.1%

N019106 001 Jul 03, 1984

SOLUTION; TOPICAL

LOCOID

YAMANOUCHI 0.1%

N019819 001 Sep 15, 1988

HYDROCORTISONE CYPIONATE

SUSPENSION; ORAL

CORTEF

PHARMACIA AND UPJOHN EQ 10MG BASE/5ML

N009900 001

HYDROCORTISONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HYDROCORTONE

+ MERCK EQ 50MG BASE/ML

N012052 001

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-HYDROCORT

ABBOTT EQ 100MG BASE/VIAL

A085928 001

EQ 100MG BASE/VIAL

A089577 001 Apr 11, 1989

EQ 250MG BASE/VIAL

A089578 001 Apr 11, 1989

EQ 500MG BASE/VIAL

A089579 001 Apr 11, 1989

EQ 1GM BASE/VIAL

A089580 001 Apr 11, 1989

HOSPIRA EQ 100MG BASE/VIAL

A040666 001 Apr 06, 2006

EQ 100MG BASE/VIAL

A085929 001

EQ 250MG BASE/VIAL

A085930 001

EQ 500MG BASE/VIAL

A085931 001

EQ 1GM BASE/VIAL

A085932 001

HYDROCORTISONE SODIUM SUCCINATE

ABRAXIS PHARM EQ 100MG BASE/VIAL

A088667 001 Jun 08, 1984

EQ 100MG BASE/VIAL

A088712 001 Jun 08, 1984

EQ 250MG BASE/VIAL

A088668 001 Jun 08, 1984

EQ 500MG BASE/VIAL

A088669 001 Jun 08, 1984

EQ 1GM BASE/VIAL

A088670 001 Jun 08, 1984

BAXTER HLTHCARE EQ 100MG BASE/VIAL

A086619 001

EQ 250MG BASE/VIAL

A087567 001

EQ 500MG BASE/VIAL

A087568 001

EQ 1GM BASE/VIAL

A087569 001

INTL MEDICATION EQ 100MG BASE/VIAL

A087532 001 Mar 19, 1982

WATSON LABS EQ 100MG BASE/VIAL

A084737 002

EQ 100MG BASE/VIAL

A084738 001

EQ 250MG BASE/VIAL

A084737 001

EQ 500MG BASE/VIAL

A084747 001

EQ 1GM BASE/VIAL

A084748 001

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE

COSETTE 0.2%

A074489 001 Aug 12, 1998

WESTCORT

+ SUN PHARM INDS INC 0.2% **

N017950 001

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE

FOUGERA PHARMS 0.2%

A075085 001 Jul 31, 2001

WESTCORT

+ SUN PHARM INDS INC 0.2% **

N018726 001 Aug 08, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-CORT-DOME

BAYER PHARMS	0.5%;EQ 3.5MG BASE/GM	N050237	006	Jun 05, 1984
	1%;EQ 3.5MG BASE/GM	N050237	005	Jun 05, 1984

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

CORTISPORIN

+ MONARCH PHARMS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **	N050479	001	
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE				
AMRING PHARMS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A065216	001	Oct 31, 2005
NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE				
PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062394	001	Sep 29, 1982

OTOCORT

WATSON LABS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A060730	002	
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SUSPENSION/DROPS; OPHTHALMIC

CORTISPORIN

MONARCH PHARMS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N050169	001	
NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE				
PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062623	001	Sep 24, 1985

SUSPENSION/DROPS; OTIC

NEOMYCIN SULFATE, POLYMYXIN B SULFATE & HYDROCORTISONE				
PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062617	001	Sep 18, 1985

OTICAIR

PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062399	001	Nov 18, 1982
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OTOBIONE

SCHERING	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A061816	001	
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OTOCORT

ACTAVIS LABS FL INC	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062521	001	Jul 11, 1985
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PEDIOTIC

MONARCH PHARMS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062822	001	Sep 29, 1987
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HYDROCORTISONE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

OTOBiotic

SCHERING	5MG/ML;EQ 10,000 UNITS BASE/ML	A062302	001	
PYOCIDIN				
FOREST LABS	5MG/ML;EQ 10,000 UNITS BASE/ML	A061606	001	

HYDROCORTISONE; TETRACYCLINE HYDROCHLORIDE

OINTMENT; OPHTHALMIC

ACHROMYCIN

LEDERLE	1.5%;1%	N050272	001	
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HYDROCORTISONE; UREA

CREAM; TOPICAL

ALPHADERM

BIOGLAN	1%;10%	A086008	001	
CALMURID HC				
PHARMACIA AND UPJOHN	1%;10%	A083947	001	

HYDROFLUMETHIAZIDE

TABLET; ORAL

DIUCARDIN

WYETH AYERST	50MG	A083383	001	
HYDROFLUMETHIAZIDE				
PAR PHARM	50MG	A088850	001	May 31, 1985
WATSON LABS	50MG	A088031	001	Apr 06, 1983
	50MG	A088528	001	Aug 15, 1984

SALURON

+ SHIRE LLC	50MG	N011949	001	
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HYDROFLUMETHIAZIDE; RESERPINE

TABLET; ORAL

HYDROFLUMETHIAZIDE AND RESERPINE

USL PHARMA	50MG;0.125MG	A088195	001	Oct 26, 1983
WATSON LABS	25MG;0.125MG	A088127	001	Mar 22, 1983
	50MG;0.125MG	A088110	001	Mar 22, 1983

RESERPINE AND HYDROFLUMETHIAZIDE

IVAX PHARMS	50MG;0.125MG	A088932	001	Jan 11, 1985
PAR PHARM	50MG;0.125MG	A088907	001	Sep 20, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROFLUMETHIAZIDE; RESERPINE

TABLET; ORAL

SALUTENSIN

SHIRE

50MG; 0.125MG

N012359 003

SALUTENSIN-DEMI

SHIRE

25MG; 0.125MG

N012359 004

HYDROGEN PEROXIDE

SOLUTION; TOPICAL

ESKATA

+ ACLARIS

40%

N209305 001 Dec 14, 2017

HYDROMORPHONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PALLADONE

PURDUE PHARMA LP

12MG

N021044 001 Sep 24, 2004

16MG

N021044 002 Sep 24, 2004

24MG

N021044 003 Sep 24, 2004

32MG

N021044 004 Sep 24, 2004

INJECTABLE; INJECTION

DILAUDID

+ FRESENIUS KABI USA

4MG/ML

N019034 005 Apr 30, 2009

DILAUDID-HP

+ FRESENIUS KABI USA

10MG/ML

N019034 001 Jan 11, 1984

250MG/VIAL

N019034 002 Aug 04, 1994

HYDROMORPHONE HYDROCHLORIDE

BARR

10MG/ML

A076444 001 Apr 25, 2003

HOSPIRA

10MG/ML

A074598 001 Jun 19, 1997

WATSON LABS

10MG/ML

A074317 001 Aug 23, 1995

SOLUTION; ORAL

HYDROMORPHONE HYDROCHLORIDE

ANDA REPOSITORY

5MG/5ML

A207108 001 Apr 22, 2020

TABLET; ORAL

HYDROMORPHONE HYDROCHLORIDE

ANDA REPOSITORY

2MG

A077471 002 Dec 09, 2009

4MG

A077471 003 Dec 09, 2009

8MG

A077471 001 Dec 09, 2009

HIKMA

4MG

A074597 003 May 29, 2009

8MG

A074597 001 Jul 29, 1998

NESHER PHARMS

2MG

A077311 001 Nov 09, 2005

4MG

A077311 002 Nov 09, 2005

8MG

A077311 003 Nov 09, 2005

NOSTRUM LABS INC

8MG

A076723 001 Oct 18, 2005

TABLET, EXTENDED RELEASE; ORAL

EXALGO

+ SPECGX LLC

8MG **

N021217 001 Mar 01, 2010

+

12MG **

N021217 002 Mar 01, 2010

+

16MG **

N021217 003 Mar 01, 2010

+

32MG **

N021217 004 Aug 24, 2012

HYDROMORPHONE HYDROCHLORIDE

ACTAVIS LABS FL INC

8MG

A202144 001 May 12, 2014

12MG

A202144 002 May 12, 2014

16MG

A202144 003 May 12, 2014

32MG

A202144 004 Jun 30, 2016

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

ALPHAREDISOL

MERCK

1MG/ML

A080778 001

CYANOKIT

BTG INTL

2.5GM/VIAL (5GM/KIT)

N022041 002 Dec 15, 2006

HYDROXOCOBALAMIN

ABRAXIS PHARM

1MG/ML

A084921 001

WATSON LABS

1MG/ML

A085528 001

HYDROXOMIN

BEL MAR

1MG/ML

A084629 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYAMPHETAMINE HYDROBROMIDESOLUTION/DROPS;OPHTHALMIC
PAREDRINE

PHARMICS 1% N000004 004

HYDROXYAMPHETAMINE HYDROBROMIDE; TROPICAMIDESOLUTION/DROPS;OPHTHALMIC
PAREMYD

+ EPIC PHARMA LLC 1%;0.25% N019261 001 Jan 30, 1992

HYDROXYCHLOROQUINE SULFATE

TABLET;ORAL

HYDROXYCHLOROQUINE SULFATE

HIKMA PHARMS	200MG	A040760 001	Aug 15, 2007
INVATECH	200MG	A040150 001	Jan 27, 1996
TEVA PHARMS	200MG	A040081 001	Sep 30, 1994
WATSON LABS	200MG	A040133 001	Nov 30, 1995

HYDROXYPROGESTERONE CAPROATE

INJECTABLE;INJECTION

HYDROXYPROGESTERONE CAPROATE

ALLERGAN	125MG/ML	N017439 001	
	250MG/ML	N017439 002	
EPIC PHARMA LLC	125MG/ML	N018004 001	

SOLUTION;INTRAMUSCULAR

DELALUTIN

+ BRISTOL MYERS SQUIBB	125MG/ML (125MG/ML) **	N010347 004	
+	125MG/ML (125MG/ML) **	N016911 001	
+	250MG/ML (250MG/ML) **	N010347 002	
+	250MG/ML (250MG/ML) **	N016911 002	

HYDROXYPROGESTERONE CAPROATE

ASPEN GLOBAL INC	1250MG/5ML (250MG/ML)	A200271 001	Aug 24, 2015
EUGIA PHARMA	1250MG/5ML (250MG/ML)	A211142 001	May 09, 2019

HYDROXYSTILBAMIDINE ISETHIONATE

INJECTABLE;INJECTION

HYDROXYSTILBAMIDINE ISETHIONATE

SANOFI AVENTIS US 225MG/AMP N009166 001

HYDROXYUREA

CAPSULE;ORAL

HYDROXYUREA

BARR	250MG	A075143 002	Sep 21, 2000
BARR LABS INC	250MG	A075020 002	Jun 26, 2000
	500MG	A075020 001	Jul 30, 1998
ROXANE	500MG	A074476 001	Aug 18, 1995

TABLET;ORAL

HYDROXYUREA

BARR 1GM A075734 001 Aug 29, 2000

HYDROXYZINE HYDROCHLORIDE

INJECTABLE;INJECTION

HYDROXYZINE

BAXTER HLTHCARE 50MG/ML A085551 002

HYDROXYZINE HYDROCHLORIDE

ALTANA	25MG/ML	A087273 001	Apr 20, 1982
	50MG/ML	A087273 002	Apr 20, 1982
BAXTER HLTHCARE	25MG/ML	A085551 001	
DR REDDYS	50MG/ML	A085779 001	
FRESENIUS KABI USA	25MG/ML	A087329 001	
	25MG/ML	A088184 001	Mar 31, 1983
	50MG/ML	A087329 002	
	50MG/ML	A088185 001	Mar 31, 1983
HOSPIRA	25MG/ML	A087416 001	
	50MG/ML	A086821 001	
	50MG/ML	A087546 001	
PHARMAFAIR	25MG/ML	A088862 001	Feb 14, 1986
	25MG/ML	A089106 001	Feb 14, 1986
	50MG/ML	A088881 001	Feb 14, 1986
	50MG/ML	A089107 001	Feb 14, 1986
SMITH AND NEPHEW	25MG/ML	A087592 001	
SOLOPAK	25MG/ML	A086822 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

	25MG/ML	A087591	001	
	50MG/ML	A087310	001	
	50MG/ML	A087593	001	
	50MG/ML	A087595	001	
	50MG/ML	A087596	001	
WATSON LABS	25MG/ML	A085778	001	
	25MG/ML	A087274	001	
	50MG/ML	A087274	002	
WYETH AYERST	25MG/ML	A086258	001	
	50MG/ML	A086258	002	
ORGATRAK				
ORGANON USA INC	25MG/ML	A087014	001	
	50MG/ML	A087014	002	
VISTARIL				
+ PFIZER	25MG/ML **	N011111	001	
+	50MG/ML **	N011111	002	
SYRUP; ORAL				
ATARAX				
ROERIG	10MG/5ML **	N010485	001	
HYDROXYZINE HYDROCHLORIDE				
ALPHARMA US PHARMS	10MG/5ML	A088785	001	Feb 03, 1988
ANIMA	10MG/5ML	A086880	001	
APOZEAL PHARMS	10MG/5ML	A210634	001	Feb 26, 2019
HIKMA	10MG/5ML	A040010	001	Oct 28, 1994
KV PHARM	10MG/5ML	A087730	001	Jul 01, 1982
PAI HOLDINGS PHARM	10MG/5ML	A040391	001	Apr 10, 2002
TABLET; ORAL				
ATARAX				
+ PFIZER	10MG **	N010392	001	
+	25MG **	N010392	004	
+	50MG **	N010392	006	
+	100MG **	N010392	005	
HYDROXYZINE HYDROCHLORIDE				
ABLE	10MG	A040559	001	Jul 22, 2004
	25MG	A040562	001	Jul 22, 2004
	50MG	A040563	001	Jul 22, 2004
ACTAVIS ELIZABETH	10MG	A089071	001	Jul 22, 1986
	25MG	A089072	001	Jul 22, 1986
	50MG	A089073	001	Jul 22, 1986
HALSEY	10MG	A089366	001	May 02, 1988
	25MG	A089117	001	May 02, 1988
	50MG	A089396	001	May 02, 1988
IVAX PHARMS	10MG	A087216	001	
	25MG	A087410	001	
	50MG	A087411	001	
KV PHARM	10MG	A087819	001	Jun 23, 1982
	25MG	A087820	001	Jun 23, 1982
	50MG	A087821	001	Jun 23, 1982
	100MG	A087822	001	Jun 23, 1982
MUTUAL PHARM	10MG	A088409	001	Nov 15, 1983
	25MG	A087857	001	Apr 18, 1983
	50MG	A087860	001	Apr 18, 1983
PLIVA	100MG	A081054	001	Sep 25, 1995
PUREPAC PHARM	10MG	A088120	001	Sep 25, 1984
	25MG	A088121	001	Sep 25, 1984
	50MG	A088122	001	Sep 25, 1984
QUANTUM PHARMICS	10MG	A088540	001	Oct 22, 1985
	25MG	A088551	001	Oct 22, 1985
	50MG	A088529	001	Oct 22, 1985
RISING	10MG	A091176	001	Jun 07, 2010
	25MG	A091176	002	Jun 07, 2010
	50MG	A091176	003	Jun 07, 2010
SANDOZ	10MG	A087246	002	
	25MG	A085247	001	
	50MG	A087245	001	
STRIDES PHARMA	10MG	A087602	001	Jan 22, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE HYDROCHLORIDE

TABLET;ORAL

HYDROXYZINE HYDROCHLORIDE

	25MG	A087603	001	Jan 22, 1982
	50MG	A087604	001	Jan 22, 1982
SUN PHARM INDS INC	10MG	A040899	001	Jun 10, 2008
	25MG	A040899	002	Jun 10, 2008
	50MG	A040899	003	Jun 10, 2008
SUN PHARM INDUSTRIES	10MG	A089381	001	May 19, 1986
	25MG	A089382	001	May 19, 1986
	50MG	A089383	001	May 19, 1986
	100MG	A087862	001	Apr 18, 1983
SUPERPHARM	10MG	A088794	001	Dec 05, 1984
	25MG	A088795	001	Dec 05, 1984
	50MG	A088796	001	Dec 05, 1984
USL PHARMA	10MG	A089121	001	Mar 20, 1986
	25MG	A089122	001	Mar 20, 1986
	50MG	A089123	001	Mar 20, 1986
WATSON LABS	10MG	A081149	001	Mar 18, 1994
	10MG	A086827	001	
	10MG	A088348	001	Sep 15, 1983
	25MG	A081150	001	Mar 18, 1994
	25MG	A086829	001	
	25MG	A088349	001	Sep 15, 1983
	50MG	A081151	001	Mar 18, 1994
	50MG	A086836	001	
	50MG	A088350	001	Sep 15, 1983

HYDROXYZINE PAMOATE

CAPSULE;ORAL

HY-PAM "25"

TEVA	EQ 25MG HYDROCHLORIDE	A088713	001	Mar 04, 1985
HYDROXYZINE PAMOATE				
BEXIMCO PHARMS USA	EQ 25MG HYDROCHLORIDE	A081127	001	Jun 28, 1991
DURAMED PHARMS BARR	EQ 25MG HYDROCHLORIDE	A088593	001	Feb 29, 1984
	EQ 50MG HYDROCHLORIDE	A088594	001	Feb 29, 1984
	EQ 100MG HYDROCHLORIDE	A088595	001	Feb 29, 1984
IVAX SUB TEVA PHARMS	EQ 25MG HYDROCHLORIDE	A087761	001	Mar 05, 1982
	EQ 50MG HYDROCHLORIDE	A087760	001	Mar 05, 1982
PAR PHARM	EQ 25MG HYDROCHLORIDE	A087656	001	Jun 11, 1982
	EQ 25MG HYDROCHLORIDE	A089145	001	Mar 17, 1986
	EQ 50MG HYDROCHLORIDE	A087657	001	Jun 11, 1982
	EQ 50MG HYDROCHLORIDE	A089146	001	Mar 17, 1986
	EQ 100MG HYDROCHLORIDE	A087658	001	Jun 11, 1982
SANDOZ	EQ 50MG HYDROCHLORIDE	A081128	001	Jun 28, 1991
	EQ 100MG HYDROCHLORIDE	A081129	001	Jun 28, 1991
SUPERPHARM	EQ 25MG HYDROCHLORIDE	A089031	001	Jan 02, 1987
	EQ 50MG HYDROCHLORIDE	A089032	001	Jan 02, 1987
	EQ 100MG HYDROCHLORIDE	A089033	001	Jan 02, 1987
VANGARD	EQ 25MG HYDROCHLORIDE	A088392	001	Sep 19, 1983
	EQ 50MG HYDROCHLORIDE	A088393	001	Sep 19, 1983
WATSON LABS	EQ 25MG HYDROCHLORIDE	A081165	001	Jul 31, 1991
	EQ 25MG HYDROCHLORIDE	A086698	001	
	EQ 25MG HYDROCHLORIDE	A086840	001	Jul 01, 1982
	EQ 50MG HYDROCHLORIDE	A086695	001	
	EQ 50MG HYDROCHLORIDE	A086705	001	Jul 01, 1982
	EQ 50MG HYDROCHLORIDE	A087767	001	Aug 16, 1982
	EQ 100MG HYDROCHLORIDE	A086697	001	
	EQ 100MG HYDROCHLORIDE	A086728	001	Oct 05, 1982
	EQ 100MG HYDROCHLORIDE	A087790	001	Aug 16, 1982
VISTARIL				
PFIZER	EQ 100MG HYDROCHLORIDE **	N011459	006	
SUSPENSION;ORAL				
VISTARIL				
PFIZER	EQ 25MG HYDROCHLORIDE/5ML	N011795	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

BONIVA

+ ROCHE EQ 3MG BASE/3ML ** N021858 001 Jan 06, 2006

IBANDRONATE SODIUM

AVET LIFESCIENCES EQ 3MG BASE/3ML A203987 001 Sep 02, 2014

NANG KUANG PHARM CO EQ 3MG BASE/3ML A204329 001 Jun 16, 2021

TABLET; ORAL

BONIVA

+ HOFFMANN LA ROCHE EQ 2.5MG BASE ** N021455 001 May 16, 2003

+ EQ 150MG BASE ** N021455 002 Mar 24, 2005

IBANDRONATE SODIUM

MYLAN PHARMS INC EQ 150MG BASE A078995 001 Mar 19, 2012

SUN PHARM INDUSTRIES EQ 150MG BASE A078996 001 Aug 15, 2012

IBRUTINIB

CAPSULE; ORAL

IBRUTINIB

ZYDUS 70MG A211344 001 Mar 31, 2021

140MG A211344 002 Mar 31, 2021

TABLET; ORAL

IMBRUVICA

+ PHARMACYCLICS LLC 560MG N210563 004 Feb 16, 2018

IBUPROFEN

CAPSULE; ORAL

IBUPROFEN

CONTRACT PHARMACAL 200MG A074782 001 Jul 06, 1998

MIDOL

BAYER 200MG ** A070626 001 Sep 02, 1987

200MG ** A071002 001 Sep 02, 1987

SOLUTION; INTRAVENOUS

CALDOLOR

CUMBERLAND PHARMS 400MG/4ML (100MG/ML) N022348 001 Jun 11, 2009

SUSPENSION; ORAL

CHILDREN'S ADVIL

GLAXOSMITHKLINE 100MG/5ML N019833 002 Sep 19, 1989

CHILDREN'S ELIXSURE

MOBERG PHARMA NORTH 100MG/5ML N021604 001 Jan 07, 2004

IBU

ABBOTT 100MG/5ML N019784 001 Dec 18, 1989

IBUPROFEN

ARISE 100MG/5ML A200457 001 Aug 18, 2011

PAI HOLDINGS PHARM 100MG/5ML A205647 001 Nov 03, 2016

STRIDES PHARMA 100MG/5ML A211666 001 Feb 22, 2021

MOTRIN

+ MCNEIL CONSUMER 100MG/5ML ** N019842 001 Sep 19, 1989

SUSPENSION/DROPS; ORAL

IBUPROFEN

STRIDES PHARMA 50MG/1.25ML A214071 001 Jun 09, 2022

MOTRIN

MCNEIL 40MG/ML N020476 001 May 25, 1995

PEDIATRIC ADVIL

+ GLAXOSMITHKLINE 100MG/2.5ML N020812 001 Jan 30, 1998

TABLET; ORAL

ACHES-N-PAIN

LEDERLE 200MG A071065 001 May 28, 1987

CAP-PROFEN

PERRIGO 200MG A072097 001 Dec 08, 1987

IBU

BASF 400MG A070083 001 Feb 22, 1985

400MG N018197 001

600MG A070088 001 Feb 08, 1985

600MG A070099 001 Mar 29, 1985

800MG A070745 001 Jul 23, 1986

IBU-TAB

ALRA 400MG A071058 001 Aug 11, 1988

600MG A071059 001 Aug 11, 1988

800MG A071965 001 Aug 11, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET;ORAL

IBU-TAB 200

ALRA	200MG	A071057	001	Aug 11, 1988
IBUPRIN				
PLIVA	200MG	A071773	001	Jul 16, 1987
IBUPROFEN				
ABBOTT	600MG	A070556	001	Jun 14, 1985
	800MG	A071264	001	Jul 25, 1986
ANI PHARMS	200MG	A071144	001	Jan 20, 1987
	200MG	A072901	001	Dec 19, 1991
	200MG	A072903	001	Dec 19, 1991
CONTRACT PHARMACAL	200MG	A071265	001	Oct 15, 1986
	200MG	A071265	002	Sep 10, 1987
	200MG	A071735	001	Sep 10, 1987
	200MG	A073691	001	Feb 25, 1994
	200MG	A074931	001	Jul 20, 1998
HALSEY	200MG	A071027	001	Sep 29, 1987
	300MG	A071028	001	Mar 23, 1987
	400MG	A071029	001	Mar 23, 1987
	600MG	A071030	001	Mar 23, 1987
	800MG	A072137	001	Feb 05, 1988
IVAX SUB TEVA PHARMS	200MG	A071154	001	Oct 27, 1987
	200MG	A072040	001	Apr 29, 1988
	400MG	A071145	001	Sep 23, 1986
	600MG	A071146	001	Sep 23, 1986
	800MG	A071769	001	May 08, 1987
J AND J CONSUMER INC	400MG	A070081	001	Jun 16, 1986
L PERRIGO CO	400MG	A077114	001	Jul 18, 2005
	600MG	A077114	002	Jul 18, 2005
	800MG	A077114	003	Jul 18, 2005
LEDERLE	400MG	A070629	001	Sep 19, 1986
	600MG	A070630	001	Sep 19, 1986
LEINER	300MG	A071266	001	Oct 15, 1986
LNK	100MG	A076741	001	Jun 17, 2004
MCNEIL	600MG	A070476	001	Jun 16, 1986
MERRO PHARM	200MG	A070985	001	Oct 02, 1987
MYLAN	200MG	A071870	001	May 05, 1988
	400MG	A070045	001	Sep 24, 1985
	600MG	A070057	001	Sep 24, 1985
	800MG	A071999	001	Dec 03, 1987
NORTHSTAR HLTHCARE	400MG	A078132	001	Sep 10, 2007
	600MG	A078132	002	Sep 10, 2007
	800MG	A078132	003	Sep 10, 2007
OHM	200MG	A071163	001	Jul 15, 1986
OHM LABS	400MG	A070818	001	Dec 26, 1985
P AND L DEV LLC	200MG	A070733	001	Sep 19, 1986
PAR PHARM	300MG	A070328	001	Aug 06, 1985
PERRIGO	200MG	A072098	001	Dec 08, 1987
	200MG	A075995	001	Mar 14, 2002
PLIVA	400MG	A071666	001	Jun 18, 1987
	600MG	A071667	001	Jun 18, 1987
	800MG	A071668	001	Jun 18, 1987
PUREPAC PHARM	200MG	A071122	001	Oct 03, 1986
	200MG	A071664	001	Feb 03, 1987
	300MG	A071123	001	Sep 19, 1986
	400MG	A071124	001	Sep 19, 1986
	600MG	A071125	001	Sep 19, 1986
	800MG	A071964	001	Feb 01, 1988
RISING	300MG	A070736	002	Jun 12, 1986
	400MG	A070736	003	Jun 12, 1986
	600MG	A070736	001	Jun 12, 1986
	800MG	A071938	001	Jan 14, 1988
SANDOZ	200MG	A071807	001	Feb 25, 1988
	200MG	A074525	001	Dec 15, 1995
	200MG	A074533	001	Dec 15, 1995
	400MG	A072064	001	Jan 14, 1988
	600MG	A072065	001	Jan 14, 1988
	800MG	A072169	001	Dec 11, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET; ORAL

IBUPROFEN

STRIDES PHARMA	200MG	A071575 001	May 08, 1987
	400MG	A070329 001	Aug 06, 1985
	600MG	A070330 001	Aug 06, 1985
	800MG	A070986 001	Jul 25, 1986
SUN PHARM INDUSTRIES	200MG	A070493 001	Dec 24, 1985
	200MG	A070908 001	Sep 26, 1986
	200MG	A071462 001	Oct 02, 1986
	400MG	A070079 001	Jul 24, 1985
	600MG	A070080 001	Jul 24, 1985
	800MG	A071448 001	Feb 18, 1987
SUNSHINE	400MG	A204062 001	Sep 10, 2018
	600MG	A204062 002	Sep 10, 2018
	800MG	A204062 003	Sep 10, 2018
SUPERPHARM	600MG	A070709 001	Apr 25, 1986
TEVA	200MG	A073141 001	May 29, 1992
	400MG	A073343 001	Jun 30, 1992
	600MG	A073344 001	Jun 30, 1992
	800MG	A073345 001	Jun 30, 1992
ULTRATAB LABS INC	200MG	A209076 001	Jan 06, 2020
VINTAGE PHARMS	200MG	A071639 001	Feb 02, 1988
	200MG	A072249 001	Jan 10, 1989
	300MG	A071230 001	Oct 22, 1986
	400MG	A071231 001	Oct 22, 1986
	400MG	A071644 001	Feb 01, 1988
	600MG	A071232 001	Oct 22, 1986
	800MG	A072004 001	Nov 18, 1987
WATSON LABS	200MG	A070435 001	Mar 05, 1986
	200MG	A071765 001	Sep 04, 1987
	200MG	A071905 001	Mar 08, 1988
	300MG	A071338 001	Dec 01, 1986
	400MG	A070038 001	Sep 06, 1985
	400MG	A070436 001	Aug 21, 1985
	600MG	A070041 001	Sep 06, 1985
	600MG	A070437 001	Aug 21, 1985
	800MG	A071547 001	Jul 02, 1987
	800MG	A071911 001	Oct 13, 1987
YICHANG HUMANWELL	200MG	A214003 001	Oct 19, 2020
IBUPROHM			
OHM LABS	200MG	A071214 001	Dec 01, 1986
	400MG	A070469 001	Aug 29, 1985
JUNIOR STRENGTH IBUPROFEN			
L PERRIGO CO	100MG	A075367 001	Apr 22, 1999
MEDIPREN			
MCNEIL	200MG	A070475 001	Feb 06, 1986
	200MG	A071215 001	Jun 26, 1986
MIDOL			
BAYER	200MG	A070591 001	Sep 02, 1987
	200MG	A071001 001	Sep 02, 1987
MOTRIN			
+ MCNEIL CONSUMER	300MG **	N017463 003	
+	400MG **	N017463 002	
+	600MG **	N017463 004	
+	800MG **	N017463 005	May 22, 1985
MCNEIL PED	100MG	N020418 001	Nov 16, 1994
MOTRIN MIGRAINE PAIN			
J AND J CONSUMER INC	200MG	N019012 004	Feb 25, 2000
NUPRIN			
BRISTOL MYERS	200MG	A072035 001	Feb 16, 1988
	200MG	A072036 001	Feb 16, 1988
J AND J CONSUMER INC	200MG	N019012 001	May 18, 1984
	200MG	N019012 002	Jul 29, 1987
RUFEN			
BASF	600MG	N018197 002	Mar 05, 1984
TAB-PROFEN			
PERRIGO	200MG	A072095 001	Dec 08, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET, CHEWABLE;ORAL

CHILDREN'S MOTRIN

+ J AND J CONSUMER INC 50MG

N020601 001 Nov 15, 1996

IBUPROFEN

PERRIGO 50MG

A076359 001 Jan 16, 2004

JUNIOR STRENGTH MOTRIN

+ J AND J CONSUMER INC 100MG

N020601 003 Nov 15, 1996

MOTRIN

MCNEIL PED 50MG

N020135 001 Nov 16, 1994

100MG

N020135 002 Nov 16, 1994

IBUPROFEN SODIUM

TABLET;ORAL

IBUPROFEN SODIUM

PERRIGO R AND D EQ 200MG BASE

A206581 001 Aug 03, 2015

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

COMBUNOX

+ FOREST LABS 400MG;5MG **

N021378 001 Nov 26, 2004

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

ACTAVIS ELIZABETH 400MG;5MG

A078769 001 Jan 04, 2008

BARR LABS INC 400MG;5MG

A078316 001 Nov 29, 2007

WATSON LABS 400MG;5MG

A078394 001 Nov 26, 2007

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION;ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

PERRIGO 100MG/5ML;15MG/5ML

A076478 001 Nov 05, 2003

TABLET;ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA LTD 200MG;30MG

A213565 001 Mar 10, 2023

CONTRACT PHARMACAL 200MG;30MG

A075588 001 Apr 08, 2002

IBUTILIDE FUMARATE

INJECTABLE;INJECTION

IBUTILIDE FUMARATE

LUITPOLD 0.1MG/ML

A090240 001 Jan 11, 2010

MYLAN INSTITUTIONAL 0.1MG/ML

A090924 001 Jan 11, 2010

ICATIBANT ACETATE

INJECTABLE;SUBCUTANEOUS

ICATIBANT ACETATE

DR REDDYS EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A213054 001 Oct 05, 2020

GLENMARK PHARMS LTD EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

A213222 001 May 21, 2021

IDARUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

IDAMYCIN

PFIZER 5MG/VIAL

N050661 002 Sep 27, 1990

10MG/VIAL

N050661 001 Sep 27, 1990

20MG/VIAL

N050661 003 Apr 25, 1995

IDARUBICIN HYDROCHLORIDE

FRESENIUS KABI USA 1MG/ML

A065440 001 Aug 04, 2009

MYLAN LABS LTD 1MG/ML

A200144 001 Oct 11, 2012

SANDOZ 1MG/ML

A091293 001 Mar 29, 2011

TEVA PARENTERAL 5MG/VIAL

A065037 003 May 01, 2002

10MG/VIAL

A065037 002 May 01, 2002

20MG/VIAL

A065037 001 May 01, 2002

IDOXURIDINE

OINTMENT;OPHTHALMIC

STOXIL

GLAXOSMITHKLINE 0.5%

N015868 001

SOLUTION/DROPS;OPHTHALMIC

DENDRID

+ ALCON 0.1%

N014169 001

HERPLEX

ALLERGAN 0.1%

N013935 002

STOXIL

GLAXOSMITHKLINE 0.1%

N013934 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IFOSFAMIDE

INJECTABLE; INJECTION

IFOSFAMIDE

FRESENIUS KABI USA	1GM/20ML (50MG/ML)	A090181 001	Sep 22, 2009
	3GM/60ML (50MG/ML)	A090181 002	Sep 22, 2009
MYLAN LABS LTD	1GM/20ML (50MG/ML)	A201689 001	Nov 26, 2012
	3GM/60ML (50MG/ML)	A201689 002	Nov 26, 2012

IFOSFAMIDE; MESNA

INJECTABLE; INJECTION

IFEX/MESNEX KIT

BAXTER HLTHCARE	1GM/VIAL;100MG/ML	N019763 003	Oct 10, 1992
	3GM/VIAL;100MG/ML	N019763 004	Oct 10, 1992

INJECTABLE; INTRAVENOUS

IFOSFAMIDE/MESNA KIT

TEVA PHARMS USA	1GM/20ML;1GM/10ML (50MG/ML;100MG/ML)	A075874 001	Feb 26, 2002
	3GM/60ML;1GM/10ML (50MG/ML;100MG/ML)	A075874 002	Feb 26, 2002

ILOPERIDONE

TABLET; ORAL

ILOPERIDONE

INVENTIA	1MG	A207231 001	Nov 28, 2016
	2MG	A207231 002	Nov 28, 2016
	4MG	A207231 003	Nov 28, 2016
	6MG	A207231 004	Nov 28, 2016
	8MG	A207231 005	Nov 28, 2016
	10MG	A207231 006	Nov 28, 2016
	12MG	A207231 007	Nov 28, 2016
LUPIN LTD	1MG	A206890 001	May 05, 2022
	2MG	A206890 002	May 05, 2022
	4MG	A206890 003	May 05, 2022
	6MG	A206890 004	May 05, 2022
	8MG	A206890 005	May 05, 2022
	10MG	A206890 006	May 05, 2022
	12MG	A206890 007	May 05, 2022
TARO	1MG	A207098 001	Jul 22, 2019
	2MG	A207098 002	Jul 22, 2019
	4MG	A207098 003	Jul 22, 2019
	6MG	A207098 004	Jul 22, 2019
	8MG	A207098 005	Jul 22, 2019
	10MG	A207098 006	Jul 22, 2019
	12MG	A207098 007	Jul 22, 2019

ILOPROST

SOLUTION; INHALATION

VENTAVIS

ACTELION	20MCG/2ML (10MCG/ML)	N021779 001	Dec 29, 2004
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IMATINIB MESYLATE

CAPSULE; ORAL

GLEEVEC

+ NOVARTIS	EQ 50MG BASE **	N021335 001	May 10, 2001
+	EQ 100MG BASE **	N021335 002	May 10, 2001

TABLET; ORAL

IMATINIB MESYLATE

AMNEAL PHARMS	EQ 100MG BASE	A207495 001	Feb 08, 2019
	EQ 400MG BASE	A207495 002	Feb 08, 2019
BRECKENRIDGE	EQ 100MG BASE	A205990 001	Feb 08, 2019
	EQ 400MG BASE	A205990 002	Feb 08, 2019
HIKMA	EQ 100MG BASE	A207586 001	Jul 13, 2018
	EQ 400MG BASE	A207586 002	Jul 13, 2018

IMIPRAMINE HYDROCHLORIDE

CONCENTRATE; ORAL

IMIPRAMINE HYDROCHLORIDE

NOVARTIS	25MG/ML	A086765 001	
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INJECTABLE; INJECTION

TOFRANIL

NOVARTIS	12.5MG/ML	N011838 002	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

CHARTWELL	10MG	A090441 002	Mar 11, 2010
	25MG	A090441 003	Mar 11, 2010
	50MG	A090441 001	Mar 11, 2010
LEDERLE	10MG	A086269 001	
	25MG	A086267 001	
	50MG	A086268 001	
PAR PHARM	10MG	A089422 001	Jul 14, 1987
	25MG	A089497 001	Jul 14, 1987
ROXANE	10MG	A083799 001	
	25MG	A083799 002	
	50MG	A083799 003	
SANDOZ	10MG	A085200 001	
	25MG	A084869 002	
	50MG	A085133 001	
SUN PHARM INDUSTRIES	10MG	A081048 001	Jun 05, 1990
TEVA	10MG	A083729 001	
	25MG	A083729 004	
	50MG	A083729 003	
USL PHARMA	25MG	A087776 001	Feb 10, 1982
VANGARD	10MG	A088036 001	Nov 03, 1982
	25MG	A087619 001	Feb 09, 1982
	50MG	A087631 001	Jan 04, 1982
WATSON LABS	10MG	A085220 001	
	10MG	A085875 001	
	25MG	A084252 002	
	25MG	A085878 001	
	50MG	A085221 001	
	50MG	A085877 001	
WEST WARD	25MG	A088222 001	May 26, 1983
	50MG	A088223 001	May 26, 1983
JANIMINE			
ABBOTT	10MG	N017895 001	
	25MG	N017895 002	
	50MG	N017895 003	
PRAMINE			
ALRA	10MG	A083827 001	
	25MG	A083827 002	
	50MG	A083827 003	
PRESAMINE			
SANOFI AVENTIS US	10MG	N011836 006	
	25MG	N011836 003	
	50MG	N011836 007	

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

RISING	EQ 75MG HYDROCHLORIDE	A202338 001	Jun 28, 2013
	EQ 100MG HYDROCHLORIDE	A202338 002	Jun 28, 2013
	EQ 125MG HYDROCHLORIDE	A202338 003	Jun 28, 2013
	EQ 150MG HYDROCHLORIDE	A202338 004	Jun 28, 2013
TOFRANIL-PM			
+	SPECGX LLC	EQ 75MG HYDROCHLORIDE **	N017090 001
+		EQ 100MG HYDROCHLORIDE **	N017090 004
+		EQ 125MG HYDROCHLORIDE **	N017090 003
+		EQ 150MG HYDROCHLORIDE **	N017090 002

IMIQUIMOD

CREAM; TOPICAL

ALDARA

+	BAUSCH	5%	N020723 001	Feb 27, 1997
	IMIQUIMOD			
	COSETTE	5%	A200481 001	Apr 18, 2011
	ENCUBE	5%	A091044 001	Feb 28, 2011
	STRIDES PHARMA	5%	A202002 001	Jun 24, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

BAXTER HLTHCARE CORP EQ 5MG BASE/ML
HOSPIRA EQ 5MG BASE/MLA075542 001 May 10, 2000
A074616 001 Aug 03, 1998

INOCOR

SANOFI AVENTIS US EQ 5MG BASE/ML

N018700 001 Jul 31, 1984

INDACATEROL MALEATE

POWDER; INHALATION

ARCAPTA NEOHALER

+ NOVARTIS EQ 75MCG BASE

N022383 001 Jul 01, 2011

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

ANI PHARMS 1.25MG
1.25MG
2.5MG
2.5MGA074498 002 Feb 12, 1998
A075201 001 Dec 04, 1998
A074498 001 Oct 31, 1996
A075201 002 Dec 04, 1998AUROBINDO PHARMA USA 1.25MG
2.5MGA075105 001 Jul 23, 1998
A075105 002 Jul 23, 1998CHARTWELL RX 1.25MG
2.5MGA074594 001 May 23, 1996
A074594 002 May 23, 1996TEVA 1.25MG
2.5MGA074665 001 Apr 04, 1997
A074665 002 Apr 04, 1997WATSON LABS 1.25MG
2.5MGA074585 001 Sep 26, 1996
A074585 002 Sep 26, 1996

LOZOL

+ SANOFI AVENTIS US 1.25MG **
+ 2.5MG **N018538 002 Apr 29, 1993
N018538 001 Jul 06, 1983INDECANIDE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

DECABID

LILLY EQ 50MG BASE
EQ 75MG BASE
EQ 100MG BASEN019693 001 Dec 29, 1989
N019693 002 Dec 29, 1989
N019693 003 Dec 29, 1989INDINAVIR SULFATE

CAPSULE; ORAL

CRIXIVAN

MERCK SHARP DOHME EQ 100MG BASE
+ EQ 200MG BASE
EQ 333MG BASE
+ EQ 400MG BASEN020685 006 Apr 19, 2000
N020685 003 Mar 13, 1996
N020685 005 Dec 17, 1998
N020685 001 Mar 13, 1996INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDICLOR

+ GE HEALTHCARE 2mCi/0.2ML

N019862 001 Dec 29, 1992

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

RENEW PHARMS 10MG/VIAL **
+ 25MG/VIAL
40MG/VIAL **
50MG/VIAL **N011525 003
N011525 001
N011525 004
N011525 002INDOMETHACIN

CAPSULE; ORAL

INDO-LEMMON

TEVA 25MG
50MGA070266 001 Nov 07, 1985
A070267 001 Nov 07, 1985

INDOCIN

+ ZYLA LIFE SCIENCES 25MG **
+ 50MG **N016059 001
N016059 002

INDOMETHACIN

ABLE 25MG
50MG
ANI PHARMS 25MG
50MGA076666 001 Dec 17, 2003
A076666 002 Dec 17, 2003
A071148 001 Mar 18, 1987
A071149 001 Mar 18, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDOMETHACIN

CAPSULE;ORAL

INDOMETHACIN

CYCLE	25MG	A070353	001	Jun 18, 1985
	50MG	A070354	001	Jun 18, 1985
DURAMED PHARMS BARR	25MG	A070326	001	Oct 18, 1985
	50MG	A070327	001	Oct 18, 1985
HALSEY	25MG	A070782	001	Jun 03, 1987
	50MG	A070635	001	Jun 03, 1987
HERITAGE PHARMA	25MG	A070719	001	Feb 12, 1986
	50MG	A070756	001	Feb 12, 1986
HERITAGE PHARMS	25MG	N018851	001	May 18, 1984
	50MG	N018851	002	May 18, 1984
IVAX SUB TEVA PHARMS	25MG	N018730	001	May 04, 1984
	50MG	N018730	002	May 04, 1984
JUBILANT GENERICS	25MG	A205215	001	Aug 25, 2017
	50MG	A205215	002	Aug 25, 2017
MUTUAL PHARM	25MG	A070067	001	Oct 03, 1986
	50MG	A070068	001	Oct 03, 1986
PARKE DAVIS	25MG	N018806	001	Nov 23, 1984
	50MG	N018806	002	Nov 23, 1984
PIONEER PHARMS	25MG	A070813	001	Aug 11, 1986
	50MG	A070592	001	Aug 11, 1986
RISING	25MG	N018858	001	Apr 20, 1984
	50MG	A070624	001	Sep 04, 1985
	50MG	N018858	002	Apr 20, 1984
SUN PHARM INDS INC	25MG	A091401	001	Mar 28, 2013
	50MG	A091401	002	Mar 28, 2013
SUN PHARM INDUSTRIES	25MG	A070900	002	Feb 09, 1987
	50MG	A070900	001	Feb 09, 1987
SUPERPHARM	25MG	A070487	001	Oct 10, 1986
	50MG	A070488	001	Oct 10, 1986
TEVA	25MG	A071342	001	Apr 18, 1988
	50MG	A071343	001	Apr 18, 1988
WATSON LABS	25MG	A070529	001	Oct 18, 1985
	25MG	A070784	001	Aug 20, 1986
	25MG	A072996	001	Jul 31, 1991
	25MG	N018690	001	Jul 31, 1984
	50MG	A070530	001	Oct 18, 1985
	50MG	A070785	001	Aug 20, 1986
	50MG	A071635	001	May 18, 1987
	50MG	A072997	001	Jul 31, 1991
	50MG	N018690	002	Jul 31, 1984

TIVORBEX

+ GENUS	20MG	N204768	001	Feb 24, 2014
+	40MG	N204768	002	Feb 24, 2014

CAPSULE, EXTENDED RELEASE;ORAL

INDOCIN SR

+ ZYLA	75MG **	N018185	001	Feb 23, 1982
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INDOMETHACIN

ABLE	75MG	A076114	001	Feb 06, 2002
AUROBINDO PHARMA	75MG	A204243	001	Dec 27, 2016
INWOOD LABS	75MG	A072410	001	Mar 15, 1989
JUBILANT GENERICS	75MG	A202706	001	Oct 05, 2015
RISING	75MG	A202139	001	Mar 20, 2014
SANDOZ	75MG	A074464	001	May 28, 1998
WATSON LABS INC	75MG	A202572	001	Dec 09, 2013

SUPPOSITORY;RECTAL

INDOCIN

+ ZYLA LIFE SCIENCES	50MG **	N017814	001	Aug 13, 1984
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SUSPENSION;ORAL

INDOMETHACIN

HIKMA	25MG/5ML	A071412	001	Mar 18, 1987
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOCIN

+ RECORDATI RARE EQ 1MG BASE/VIAL ** N018878 001 Jan 30, 1985

INFIGRATINIB PHOSPHATE

CAPSULE; ORAL

TRUSELTIQ

+ HELSINN HLTHCARE 25MG N214622 001 May 28, 2021

+ 100MG N214622 002 May 28, 2021

INGENOL MEBUTATE

GEL; TOPICAL

INGENOL MEBUTATE

PADAGIS ISRAEL 0.015% A209018 001 Jan 07, 2019

0.05% A209019 001 Jan 09, 2019

PICATO

+ LEO LABS 0.015% N202833 001 Jan 23, 2012

+ 0.05% N202833 002 Jan 23, 2012

INULIN

INJECTABLE; INJECTION

INULIN AND SODIUM CHLORIDE

ISO TEX 100MG/ML N002282 001

INVERT SUGAR

INJECTABLE; INJECTION

TRAVERT 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 10GM/100ML N016717 001

IOBENGUANE SULFATE I-131

INJECTABLE; INJECTION

IOBENGUANE SULFATE I 131

PHARMALUCENCE 2.3mCi/ML N020084 001 Mar 25, 1994

IO CETAMIC ACID

TABLET; ORAL

CHOLEBRINE

MALLINCKRODT 750MG N017129 001

IODAMIDE MEGLUMINE

INJECTABLE; INJECTION

RENOVUE-65

BRACCO 65% N017902 001

RENOVUE-DIP

BRACCO 24% N017903 001

IODIPAMIDE MEGLUMINE

INJECTABLE; INJECTION

CHOLOGRAFIN MEGLUMINE

BRACCO 10.3% N009321 007

+ 52% N009321 003

IODIPAMIDE SODIUM

INJECTABLE; INJECTION

CHOLOGRAFIN SODIUM

BRACCO 20% N009321 001

IODIXANOL

INJECTABLE; INJECTION

VISIPAQUE 270

GE HEALTHCARE 55% N020808 001 Aug 29, 1997

IODOHIPPURATE SODIUM I-123

INJECTABLE; INJECTION

NEPHROFLOW

GE HEALTHCARE 1mCi/ML N018289 001 Dec 28, 1984

IODOHIPPURATE SODIUM I-131

INJECTABLE; INJECTION

HIPURAN I 131

MALLINCKRODT 0.25mCi/ML N016666 001

HIPPUTOPE

BRACCO 1-2mCi/VIAL N015419 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IODOHIPPURATE SODIUM I-131

INJECTABLE; INJECTION

IODOHIPPURATE SODIUM I 131

PHARMALUCENCE

0.2mCi/ML

N017313 001

IODOXAMATE MEGLUMINE

INJECTABLE; INJECTION

CHOLOVUE

BRACCO

9.9%

N018077 001

40.3%

N018076 001

IOFETAMINE HYDROCHLORIDE I-123

INJECTABLE; INJECTION

SPECTAMINE

IMP

1mCi/ML

N019432 001 Dec 24, 1987

IOHEXOL

FOR SOLUTION; ORAL

ORALTAG

INTERPHARMA PRAHA AS 9.7GM/BOT

N205383 001 Mar 26, 2015

INJECTABLE; INJECTION

OMNIPAQUE 210

GE HEALTHCARE 45.3%

N018956 006 Jun 30, 1989

SOLUTION; INJECTION, ORAL, RECTAL

OMNIPAQUE 240

GE HEALTHCARE 51.8%

N020608 001 Oct 24, 1995

SOLUTION; URETHRAL

OMNIPAQUE 70

GE HEALTHCARE 15.1%

N018956 007 Jun 01, 1994

IOPAMIDOL

INJECTABLE; INJECTION

IOPAMIDOL

BAXTER HLTHCARE

41%

A074629 001 Nov 06, 1996

51%

A074629 004 Mar 31, 1998

61%

A074629 002 Nov 06, 1996

76%

A074629 003 Nov 06, 1996

HOSPIRA

61%

A074734 001 Dec 10, 1996

76%

A074734 002 Dec 10, 1996

IOPAMIDOL-200

COOK IMAGING

41%

A074881 001 Jul 28, 2000

HOSPIRA

41%

A074898 001 Dec 30, 1997

IOPAMIDOL-200 IN PLASTIC CONTAINER

HOSPIRA

41%

A074636 001 Dec 30, 1997

IOPAMIDOL-250

COOK IMAGING

51%

A074881 002 Jul 28, 2000

FRESENIUS KABI USA

51%

A074679 001 Apr 02, 1997

HOSPIRA

51%

A074898 002 Dec 30, 1997

51%

A075005 001 Feb 24, 1998

IOPAMIDOL-250 IN PLASTIC CONTAINER

HOSPIRA

51%

A074636 002 Dec 30, 1997

IOPAMIDOL-300

ABBVIE

61%

A074638 001 Apr 30, 1997

COOK IMAGING

61%

A074881 003 Jul 28, 2000

FRESENIUS KABI USA

61%

A074679 002 Apr 02, 1997

HOSPIRA

61%

A074898 003 Dec 30, 1997

61%

A075005 002 Feb 24, 1998

IOPAMIDOL-300 IN PLASTIC CONTAINER

HOSPIRA

61%

A074636 003 Dec 30, 1997

61%

A074637 001 Apr 03, 1997

IOPAMIDOL-370

COOK IMAGING

76%

A074881 004 Jul 28, 2000

FRESENIUS KABI USA

76%

A074679 003 Apr 02, 1997

HOSPIRA

76%

A074898 004 Dec 30, 1997

76%

A075005 003 Feb 24, 1998

IOPAMIDOL-370 IN PLASTIC CONTAINER

HOSPIRA

76%

A074636 004 Dec 30, 1997

ISOVUE-128

BRACCO

26%

N018735 005 Oct 21, 1986

ISOVUE-200

BRACCO

41%

N020327 001 Oct 12, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IOPAMIDOL

INJECTABLE; INJECTION

ISOVUE-250

+ BRACCO

51%

N020327 002 Oct 12, 1994

SCANLUX-300

SANOCHEMIA CORP USA

61%

A090394 001 Jun 18, 2010

SCANLUX-370

SANOCHEMIA CORP USA

76%

A090394 002 Jun 18, 2010

IOPANOIC ACID

TABLET; ORAL

TELEPAQUE

GE HEALTHCARE

500MG

N008032 001

IOPHENDYLATE

INJECTABLE; INJECTION

PANTOPAQUE

ALCON

100%

N005319 001

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+ BAYER HLTHCARE

49.9%

N021425 003 Mar 12, 2004

ULTRAVIST 150

+ BAYER HLTHCARE

31.2%

N020220 004 May 10, 1995

ULTRAVIST 240

+ BAYER HLTHCARE

49.9%

N020220 003 May 10, 1995

ULTRAVIST 300 IN PLASTIC CONTAINER

+ BAYER HLTHCARE

62.3%

N020220 005 Nov 18, 2008

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY 30

+ LIEBEL-FLARSHEIM

30%

N016983 001

CONRAY 43

+ LIEBEL-FLARSHEIM

43%

N013295 002

IOTHALAMATE MEGLUMINE; IOTHALAMATE SODIUM

INJECTABLE; INJECTION

VASCORAY

MALLINCKRODT

52%;26%

N016783 001

IOTHALAMATE SODIUM

INJECTABLE; INJECTION

ANGIO-CONRAY

MALLINCKRODT

80%

N013319 001

CONRAY 325

MALLINCKRODT

54.3%

N017685 001

CONRAY 400

MALLINCKRODT

66.8%

N014295 001

IOTROLAN

INJECTABLE; INTRATHECAL

OSMOVIST 190

BAYER HLTHCARE

40.6%

N019580 001 Dec 07, 1989

OSMOVIST 240

BAYER HLTHCARE

51.3%

N019580 002 Dec 07, 1989

IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 160

LIEBEL-FLARSHEIM

34%

N019710 003 Dec 30, 1988

OPTIRAY 240

+ LIEBEL-FLARSHEIM

51%

N019710 002 Dec 30, 1988

51%

N020923 001 May 28, 1998

IOXAGLATE MEGLUMINE; IOXAGLATE SODIUM

INJECTABLE; INJECTION

HEXABRIX

GUERBET

39.3%;19.6%

N018905 002 Jul 26, 1985

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IOXILAN

INJECTABLE; INJECTION

OXILAN-300

GUERBET

62%

N020316 001 Dec 21, 1995

OXILAN-350

GUERBET

73%

N020316 002 Dec 21, 1995

IPODATE CALCIUM

GRANULE; ORAL

ORAGRAFIN CALCIUM

BRACCO

3GM/PACKET

N012968 001

IPODATE SODIUM

CAPSULE; ORAL

BILIVIST

BAYER HLTHCARE

500MG

A087768 001 Aug 11, 1982

ORAGRAFIN SODIUM

BRACCO

500MG

N012967 001

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT

BOEHRINGER INGELHEIM 0.018MG/INH

N019085 001 Dec 29, 1986

SOLUTION; INHALATION

ATROVENT

+ BOEHRINGER INGELHEIM 0.02% **

N020228 001 Sep 29, 1993

IPRATROPIUM BROMIDE

ACTAVIS MID ATLANTIC

0.02%

A075111 001 Apr 22, 1999

APOTEX INC

0.02%

A075441 001 Mar 28, 2001

BAUSCH

0.02%

A075835 001 Oct 15, 2001

LANDELA PHARM

0.02%

A077072 001 Jul 19, 2005

MYLAN SPECIALITY LP

0.02%

A074755 001 Jan 10, 1997

ROXANE

0.02%

A075867 001 Jul 22, 2002

TEVA PHARMS USA

0.02%

A075313 001 Feb 07, 2000

WATSON LABS

0.02%

A076291 001 May 09, 2005

ZENNOVA

0.02%

A075507 001 Jan 19, 2001

SPRAY, METERED; NASAL

ATROVENT

+ BOEHRINGER INGELHEIM 0.021MG/SPRAY **

N020393 001 Oct 20, 1995

+ 0.042MG/SPRAY **

N020394 001 Oct 20, 1995

IPRATROPIUM BROMIDE

AUROBINDO PHARMA USA 0.021MG/SPRAY

A075552 001 Mar 31, 2003

0.042MG/SPRAY

A075553 001 Mar 31, 2003

IRBESARTAN

TABLET; ORAL

IRBESARTAN

AJANTA PHARMA LTD

75MG

A203685 001 Dec 10, 2015

150MG

A203685 002 Dec 10, 2015

300MG

A203685 003 Dec 10, 2015

APOTEX INC

75MG

A200832 001 Oct 15, 2012

150MG

A200832 002 Oct 15, 2012

300MG

A200832 003 Oct 15, 2012

CHARTWELL RX

75MG

A203161 001 Sep 27, 2012

150MG

A203161 002 Sep 27, 2012

300MG

A203161 003 Sep 27, 2012

HIKMA

75MG

A090201 001 Oct 15, 2012

150MG

A090201 002 Oct 15, 2012

300MG

A090201 003 Oct 15, 2012

UPSHER SMITH LABS

75MG

A200461 001 Sep 27, 2012

150MG

A200461 002 Sep 27, 2012

300MG

A200461 003 Sep 27, 2012

WATSON LABS INC

75MG

A090720 001 Oct 12, 2012

150MG

A090720 002 Oct 12, 2012

300MG

A090720 003 Oct 12, 2012

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

CHARTWELL RX	40MG/2ML (20MG/ML)	A078953 001	Apr 15, 2010
	100MG/5ML (20MG/ML)	A078953 002	Apr 15, 2010
CIPLA LTD	40MG/2ML (20MG/ML)	A077219 001	Feb 20, 2008
	100MG/5ML (20MG/ML)	A077219 002	Feb 20, 2008
EMCURE PHARMS LTD	40MG/2ML (20MG/ML)	A200771 001	Feb 14, 2012
	100MG/5ML (20MG/ML)	A200771 002	Feb 14, 2012
FRESENIUS KABI USA	40MG/2ML (20MG/ML)	A078188 001	Feb 27, 2008
	100MG/5ML (20MG/ML)	A078188 002	Feb 27, 2008
PLIVA LACHEMA	40MG/2ML (20MG/ML)	A078122 001	Oct 31, 2008
	100MG/5ML (20MG/ML)	A078122 002	Oct 31, 2008
SANDOZ	40MG/2ML (20MG/ML)	A077994 001	Feb 27, 2008
	40MG/2ML (20MG/ML)	A090137 001	Nov 12, 2009
	100MG/5ML (20MG/ML)	A077994 002	Feb 27, 2008
	100MG/5ML (20MG/ML)	A090137 002	Nov 12, 2009
SUN PHARMA GLOBAL	40MG/2ML (20MG/ML)	A078805 001	Apr 21, 2008
	100MG/5ML (20MG/ML)	A078805 002	Apr 21, 2008
TEVA PHARMS USA	40MG/2ML (20MG/ML)	A090101 002	Feb 27, 2008
	100MG/5ML (20MG/ML)	A090101 003	Feb 27, 2008

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

BETA-2

NEPHRON	1%	A086711 001	
BRONKOSOL			
SANOFI AVENTIS US	0.25%	N012339 009	
	1%	N012339 008	

ISOETHARINE HYDROCHLORIDE

ALPHARMA US PHARMS	1%	A087101 001	
+ ASTRAZENECA	0.062%	A087937 001	Nov 15, 1982
	0.062%	A089614 001	Jun 13, 1991
+	0.125%	A087938 001	Nov 15, 1982
	0.125%	A089615 001	Jun 13, 1991
+	0.167%	A088470 001	Mar 14, 1984
	0.167%	A089616 001	Jun 13, 1991
+	0.2%	A088471 001	Mar 14, 1984
	0.2%	A089617 001	Jun 13, 1991
+	0.25%	A088472 001	Mar 14, 1984
	0.25%	A089618 001	Jun 13, 1991
BAXTER HLTHCARE	0.08%	A088144 001	Jul 29, 1983
	0.14%	A088145 001	Mar 26, 1984
	0.25%	A088146 001	Aug 01, 1983
DEY	0.08%	A088187 001	Dec 03, 1982
	0.1%	A087389 001	
	0.17%	A087390 001	
	0.25%	A088188 001	Dec 03, 1982
+	1%	A086763 001	
INTL MEDICATION	0.077%	A086651 001	
	0.08%	A086651 002	
	0.1%	A086651 003	
	0.143%	A086651 004	
	0.167%	A086651 005	
	0.2%	A086651 006	
	0.25%	A086651 007	
	1%	A086651 008	
PARKE DAVIS	0.5%	A085997 001	
	1%	A085889 001	
ROXANE	0.1%	A087396 001	
	0.125%	A087025 001	
	0.167%	A088226 001	Sep 16, 1983
	0.2%	A087324 001	
	0.25%	A088275 001	Jun 03, 1983
	1%	A086899 001	
ISOETHARINE HYDROCHLORIDE S/F			
DEY	0.08%	A089817 001	Nov 22, 1988
	0.1%	A089818 001	Nov 22, 1988
	0.17%	A089819 001	Nov 22, 1988
	0.25%	A089820 001	Nov 22, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

ISOETHARINE HYDROCHLORIDE S/F
1%

A089252 001 Sep 15, 1986

ISOETHARINE MESYLATEAEROSOL, METERED; INHALATION
BRONKOMETERSANOFI AVENTIS US 0.34MG/INH
ISOETHARINE MESYLATE
ALPHARMA US PHARMS 0.34MG/INH

N012339 007

A087858 001 Aug 21, 1984

ISOFLURANE

LIQUID; INHALATION

ISOFLURANE

HOSPIRA 99.9%
WATSON LABS INC 99.9%

A074097 001 Jan 25, 1993

A074393 001 May 12, 1995

ISOFLUROPHATE

OINTMENT; OPHTHALMIC

FLOROPRYL

MERCK 0.025%

N010656 001

ISONIAZID

INJECTABLE; INJECTION

NYDRAZID

SANDOZ 100MG/ML **
RIMIFON
ROCHE 25MG/ML
100MG/ML

N008662 001

N008420 002

N008420 003

SYRUP; ORAL

ISONIAZID

CHARTWELL RX 50MG/5ML

A081118 001 Jul 21, 1997

LANIAZID

LANNETT 50MG/5ML

A089243 001 Feb 03, 1986

RIMIFON

ROCHE 50MG/5ML

N008420 001

TABLET; ORAL

DOW-ISONIAZID

DOW PHARM 300MG

A080330 002

HYZYD

MEDPOINTE PHARM HLC 100MG

A080134 003

300MG

A080134 004

INH

NOVARTIS 300MG

A080935 001

ISONIAZID

DURAMED PHARMS BARR 100MG

A088231 001 Mar 17, 1983

300MG

A088119 001 Mar 17, 1983

+ EPIC PHARMA LLC 100MG **

N008678 002

+ 300MG **

N008678 003

HALSEY 50MG

A083632 001

HIKMA INTL PHARMS 100MG

A080212 001

300MG

A087425 001

IMPAX LABS 100MG

A080153 001

IVAX SUB TEVA PHARMS 100MG

A080270 001

300MG

A083610 001

LILLY 100MG

N008499 002

300MG

N008499 003

MK LABS 100MG

A080941 001

NEXGEN PHARMA INC 100MG

A084050 001

PANRAY 50MG

N008428 001

100MG

N008428 002

300MG

N008428 003

PERRIGO 100MG

A083060 001

PHARMAVITE 100MG

A085091 001

PHOENIX LABS NY 50MG

A080368 001

100MG

A080368 002

PUREPAC PHARM 50MG

A080132 003 Jul 14, 1982

100MG

A080132 004 Jul 14, 1982

SUN PHARM INDUSTRIES 100MG

A080136 001

300MG

A083633 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISONIAZID

TABLET; ORAL

ISONIAZID

WATSON LABS

50MG

A080522 001

100MG

A080401 001

100MG

A080523 001

100MG

A085790 001

300MG

A080521 001

300MG

A083178 001

300MG

A085784 001

WHITEWORTH TOWN PLSN

100MG

A080120 002

LANIAZID

CHARTWELL MOLECULAR

50MG

A080140 001

100MG

A080140 002

300MG

A089776 001 Jun 13, 1988

NYDRAZID

BRISTOL MYERS SQUIBB

100MG

N008392 003

STANOZIDE

EVERYLIFE

100MG

A080126 001

300MG

A080126 002

ISONIAZID; PYRAZINAMIDE; RIFAMPIN

TABLET; ORAL

RIFATER

+ SANOFI AVENTIS US

50MG; 300MG; 120MG

N050705 001 May 31, 1994

ISONIAZID; RIFAMPIN

CAPSULE; ORAL

RIFAMATE

SANOFI AVENTIS US

150MG; 300MG

A061884 001

RIFAMPIN AND ISONIAZID

HIKMA INTL PHARMS

150MG; 300MG

A065221 001 Jul 29, 2005

ISOPROPAMIDE IODIDE

TABLET; ORAL

DARBID

GLAXOSMITHKLINE

EQ 5MG BASE

N010744 001

ISOPROTERENOL HYDROCHLORIDE

AEROSOL, METERED; INHALATION

ISOPROTERENOL HYDROCHLORIDE

3M

0.12MG/INH

N010375 004

ALPHARMA US PHARMS

0.12MG/INH

A085904 001

ISUPREL

SANOFI AVENTIS US

0.103MG/INH

N011178 001

DISC; INHALATION

NORISODRINE AEROTROL

ABBOTT

0.25%

N016814 001

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

ABRAXIS PHARM

0.2MG/ML

A083431 001

AM REGENT

0.2MG/ML

A214775 001 Oct 19, 2022

BAXTER HLTHCARE

0.2MG/ML

A083486 001

CIPLA

0.2MG/ML

A211738 001 Jun 28, 2019

HOSPIRA

0.02MG/ML

A083283 001

0.2MG/ML

A083346 001

INTL MEDICATION

0.2MG/ML

A083724 001

MYLAN LABS LTD

0.2MG/ML

A212573 001 Sep 06, 2022

ZYDUS PHARMS

0.2MG/ML

A215557 001 Apr 14, 2023

ISUPREL

+ BAUSCH

0.2MG/ML

N010515 001

SOLUTION; INHALATION

AEROLONE

LILLY

0.25%

N007245 001

ISOPROTERENOL HYDROCHLORIDE

ARMOUR PHARM

0.031%

A087935 001 Nov 18, 1982

0.062%

A087936 001 Nov 18, 1982

DEY

0.5%

A086764 001 Jan 04, 1982

PARKE DAVIS

0.25%

A085994 001

0.5%

A085540 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOPROTERENOL HYDROCHLORIDE

SOLUTION; INHALATION

ISUPREL

SANOFI AVENTIS US

0.5%

N006327 002

1%

N006327 003

VAPO-ISO

FISONS

0.5%

N016813 001

TABLET; RECTAL, SUBLINGUAL

ISUPREL

SANOFI AVENTIS US

10MG

N006328 001

15MG

N006328 002

ISOPROTERENOL HYDROCHLORIDE; PHENYLEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

DUO-MEDIHALER

3M

0.16MG/INH; 0.24MG/INH

N013296 001

ISOPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

MEDIHALER-ISO

3M

0.08MG/INH

N010375 003

POWDER; INHALATION

NORISODRINE

ABBVIE

10%

N006905 003

25%

N006905 002

ISOSORBIDE

SOLUTION; ORAL

ISMOTIC

ALCON

100GM/220ML

N017063 001

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL

DILATRATE-SR

+ AUXILIUM PHARMS LLC

40MG

N019790 001 Sep 02, 1988

ISORDIL

WYETH AYERST

40MG

N012882 002 Jul 29, 1988

TABLET; ORAL

ISORDIL

+ BAUSCH

10MG **

N012093 002 Jul 29, 1988

+

20MG **

N012093 006 Jul 29, 1988

+

30MG **

N012093 005 Jul 29, 1988

ISOSORBIDE DINITRATE

ANI PHARMS

10MG

A086032 001 Jan 07, 1988

HIKMA INTL PHARMS

30MG

A040591 001 Jan 10, 2007

SUN PHARM INDUSTRIES

5MG

A086166 002 Sep 19, 1986

10MG

A086169 001 Sep 19, 1986

20MG

A086167 001 Sep 19, 1986

30MG

A087564 001 Sep 18, 1986

SUPERPHARM

5MG

A089190 001 Feb 17, 1987

10MG

A089191 001 Feb 17, 1987

20MG

A089192 001 Feb 17, 1987

WATSON LABS

5MG

A086034 001 Jan 06, 1988

SORBITRATE

ASTRAZENECA

5MG

N016192 001 Apr 01, 1996

10MG

N016192 002 Apr 01, 1996

20MG

A086405 002 Aug 21, 1990

30MG

A088124 001 Aug 21, 1990

40MG

A088125 001 Aug 21, 1990

TABLET; SUBLINGUAL

ISORDIL

+ BIOVAIL

2.5MG **

N012940 004 Jul 29, 1988

+

5MG **

N012940 003 Jul 29, 1988

+

10MG **

N012940 005 Jul 29, 1988

ISOSORBIDE DINITRATE

HIKMA INTL PHARMS

2.5MG

A086054 001 Oct 29, 1987

5MG

A086055 001 Nov 02, 1987

SANDOZ

2.5MG

A086225 001 Feb 19, 1988

5MG

A086222 001 Feb 19, 1988

SUN PHARM INDUSTRIES

2.5MG

A084204 001 Sep 18, 1986

5MG

A086168 001 Sep 18, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOSORBIDE DINITRATE

TABLET;SUBLINGUAL

ISOSORBIDE DINITRATE

	10MG	A087545 001	Sep 18, 1986
WATSON LABS	2.5MG **	A086033 001	Feb 26, 1988
WATSON LABS TEVA	5MG **	A086031 001	Sep 29, 1987

SORBITRATE

ASTRAZENECA	2.5MG	N016191 002	Apr 01, 1996
	5MG	N016191 001	Apr 01, 1996

TABLET, CHEWABLE;ORAL

SORBITRATE

ASTRAZENECA	5MG	N016776 002	Apr 01, 1996
	10MG	N016776 003	Apr 01, 1996

TABLET, EXTENDED RELEASE;ORAL

ISORDIL

WYETH AYERST	40MG	N012882 001	Jul 29, 1988
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ISOSORBIDE DINITRATE

IMPAX LABS INC	40MG	A040723 001	Mar 17, 2008
SUN PHARM INDS INC	40MG	A040009 001	Dec 30, 1998

ISOSORBIDE MONONITRATE

TABLET;ORAL

ISMO

PROMIUS PHARMA	20MG	N019091 001	Dec 30, 1991
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ISOSORBIDE MONONITRATE

ANI PHARMS	20MG	A075147 001	Nov 27, 1998
HIKMA PHARMS	20MG	A075361 001	Oct 05, 2000

TABLET, EXTENDED RELEASE;ORAL

IMDUR

+ SCHERING PLOUGH	30MG **	N020225 001	Aug 12, 1993
	60MG **	N020225 002	Aug 12, 1993
	120MG **	N020225 003	Mar 30, 1995

ISOSORBIDE MONONITRATE

ACCORD HLTHCARE	30MG	A209684 001	Oct 24, 2017
	60MG	A209684 002	Oct 24, 2017
	120MG	A209684 003	Oct 24, 2017
ACTAVIS ELIZABETH	30MG	A075306 001	Dec 31, 1998
	60MG	A075306 002	Dec 31, 1998
ALKERMES GAINESVILLE	60MG	A075041 001	Sep 22, 1998
HIKMA INTL PHARMS	30MG	A076813 002	Mar 30, 2006
	60MG	A076813 001	Jan 07, 2005
IVAX SUB TEVA PHARMS	30MG	A075448 002	Aug 07, 2001
	60MG	A075448 001	Jun 19, 2000
	120MG	A075448 003	Aug 07, 2001
SHANDONG	120MG	A214115 001	Apr 29, 2022
SKYEPHARMA AG	60MG	A075166 001	Oct 07, 1999
STRIDES PHARMA	30MG	A090598 001	Aug 11, 2010
	60MG	A090598 002	Aug 11, 2010
	120MG	A090598 003	Aug 11, 2010

ISOSULFAN BLUE

INJECTABLE;INJECTION

ISOSULFAN BLUE

BELOTECA	1%	A210714 001	Jan 16, 2019
EUGIA PHARMA	1%	A206831 001	Feb 02, 2016
FRESENIUS KABI USA	1%	A211869 001	Sep 08, 2022
SOMERSET THERAPS LLC	1%	A210558 001	Jul 12, 2019

LYMPHAZURIN

+ COVIDIEN	1% **	N018310 001	
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ISOTRETINOIN

CAPSULE;ORAL

ABSORICA LD

+ SUN PHARM	20MG	N211913 003	Nov 05, 2019
	28MG	N211913 005	Nov 05, 2019

ACCUATANE

+ HOFFMANN LA ROCHE	10MG **	N018662 002	May 07, 1982
	20MG **	N018662 004	Mar 28, 1983
	40MG **	N018662 003	May 07, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOTRETINOIN

CAPSULE;ORAL

SOTRET

SUN PHARM INDS LTD	10MG	A076041 001	Dec 24, 2002
	20MG	A076041 002	Dec 24, 2002
	30MG	A076503 001	Jun 20, 2003
	40MG	A076041 003	Dec 24, 2002

ISRADIPINE

CAPSULE;ORAL

DYNACIRC

+ SMITHKLINE BEECHAM	2.5MG **	N019546 001	Dec 20, 1990
+	5MG **	N019546 002	Dec 20, 1990

TABLET, EXTENDED RELEASE;ORAL

DYNACIRC CR

+ GLAXOSMITHKLINE LLC	5MG **	N020336 001	Jun 01, 1994
+	10MG **	N020336 002	Jun 01, 1994

ISRADIPINE

MYLAN

	5MG	A201067 001	Nov 27, 2015
	10MG	A201067 002	Nov 27, 2015

ITRACONAZOLE

CAPSULE;ORAL

ITRACONAZOLE

JUBILANT GENERICS	100MG	A203445 001	Feb 23, 2017
MYLAN PHARMS INC	100MG	A200463 001	Jul 20, 2012
PAR PHARM INC	100MG	A205724 001	Dec 13, 2016
STRIDES PHARMA	100MG	A206410 001	Jul 02, 2019

INJECTABLE; INJECTION

SPORANOX

JANSSEN PHARMS	10MG/ML	N020966 001	Mar 30, 1999
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SOLUTION;ORAL

ITRACONAZOLE

APOTEX	10MG/ML	A208481 001	Aug 02, 2019
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TABLET;ORAL

ONMEL

+ SEBELA IRELAND LTD	200MG	N022484 001	Apr 29, 2010
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IVABRADINE HYDROCHLORIDE

TABLET;ORAL

IVABRADINE HYDROCHLORIDE

INGENUS PHARMS LLC	EQ 5MG BASE	A214051 001	Dec 30, 2021
	EQ 7.5MG BASE	A214051 002	Dec 30, 2021

IVERMECTIN

LOTION;TOPICAL

IVERMECTIN

TEVA PHARMS USA	0.5%	A212485 001	Mar 21, 2022
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TABLET;ORAL

STROMEKTOL

+ MERCK SHARP DOHME	6MG **	N050742 001	Nov 22, 1996
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KANAMYCIN SULFATE

CAPSULE;ORAL

KANTREX

APOTHECON	EQ 500MG BASE	A060516 001	
	EQ 500MG BASE	A061911 001	
	EQ 500MG BASE	A062726 001	Mar 06, 1987

INJECTABLE; INJECTION

KANAMYCIN

HIKMA	EQ 75MG BASE/2ML	A062324 001	
	EQ 500MG BASE/2ML	A062324 002	
	EQ 1GM BASE/3ML	A062324 003	

KANAMYCIN SULFATE

ABRAXIS PHARM	EQ 75MG BASE/2ML	A062504 001	Apr 05, 1984
	EQ 500MG BASE/2ML	A062504 002	Apr 05, 1984
	EQ 1GM BASE/3ML	A062504 003	Apr 05, 1984
FRESENIUS KABI USA	EQ 500MG BASE/2ML	A065111 001	Dec 17, 2002
	EQ 1GM BASE/3ML	A065111 002	Dec 17, 2002
INTL MEDICATION	EQ 500MG BASE/2ML	A062466 001	Sep 30, 1983
	EQ 1GM BASE/3ML	A062466 002	Sep 30, 1983
LOCH	EQ 75MG BASE/2ML	A063021 001	Jul 31, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KANAMYCIN SULFATE

INJECTABLE; INJECTION

KANAMYCIN SULFATE

	EQ 500MG BASE/2ML	A063022 001	Jul 31, 1992
	EQ 1GM BASE/3ML	A063025 001	Jul 31, 1992
PHARMAFAIR	EQ 75MG BASE/2ML	A062668 001	May 07, 1987
	EQ 500MG BASE/2ML	A062672 001	May 07, 1987
	EQ 1GM BASE/3ML	A062669 001	May 07, 1987
SOLOPAK	EQ 75MG BASE/2ML	A062605 003	Feb 26, 1986
	EQ 500MG BASE/2ML	A062605 001	Feb 26, 1986
	EQ 1GM BASE/3ML	A062605 002	Feb 26, 1986
WARNER CHILCOTT	EQ 1GM BASE/3ML	A063092 001	Oct 11, 1989
WATSON LABS	EQ 1GM BASE/3ML	A062520 003	May 09, 1985
KANTREX			
APOTHECON	EQ 75MG BASE/2ML	A061655 003	
	EQ 75MG BASE/2ML	A061901 003	
	EQ 75MG BASE/2ML	A062564 001	Sep 21, 1984
	EQ 500MG BASE/2ML	A061655 001	
	EQ 500MG BASE/2ML	A061901 001	
	EQ 500MG BASE/2ML	A062564 002	Sep 21, 1984
	EQ 1GM BASE/3ML	A061655 002	
	EQ 1GM BASE/3ML	A061901 002	
	EQ 1GM BASE/3ML	A062564 003	Sep 21, 1984
KLEBCIL			
KING PHARMS	EQ 75MG BASE/2ML	A062170 001	
	EQ 500MG BASE/2ML	A062170 002	
	EQ 1GM BASE/3ML	A062170 003	

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETAMINE HYDROCHLORIDE

FRESENIUS KABI USA	EQ 10MG BASE/ML	A215808 001	Jan 13, 2023
	EQ 50MG BASE/ML	A215808 002	Jan 13, 2023

KETOCONAZOLE

CREAM; TOPICAL

NIZORAL

+ JANSSEN PHARMA	2% **	N019084 001	Dec 31, 1985
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GEL; TOPICAL

XOLEGEL

+ ALMIRALL	2%	N021946 001	Jul 28, 2006
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SHAMPOO; TOPICAL

NIZORAL

+ JANSSEN PHARMS	2% **	N019927 001	Aug 31, 1990
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SUSPENSION; ORAL

NIZORAL

JANSSEN PHARMA	100MG/5ML	A070767 001	Nov 07, 1986
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TABLET; ORAL

KETOCONAZOLE

AAIPHARMA LLC	200MG	A075341 001	Jul 27, 1999
AUROBINDO PHARMA USA	200MG	A075597 001	Dec 23, 1999
CHARTWELL RX	200MG	A074971 001	Jun 15, 1999
HERITAGE PHARMA	200MG	A075362 001	Jun 15, 1999
SUN PHARM INDUSTRIES	200MG	A075314 001	Jun 15, 1999
TEVA	200MG	A075273 001	Jun 15, 1999

NIZORAL

+ JANSSEN PHARMS	200MG **	N018533 001	
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KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

RISING	50MG	A074024 001	Dec 29, 1995
	50MG	A074035 002	Dec 31, 1996
	75MG	A074024 002	Dec 29, 1995
	75MG	A074035 003	Dec 31, 1996
TEVA	25MG	A073515 001	Dec 22, 1992
	75MG	A073517 001	Dec 22, 1992

ORUDIS

+ WYETH AYERST	25MG **	N018754 001	Jul 31, 1987
+	50MG **	N018754 002	Jan 09, 1986
+	75MG **	N018754 003	Jan 09, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOPROFEN

CAPSULE, EXTENDED RELEASE;ORAL

KETOPROFEN

ACTAVIS LABS FL INC	100MG	A075270 002	Mar 24, 1999
	150MG	A075270 003	Mar 24, 1999
	200MG	A075270 001	Mar 24, 1999
ALKERMES GAINESVILLE	200MG	A074879 001	Dec 10, 1997
MYLAN	100MG	A075679 003	Feb 20, 2002
	150MG	A075679 002	Feb 20, 2002

ORUVAIL

+ WYETH PHARMS INC	100MG **	N019816 003	Feb 08, 1995
+	150MG **	N019816 002	Feb 08, 1995
+	200MG **	N019816 001	Sep 24, 1993

FILM;ORAL

NEXCEDE

NOVARTIS	12.5MG	N022470 001	Nov 25, 2009
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TABLET;ORAL

ACTRON

BAYER	12.5MG	N020499 001	Oct 06, 1995
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KETOPROFEN

PERRIGO	12.5MG	A075364 001	Feb 07, 2002
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ORUDIS KT

+ WYETH CONS	12.5MG **	N020429 001	Oct 06, 1995
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KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

AMPHASTAR PHARM	15MG/ML	A076209 001	Jul 21, 2004
	30MG/ML	A076209 002	Jul 21, 2004
APOTEX INC	30MG/ML	A075626 001	Jul 24, 2001
	30MG/ML	A077201 001	Oct 14, 2005
APOTHECON	15MG/ML	A075348 001	Nov 28, 2000
	30MG/ML	A075348 002	Nov 28, 2000
BAXTER HLTHCARE CORP	15MG/ML	A075631 002	Jun 29, 2001
	30MG/ML	A075631 001	Jun 29, 2001
BEDFORD	15MG/ML	A075230 002	Oct 25, 1999
	30MG/ML	A075230 001	Oct 25, 1999
EUGIA PHARMA	15MG/ML	A212939 001	Oct 20, 2020
	30MG/ML	A212939 002	Oct 20, 2020
GLAND PHARMA LTD	15MG/ML	A076722 001	Jul 27, 2004
	30MG/ML	A076722 002	Jul 27, 2004
HIKMA	15MG/ML **	A075222 001	Apr 26, 1999
	15MG/ML	A075299 001	Nov 03, 1999
	30MG/ML **	A075222 002	Apr 26, 1999
	30MG/ML **	A075228 001	Apr 26, 1999
	30MG/ML	A075299 002	Nov 03, 1999
HOSPIRA	15MG/ML	A074801 001	Jun 05, 1997
	15MG/ML	A074993 001	Jan 27, 1999
	30MG/ML	A074801 002	Jun 05, 1997
LUITPOLD	15MG/ML	A078145 001	Jan 14, 2008
	30MG/ML	A078145 002	Jan 14, 2008
SANDOZ	15MG/ML	A076271 001	Oct 06, 2004
STERISCIENCE	15MG/ML	A078299 001	Jul 16, 2007
	15MG/ML	A201155 001	Aug 04, 2014
	30MG/ML	A078299 002	Jul 16, 2007
	30MG/ML	A201155 002	Aug 04, 2014
WOCKHARDT BIO AG	30MG/ML	A077943 001	Mar 27, 2007

TORADOL

+ ROCHE PALO	15MG/ML **	N019698 001	Nov 30, 1989
+	30MG/ML **	N019698 002	Nov 30, 1989

SOLUTION/DROPS;OPHTHALMIC

ACULAR PRESERVATIVE FREE

ALLERGAN	0.5%	N020811 001	Nov 03, 1997
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KETOROLAC TROMETHAMINE

AKORN	0.4%	A078399 001	Nov 05, 2009
	0.45%	A203376 001	Feb 10, 2014
CHARTWELL RX	0.5%	A078434 001	Nov 05, 2009
EUGIA PHARMA	0.4%	A205191 001	Nov 15, 2018
	0.5%	A205190 001	Dec 03, 2020
SANDOZ	0.4%	A078721 001	Nov 05, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOROLAC TROMETHAMINE

TABLET;ORAL

KETOROLAC TROMETHAMINE

CHARTWELL RX	10MG	A074790	001	Jun 26, 1997
PLIVA	10MG	A075284	001	Jun 23, 1999
WATSON LABS	10MG	A074955	001	Sep 19, 1997

TORADOL

+ ROCHE PALO	10MG **	N019645	001	Dec 20, 1991
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KETOTIFEN FUMARATE

DRUG-ELUTING CONTACT LENS;OPHTHALMIC

ACUVUE THERAVISION WITH KETOTIFEN

+ JOHNSON JOHNSON VISN EQ 19MCG BASE		N022388	001	Feb 25, 2022
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SOLUTION/DROPS;OPHTHALMIC

ZADITOR

+ ALCON PHARMA	EQ 0.025% BASE **	N021066	002	Oct 19, 2006
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KRYPTON, KR-81M

GAS;INHALATION

MPI KRYPTON 81M GENERATOR

GE HEALTHCARE	N/A	N018088	001	
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L-GLUTAMINE

FOR SOLUTION;ORAL

NUTRESTORE

+ EMMAUS MEDCL	5GM/PACKET	N021667	001	Jun 10, 2004
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LABETALOL HYDROCHLORIDE

INJECTABLE;INJECTION

LABETALOL HYDROCHLORIDE

APOTHECON	5MG/ML	A075355	001	Nov 29, 1999
HOSPIRA	5MG/ML	A075242	001	Sep 30, 1999
RISING	5MG/ML	A075524	001	Nov 29, 1999
STERISCIENCE	5MG/ML	A079134	001	Feb 03, 2010

NORMODYNE

+ SCHERING	5MG/ML **	N018686	001	Aug 01, 1984
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TRANDATE

+ SEBELA IRELAND LTD	5MG/ML **	N019425	001	Dec 31, 1985
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TABLET;ORAL

LABETALOL HYDROCHLORIDE

APOTHECON	100MG	A075223	001	Nov 20, 1998
	200MG	A075223	002	Nov 20, 1998
	300MG	A075223	003	Nov 20, 1998
TEVA	100MG	A074989	001	Sep 30, 1998
	200MG	A074989	002	Sep 30, 1998
	300MG	A074989	003	Sep 30, 1998
UNICHEM	100MG	A212719	001	Aug 08, 2022
	200MG	A212719	002	Aug 08, 2022
	300MG	A212719	003	Aug 08, 2022

NORMODYNE

+ SCHERING	100MG **	N018687	001	Aug 31, 1987
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+	200MG **	N018687	002	Aug 01, 1984
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+	300MG **	N018687	003	Aug 01, 1984
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+	400MG **	N018687	004	Aug 01, 1984
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TRANDATE

+ ALVOGEN	100MG	N018716	001	May 24, 1985
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+	200MG	N018716	002	Aug 01, 1984
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+	300MG	N018716	003	Aug 01, 1984
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+	400MG **	N018716	004	Aug 01, 1984
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LACOSAMIDE

TABLET;ORAL

LACOSAMIDE

ACCORD HLTHCARE	50MG	A205011	001	Jul 12, 2022
	100MG	A205011	002	Jul 12, 2022
	150MG	A205011	003	Jul 12, 2022
	200MG	A205011	004	Jul 12, 2022
ACTAVIS LABS FL INC	50MG	A204855	001	Jan 05, 2023
	100MG	A204855	002	Jan 05, 2023
	150MG	A204855	003	Jan 05, 2023
	200MG	A204855	004	Jan 05, 2023
APOTEX	50MG	A214567	001	Sep 23, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LACOSAMIDETABLET; ORAL
LACOSAMIDE

100MG	A214567 002	Sep 23, 2022
150MG	A214567 003	Sep 23, 2022
200MG	A214567 004	Sep 23, 2022

LACTITOL

FOR SOLUTION; ORAL

PIZENSY

+ BRAINTREE LABS 10GM

N211281 001 Feb 12, 2020

LACTULOSE

SOLUTION; ORAL

CHRONULAC

+ SANOFI AVENTIS US 10GM/15ML **

N017884 001

CONSTILAC

ALRA 10GM/15ML

A071054 001 Jul 26, 1988

CONSTULOSE

ACTAVIS MID ATLANTIC 10GM/15ML

A070288 001 Aug 15, 1988

DUPHALAC

SOLVAY 10GM/15ML

A072372 001 Mar 22, 1989

EVALOSE

TEVA PHARMS 10GM/15ML

A073497 001 May 28, 1993

LACTULOSE

ANI PHARMS 10GM/15ML

A078430 001 Nov 28, 2007

APOZEAL PHARMS 10GM/15ML

A207786 001 Jun 11, 2018

HIKMA 10GM/15ML

A073591 001 May 29, 1992

MORTON GROVE 10GM/15ML

A071841 001 Sep 22, 1988

PACO 10GM/15ML

A073160 001 Aug 25, 1992

VISTAPHARM 10GM/15ML

A074138 001 Sep 30, 1992

LAXILOSE

NOSTRUM LABS 10GM/15ML

A073686 001 May 28, 1993

SOLUTION; ORAL, RECTAL

ACILAC

NOSTRUM LABS 10GM/15ML

A073685 001 May 28, 1993

CEPHULAC

+ SANOFI AVENTIS US 10GM/15ML **

N017657 001

CHOLAC

ALRA 10GM/15ML

A071331 001 Jul 26, 1988

ENULOSE

ACTAVIS MID ATLANTIC 10GM/15ML

A071548 001 Aug 15, 1988

GENERLAC

MORTON GROVE 10GM/15ML

A071842 001 Sep 27, 1988

HEPTALAC

TEVA PHARMS 10GM/15ML

A073504 001 May 28, 1993

LACTULOSE

ANI PHARMS 10GM/15ML

A090426 001 Nov 21, 2008

APOZEAL PHARMS 10GM/15ML

A203762 001 Mar 27, 2015

PACO 10GM/15ML

A072029 001 Aug 25, 1992

ROXANE 10GM/15ML

A073590 001 May 29, 1992

SOLVAY 10GM/15ML

N017906 001

PORTALAC

SOLVAY 10GM/15ML

A072374 001 Mar 22, 1989

LAMIVUDINE

TABLET; ORAL

LAMIVUDINE

AUROBINDO PHARMA LTD 150MG

A202032 001 Nov 17, 2011

300MG

A202032 002 Nov 17, 2011

AUROBINDO PHARMA USA 100MG

A204002 001 Dec 31, 2014

MYLAN 150MG

A204528 001 Mar 04, 2016

300MG

A204528 002 Mar 04, 2016

LAMIVUDINE; NEVIRAPINE; ZIDOVUDINE

TABLET; ORAL

LAMIVUDINE, NEVIRAPINE AND ZIDOVUDINE

+ MICRO LABS 150MG; 200MG; 300MG

N205626 001 Aug 13, 2018

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LAMIVUDINE; RALTEGRAVIR POTASSIUM

TABLET;ORAL

DUTREBIS

MERCK SHARP DOHME	150MG;EQ 300MG BASE	N206510	001	Feb 06, 2015
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LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

+ AUROBINDO PHARMA LTD	300MG;300MG	N022344	001	May 15, 2018
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TEMIXYS

+ CHARTWELL RX	300MG;300MG	N211284	001	Nov 16, 2018
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LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

LAMIVUDINE AND ZIDOVUDINE

AUROBINDO PHARMA LTD	150MG;300MG	A202418	001	May 15, 2012
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CHARTWELL RX	150MG;300MG	A079081	001	May 25, 2011
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MYLAN	150MG;300MG	A204005	001	Aug 28, 2014
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MYLAN LABS LTD	150MG;300MG	A079079	001	Aug 12, 2019
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PHARMACARE	150MG;300MG	N022018	001	Mar 17, 2017
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LAMOTRIGINE

TABLET;ORAL

LAMICTAL

+ GLAXOSMITHKLINE LLC	50MG **	N020241	006	Dec 27, 1994
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+	250MG **	N020241	004	Dec 27, 1994
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LAMOTRIGINE

ACTAVIS TOTOWA	25MG	A078669	001	Apr 08, 2011
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	100MG	A078669	002	Apr 08, 2011
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	150MG	A078669	003	Apr 08, 2011
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	200MG	A078669	004	Apr 08, 2011
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CHARTWELL MOLECULAR	25MG	A077783	001	Nov 01, 2010
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	100MG	A077783	002	Nov 01, 2010
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	150MG	A077783	003	Nov 01, 2010
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	200MG	A077783	004	Nov 01, 2010
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GRANULES	25MG	A078982	001	Jan 27, 2009
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	100MG	A078982	002	Jan 27, 2009
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	150MG	A078982	003	Jan 27, 2009
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	200MG	A078982	004	Jan 27, 2009
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HIKMA PHARMS	25MG	A078134	001	Apr 19, 2011
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	100MG	A078134	002	Apr 19, 2011
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	150MG	A078134	003	Apr 19, 2011
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	200MG	A078134	004	Apr 19, 2011
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MYLAN	25MG	A077428	001	Jan 27, 2009
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	100MG	A077428	002	Jan 27, 2009
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	150MG	A077428	003	Jan 27, 2009
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	200MG	A077428	004	Jan 27, 2009
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MYLAN LABS LTD	25MG	A078443	001	Feb 11, 2009
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	100MG	A078443	002	Feb 11, 2009
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	150MG	A078443	003	Feb 11, 2009
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	200MG	A078443	004	Feb 11, 2009
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RISING	25MG	A077420	001	Jan 27, 2009
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	100MG	A077420	002	Jan 27, 2009
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	150MG	A077420	003	Jan 27, 2009
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	200MG	A077420	004	Jan 27, 2009
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ROXANE	25MG	A077392	001	Jan 27, 2009
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	100MG	A077392	002	Jan 27, 2009
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	150MG	A077392	003	Jan 27, 2009
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	200MG	A077392	004	Jan 27, 2009
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SANDOZ	25MG	A078645	001	Jan 27, 2009
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	100MG	A078645	002	Jan 27, 2009
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	150MG	A078645	003	Jan 27, 2009
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	200MG	A078645	004	Jan 27, 2009
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TEVA	25MG	A076388	001	Aug 30, 2006
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	100MG	A076388	002	Aug 30, 2006
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	150MG	A076388	003	Aug 30, 2006
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	200MG	A076388	004	Aug 30, 2006
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ZENNOVA	25MG	A078310	001	Feb 04, 2009
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	100MG	A078310	002	Feb 04, 2009
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	150MG	A078310	003	Feb 04, 2009
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	200MG	A078310	004	Feb 04, 2009
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LAMOTRIGINE

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

RUBICON	25MG	A202887 001	Jun 17, 2013
	50MG	A202887 002	Jun 17, 2013
TORRENT	200MG	A203370 004	Dec 23, 2013

TABLET, FOR SUSPENSION;ORAL

LAMICTAL CD

GLAXOSMITHKLINE LLC	100MG	N020764 003	Aug 24, 1998
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LAMOTRIGINE

JUBILANT GENERICS	5MG	A200220 001	Feb 28, 2011
	25MG	A200220 002	Feb 28, 2011
MYLAN	5MG	A076630 001	Jan 22, 2009
	25MG	A076630 002	Jan 22, 2009
SANDOZ	5MG	A078409 002	Jan 22, 2009
	25MG	A078409 003	Jan 22, 2009
TEVA	5MG	A076420 001	Jun 21, 2006
	25MG	A076420 002	Jun 21, 2006

TABLET, ORALLY DISINTEGRATING;ORAL

LAMOTRIGINE

IMPAX LABS INC	25MG	A200828 001	Jul 15, 2013
	50MG	A200828 002	Jul 15, 2013
	100MG	A200828 003	Jul 15, 2013
	200MG	A200828 004	Jul 15, 2013

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

AJANTA PHARMA LTD	15MG	A203957 001	Oct 14, 2016
	30MG	A203957 002	Oct 14, 2016
BRECKENRIDGE	15MG	A203964 001	Oct 17, 2018
	30MG	A203964 002	Oct 17, 2018
KRKA TOVARNA ZDRAVIL	15MG	A091212 001	Sep 16, 2013
	30MG	A091212 002	Sep 16, 2013
LANNETT CO INC	15MG	A207157 001	Sep 29, 2017
MYLAN	15MG	A203187 001	Jun 01, 2016

PREVACID

+ TAKEDA PHARMS USA	15MG	N020406 001	May 10, 1995
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FOR SUSPENSION, DELAYED RELEASE;ORAL

PREVACID

TAKEDA PHARMS NA	15MG/PACKET	N021281 001	May 03, 2001
	30MG/PACKET	N021281 002	May 03, 2001

INJECTABLE;INTRAVENOUS

PREVACID IV

+ TAKEDA PHARMS NA	30MG/VIAL **	N021566 001	May 27, 2004
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TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

ANI PHARMS	15MG	A078730 001	Oct 15, 2010
	30MG	A078730 002	Oct 15, 2010

LANSOPRAZOLE; NAPROXEN

CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

PREVACID NAPRAPAC 250 (COPACKAGED)

+ TAKEDA PHARMS NA	15MG,N/A;N/A,250MG **	N021507 002	Nov 14, 2003
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PREVACID NAPRAPAC 375 (COPACKAGED)

TAKEDA PHARMS NA	15MG,N/A;N/A,375MG	N021507 003	Nov 14, 2003
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PREVACID NAPRAPAC 500 (COPACKAGED)

TAKEDA PHARMS NA	15MG,N/A;N/A,500MG	N021507 004	Nov 14, 2003
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LANTHANUM CARBONATE

TABLET, CHEWABLE;ORAL

FOSRENOL

TAKEDA PHARMS USA	EQ 250MG BASE	N021468 001	Oct 26, 2004
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LAPYRIUM CHLORIDE; UNDECOYLUM CHLORIDE IODINE COMPLEX

SOLUTION;TOPICAL

VIRAC REX

CHESEBROUGH PONDS	0.5%;1.8%	N011914 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LASMIDITAN SUCCINATE

TABLET; ORAL

REYVOW

+ ELI LILLY AND CO EQ 200MG BASE N211280 003 Dec 18, 2020

LATANOPROST

SOLUTION/DROPS; OPHTHALMIC

LATANOPROST

APOTEX INC 0.005% A077697 001 Mar 22, 2011

EPIC PHARMA LLC 0.005% A090887 001 Jul 19, 2011

EUGIA PHARMA 0.005% A206519 001 Sep 03, 2019

LEFAMULIN ACETATE

SOLUTION; INTRAVENOUS

XENLETA

+ NABRIVA EQ 150MG BASE/15ML (EQ 10MG BASE/ML) N211673 001 Aug 19, 2019

LEFLUNOMIDE

TABLET; ORAL

LEFLUNOMIDE

BARR 10MG A077083 001 Sep 13, 2005

20MG A077083 002 Sep 13, 2005

SANDOZ 10MG A077085 001 Sep 13, 2005

20MG A077085 002 Sep 13, 2005

TEVA PHARMS 10MG A077084 001 Sep 13, 2005

20MG A077084 002 Sep 13, 2005

LENALIDOMIDE

CAPSULE; ORAL

LENALIDOMIDE

TORRENT 10MG A213405 001 Feb 17, 2023

20MG A213405 003 Aug 03, 2023

25MG A213405 002 Feb 17, 2023

LESINURAD

TABLET; ORAL

ZURAMPIC

+ IRONWOOD PHARMS INC 200MG N207988 001 Dec 22, 2015

LETROZOLE

TABLET; ORAL

LETROZOLE

ACTAVIS TOTOWA 2.5MG A090292 001 Jul 13, 2011

APOTEX INC 2.5MG A091303 001 Apr 19, 2012

CHARTWELL RX 2.5MG A202716 001 May 16, 2013

FRESENIUS KABI USA 2.5MG A090491 001 Jun 03, 2011

HIKMA 2.5MG A090838 001 Jun 03, 2011

HIKMA PHARMS 2.5MG A203796 001 Jun 03, 2016

IMPAX LABS 2.5MG A091638 001 Jun 03, 2011

INDICUS PHARMA 2.5MG A201804 001 Jun 03, 2011

LANNETT CO INC 2.5MG A091098 001 Jun 03, 2011

2.5MG A202048 001 Oct 29, 2014

MYLAN 2.5MG A078190 001 Dec 24, 2008

STRIDES PHARMA 2.5MG A090789 001 Jun 03, 2011

SUN PHARM INDS LTD 2.5MG A091466 001 Jun 03, 2011

SYNTHON PHARMS 2.5MG A090196 001 Jun 03, 2011

LEUCOVORIN CALCIUM

FOR SOLUTION; ORAL

LEUCOVORIN CALCIUM

HOSPIRA EQ 60MG BASE/VIAL N008107 003 Jan 30, 1987

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

ABIC EQ 3MG BASE/ML A089352 001 Jun 01, 1988

EQ 50MG BASE/VIAL A089353 001 Jun 01, 1988

ABRAXIS PHARM EQ 50MG BASE/VIAL A088939 001 Dec 01, 1986

ELKINS SINN EQ 50MG BASE/VIAL A070480 001 Jan 02, 1987

EQ 100MG BASE/VIAL A081224 001 Jun 03, 1994

+ HOSPIRA EQ 3MG BASE/ML ** N008107 001

+ EQ 50MG BASE/VIAL ** N008107 002

+ EQ 100MG BASE/VIAL ** N008107 004 May 23, 1988

+ EQ 350MG BASE/VIAL ** N008107 005 Apr 05, 1989

NOVAST LABS EQ 10MG BASE/ML A210917 001 Nov 23, 2018

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

PHARMACHEMIE	EQ 350MG BASE/VIAL	A040262 001	Dec 15, 1999
PHARMACHEMIE USA	EQ 50MG BASE/VIAL	A089628 001	Apr 17, 1997
	EQ 100MG BASE/VIAL	A089915 001	Apr 17, 1997
TEVA PARENTERAL	EQ 50MG BASE/VIAL	A081278 001	Sep 28, 1993
TEVA PHARMS USA	EQ 100MG BASE/VIAL	A081277 001	Sep 28, 1993
	EQ 350MG BASE/VIAL	A040174 001	Jun 12, 1997
LEUCOVORIN CALCIUM PRESERVATIVE FREE			
AM REGENT	EQ 50MG BASE/VIAL	A040338 001	Jan 31, 2001
HIKMA	EQ 10MG BASE/ML	A040347 001	Apr 25, 2000
HOSPIRA	EQ 10MG BASE/ML **	A040147 001	Jun 25, 1997
TEVA PARENTERAL	EQ 10MG BASE/ML	A040332 001	Jun 28, 1999
WELLCOVORIN			
GLAXOSMITHKLINE	EQ 5MG BASE/ML	A087439 001	Oct 19, 1982
	EQ 25MG BASE/VIAL	A089833 001	Jan 23, 1989
	EQ 50MG BASE/VIAL	A089465 001	Jan 23, 1989
	EQ 100MG BASE/VIAL	A089834 001	Jan 23, 1989

TABLET; ORAL

LEUCOVORIN CALCIUM

ANI PHARMS	EQ 15MG BASE	A075327 001	Mar 24, 1999
PAR PHARM	EQ 5MG BASE	A071600 001	Oct 14, 1987
	EQ 25MG BASE	A071598 001	Oct 14, 1987
PHARMACHEMIE	EQ 5MG BASE	A073099 001	Mar 28, 1997
	EQ 25MG BASE	A073101 001	Mar 28, 1997
XANODYNE PHARM	EQ 5MG BASE	N018459 001	Jan 30, 1986
	EQ 10MG BASE	A071962 001	Nov 19, 1987
	EQ 15MG BASE	A071104 001	Mar 04, 1987

WELLCOVORIN

+	GLAXOSMITHKLINE	EQ 5MG BASE **	N018342 001	Jul 08, 1983
+		EQ 25MG BASE **	N018342 002	Jul 08, 1983

LEUPROLIDE ACETATE

IMPLANT; IMPLANTATION

VIADUR

ORTHO MCNEIL JANSSEN	EQ 65MG BASE	N021088 001	Mar 03, 2000
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INJECTABLE; INJECTION

LEUPROLIDE ACETATE

GENZYME	1MG/0.2ML	A075721 001	Nov 29, 2001	
LUPRON				
+	ABBVIE ENDOCRINE INC	1MG/0.2ML **	N019010 001	Apr 09, 1985
LUPRON DEPOT				
+	ABBVIE ENDOCRINE INC	3.75MG/VIAL **	N020011 001	Oct 22, 1990

POWDER; INTRAMUSCULAR

LUPRON DEPOT-PED KIT

+	ABBVIE ENDOCRINE INC	3.75MG, 7.5MG **	N020263 003	Apr 16, 1993
+		7.5MG, 7.5MG **	N020263 004	Apr 16, 1993

LEUPROLIDE ACETATE; NORETHINDRONE ACETATE

INJECTABLE, TABLET; INTRAMUSCULAR, ORAL

LUPANETA PACK

+	ABBVIE ENDOCRINE	3.75MG/VIAL, N/A; N/A, 5MG	N203696 001	Dec 14, 2012
+		11.25MG/VIAL, N/A; N/A, 5MG	N203696 002	Dec 14, 2012

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

MYLAN SPECIALITY LP	EQ 0.0103% BASE	A077800 001	Mar 15, 2013
	EQ 0.021% BASE	A077800 002	Mar 15, 2013
	EQ 0.042% BASE	A077800 003	Mar 15, 2013

LEVALLORPHAN TARTRATE

INJECTABLE; INJECTION

LORFAN

ROCHE	1MG/ML	N010423 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVAMISOLE HYDROCHLORIDE

TABLET; ORAL

ERGAMISOL

JANSSEN PHARMA EQ 50MG BASE N020035 001 Jun 18, 1990

LEVAMLODIPINE MALEATE

TABLET; ORAL

CONJUPRI

+	CSPC OUYI	EQ 1.25MG BASE	N212895 001	Dec 19, 2019
+		EQ 2.5MG BASE	N212895 002	Dec 19, 2019
+		EQ 5MG BASE	N212895 003	Dec 19, 2019

LEVETIRACETAM

INJECTABLE; INTRAVENOUS

LEVETIRACETAM

AKORN	500MG/5ML (100MG/ML)	A209934 001	May 04, 2018
AM REGENT	500MG/5ML (100MG/ML)	A202143 001	Jan 31, 2012
FRESENIUS KABI USA	500MG/5ML (100MG/ML)	A090813 001	May 26, 2010
JUBILANT GENERICS	500MG/5ML (100MG/ML)	A206838 001	Jun 02, 2016

SOLUTION; ORAL

LEVETIRACETAM

APOTEX INC	100MG/ML	A090187 001	Aug 05, 2011
PHARMOBEDIENT CNSLTG	100MG/ML	A090028 001	Mar 03, 2010
TOLMAR	100MG/ML	A079107 001	Jan 15, 2009

TABLET; ORAL

LEVETIRACETAM

ACTAVIS LABS FL INC	250MG	A077408 001	Mar 02, 2009
	500MG	A077408 002	Mar 02, 2009
	750MG	A077408 003	Mar 02, 2009
FOSUN PHARMA	250MG	A077324 001	Jan 15, 2009
	500MG	A077324 002	Jan 15, 2009
	750MG	A077324 003	Jan 15, 2009
	1GM	A077324 004	Jan 15, 2009
LOTUS PHARM CO LTD	250MG	A090906 002	Oct 31, 2016
	500MG	A090906 001	Nov 05, 2010
	750MG	A090906 003	Oct 31, 2016
	1GM	A090906 004	Oct 31, 2016
MYLAN	250MG	A076919 001	Nov 04, 2008
	250MG	A078731 001	Feb 10, 2009
	500MG	A078731 002	Feb 10, 2009
	750MG	A078731 003	Feb 10, 2009
	1GM	A078731 004	Feb 10, 2009
NOSTRUM LABS INC	250MG	A090511 001	Aug 18, 2011
	500MG	A090511 002	Aug 18, 2011
	750MG	A090511 003	Aug 18, 2011
	1GM	A090511 004	Aug 18, 2011
SECAN PHARMS	500MG	A205102 004	Dec 16, 2015
	1GM	A205102 003	Dec 16, 2015
TEVA PHARMS	250MG	A078101 001	Jan 15, 2009
	500MG	A078101 002	Jan 15, 2009
	750MG	A078101 003	Jan 15, 2009
	1GM	A078101 004	Jan 15, 2009
WATSON LABS INC	250MG	A078797 002	Jan 15, 2009
	500MG	A078797 003	Jan 15, 2009
	750MG	A078797 004	Jan 15, 2009
	1GM	A078797 001	Jan 15, 2009

TABLET, EXTENDED RELEASE; ORAL

ELEPSIA XR

+	TRIPOINT	1GM	N204417 001	Dec 20, 2018
+		1.5GM	N204417 002	Dec 20, 2018

LEVETIRACETAM

ACTAVIS ELIZABETH	500MG	A091557 001	Sep 12, 2011
	750MG	A091557 002	Sep 12, 2011
AUROBINDO PHARMA USA	500MG	A200475 001	Dec 19, 2011
	750MG	A200475 002	Dec 19, 2011
	1GM	A200475 003	Dec 07, 2015
CHARTWELL RX	500MG	A091291 001	Sep 12, 2011
	750MG	A091291 002	Sep 12, 2011
DEXCEL	500MG	A202167 001	Sep 04, 2015
	750MG	A202167 002	Sep 04, 2015

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVETIRACETAM

TABLET, EXTENDED RELEASE;ORAL

LEVETIRACETAM

LOTUS PHARM CO LTD	500MG	A202095 002	Jun 06, 2016
	750MG	A202095 001	Jun 06, 2016
ROUSES POINT PHARMS	500MG	A202524 001	Aug 27, 2012
	750MG	A202524 002	Aug 27, 2012
SANDOZ	500MG	A091668 001	Nov 01, 2012
	750MG	A091668 002	Nov 01, 2012
SUN PHARM	500MG	A203059 001	Sep 09, 2013
	750MG	A203059 002	Sep 09, 2013
SUN PHARM INDUSTRIES	500MG	A091285 001	Sep 12, 2011
	750MG	A091285 002	Sep 12, 2011
TEVA PHARMS	500MG	A091430 001	Sep 12, 2011
	750MG	A091430 002	Sep 12, 2011
TORRENT PHARMS LTD	500MG	A091338 001	May 29, 2012
	750MG	A091338 002	May 29, 2012

LEVOBETAXOLOL HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

BETAXON

ALCON PHARMS LTD	EQ 0.5% BASE	N021114 001	Feb 23, 2000
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LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AKBETA

AKORN	0.5%	A074780 001	Oct 29, 1996
EPIC PHARMA LLC	0.25%	A074779 001	Oct 29, 1996

BETAGAN

+ ALLERGAN

0.25% **

N019814 001 Jun 28, 1989

LEVOBUNOLOL HYDROCHLORIDE

ALCON LABS INC	0.25%	A074851 001	Oct 28, 1996
APOTEX INC	0.25%	A075473 001	Aug 03, 2000
	0.5%	A075475 001	Aug 03, 2000
BAUSCH AND LOMB	0.25%	A074307 001	Mar 04, 1994
SANDOZ	0.5%	A074850 001	Oct 28, 1996

LEVOBUPIVACAINE HYDROCHLORIDE

INJECTABLE;INJECTION

CHIROCAINE

PURDUE PHARMA LP	EQ 2.5MG BASE/ML **	N020997 001	Aug 05, 1999
	EQ 5MG BASE/ML **	N020997 002	Aug 05, 1999
	EQ 7.5MG BASE/ML **	N020997 003	Aug 05, 1999

LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

LIVOSTIN

NOVARTIS	EQ 0.05% BASE	N020219 001	Nov 10, 1993
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LEVOCARNITINE

INJECTABLE;INJECTION

LEVOCARNITINE

TEVA PHARMS USA	200MG/ML	A075881 001	Mar 29, 2001
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SOLUTION;ORAL

CARNITOR

LEADIANT BIOSCI INC	1GM/10ML	N018948 002	Apr 27, 1988
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LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX	2.5MG/5ML	A202915 001	Aug 21, 2014
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XYZAL

+ CHATTEM SANOFI

2.5MG/5ML **

N022157 001 Jan 28, 2008

TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

APOTEX	5MG	A203027 001	Feb 13, 2015
GRANULES	5MG	A090486 001	Mar 26, 2013
SUN PHARM INDS LTD	5MG	A201653 001	Jun 26, 2015
US ANTIBIOTICS	5MG	A204323 001	Dec 20, 2016

XYZAL

+ CHATTEM SANOFI

5MG **

N022064 001 May 25, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVODOPA

CAPSULE; ORAL

BENDOPA

VALEANT PHARM INTL	100MG	N016948 003
	250MG	N016948 001
	500MG	N016948 002

DOPAR

SHIRE	100MG	N016913 003
	250MG	N016913 001
	500MG	N016913 002

LARODOPA

ROCHE	100MG	N016912 002
	250MG	N016912 001
	500MG	N016912 006

TABLET; ORAL

DOPAR

SHIRE	250MG	N016913 004
	500MG	N016913 005

LARODOPA

ROCHE	100MG	N016912 005
	250MG	N016912 003
	500MG	N016912 004

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVAQUIN

+	JANSSEN PHARMS	EQ 500MG/20ML (EQ 25MG/ML) **	N020635 001	Dec 20, 1996
+		EQ 750MG/30ML (EQ 25MG/ML) **	N020635 004	Dec 20, 1996

LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER

+	JANSSEN PHARMS	EQ 250MG/50ML (EQ 5MG/ML) **	N020635 002	Dec 20, 1996
+		EQ 500MG/100ML (EQ 5MG/ML) **	N020635 003	Dec 20, 1996
+		EQ 750MG/150ML (EQ 5MG/ML) **	N020635 005	Dec 20, 1996

LEVOFLOXACIN

AVET LIFESCIENCES	EQ 500MG/20ML (EQ 25MG/ML)	A202590 001	Jan 24, 2013
	EQ 750MG/30ML (EQ 25MG/ML)	A202590 002	Jan 24, 2013
EUGIA PHARMA	EQ 500MG/20ML (EQ 25MG/ML)	A202328 001	Jan 24, 2013
	EQ 750MG/30ML (EQ 25MG/ML)	A202328 002	Jan 24, 2013
HOSPIRA	EQ 500MG/20ML (EQ 25MG/ML)	A078577 001	Aug 12, 2015
	EQ 750MG/30ML (EQ 25MG/ML)	A078577 002	Aug 12, 2015
MYLAN ASI	EQ 500MG/20ML (EQ 25MG/ML)	A200560 001	Jun 20, 2011
	EQ 750MG/30ML (EQ 25MG/ML)	A200560 002	Jun 20, 2011
ZYDUS PHARMS	EQ 500MG/20ML (EQ 25MG/ML)	A205968 001	Jun 01, 2017
	EQ 750MG/30ML (EQ 25MG/ML)	A205968 002	Jun 01, 2017

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

EUGIA PHARMA	EQ 250MG/50ML (EQ 5MG/ML)	A206919 001	Feb 10, 2016
	EQ 500MG/100ML (EQ 5MG/ML)	A206919 002	Feb 10, 2016
	EQ 750MG/150ML (EQ 5MG/ML)	A206919 003	Feb 10, 2016

SOLUTION; ORAL

LEVAQUIN

+	JANSSEN PHARMS	250MG/10ML	N021721 001	Oct 21, 2004
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SOLUTION/DROPS; OPHTHALMIC

IQUIX

+	SANTEN	1.5% **	N021571 001	Mar 01, 2004
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LEVOFLOXACIN

AKORN	0.5%	A090268 001	Dec 20, 2010
MYLAN LABS LTD	0.5%	A204899 001	Dec 08, 2017
RUBICON	0.5%	A078282 001	Dec 20, 2010
WATSON LABS TEVA	0.5%	A076826 001	Feb 10, 2011

QUIXIN

+	SANTEN	0.5% **	N021199 001	Aug 18, 2000
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TABLET; ORAL

LEVAQUIN

+	JANSSEN PHARMS	250MG **	N020634 001	Dec 20, 1996
+		500MG **	N020634 002	Dec 20, 1996
+		750MG **	N020634 003	Sep 08, 2000

LEVOFLOXACIN

JUBILANT GENERICS	250MG	A203613 001	Jun 19, 2015
	500MG	A203613 002	Jun 19, 2015
MYLAN	250MG	A076276 001	Jun 20, 2011
	500MG	A076276 002	Jun 20, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOFLOXACIN

TABLET; ORAL

LEVOFLOXACIN

	750MG	A077097 001	Jun 20, 2011
TORRENT PHARMS	250MG	A090722 001	Jun 20, 2011
	500MG	A090722 002	Jun 20, 2011
	750MG	A090722 003	Jun 20, 2011
WATSON LABS INC	250MG	A201484 001	Nov 22, 2013
	500MG	A201484 002	Nov 22, 2013
	750MG	A201484 003	Nov 22, 2013

LEVOLEUCOVORIN

POWDER; INTRAVENOUS

KHAPZORY

+ ACROTECH BIOPHARMA	300MG/VIAL	N211226 002	Oct 19, 2018
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LEVOLEUCOVORIN CALCIUM

POWDER; INTRAVENOUS

LEVOLEUCOVORIN CALCIUM

ACTAVIS LLC	EQ 50MG BASE/VIAL	A206516 001	Feb 13, 2017
+	EQ 175MG BASE/VIAL	N208723 001	Sep 29, 2016
AMNEAL	EQ 50MG BASE/VIAL	A207547 001	Feb 13, 2017
HIKMA	EQ 50MG BASE/VIAL	A206263 001	Jun 16, 2016

SOLUTION; INTRAVENOUS

FUSILEV

+ ACROTECH BIOPHARMA	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	N020140 002	Apr 29, 2011
	**		
+	EQ 250MG BASE/25ML (EQ 10MG BASE/ML) **	N020140 003	Apr 29, 2011

LEVOLEUCOVORIN CALCIUM

AILEX PHARMS LLC	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A213797 001	Nov 02, 2021
MEITHEAL	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A211002 002	Aug 16, 2019
MYLAN TEORANTA	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A203576 001	Oct 20, 2015
	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A203576 002	Oct 20, 2015
NOVAST LABS	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	A210623 001	May 03, 2018
	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A210623 002	May 03, 2018
SANDOZ	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	A203563 002	Mar 09, 2015

LEVOMEPRMAZINE

INJECTABLE; INJECTION

LEVOPROME

IMMUNEX	20MG/ML	N015865 001	
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LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE; ORAL

ORLAAM

+ ROXANE	10MG/ML **	N020315 001	Jul 09, 1993
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LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

LEVOMILNACIPRAN HYDROCHLORIDE

AMNEAL PHARMS CO	EQ 20MG BASE	A210790 001	Feb 04, 2019
	EQ 40MG BASE	A210790 002	Feb 04, 2019
	EQ 80MG BASE	A210790 003	Feb 04, 2019
	EQ 120MG BASE	A210790 004	Feb 04, 2019
AUROBINDO PHARMA LTD	EQ 20MG BASE	A210826 001	Jan 06, 2023
	EQ 40MG BASE	A210826 002	Jan 06, 2023
	EQ 80MG BASE	A210826 003	Jan 06, 2023
	EQ 120MG BASE	A210826 004	Jan 06, 2023
HIKMA	EQ 20MG BASE	A210732 001	Nov 05, 2020
	EQ 40MG BASE	A210732 002	Nov 05, 2020
	EQ 80MG BASE	A210732 003	Nov 05, 2020
	EQ 120MG BASE	A210732 004	Nov 05, 2020

LEVONORDEFRIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE W/ LEVONORDEFRIN

SOLVAY	0.05MG/ML; 2%	A085010 001	
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CARBOCAINE W/ NEO-COBEFRIN

EASTMAN KODAK	0.05MG/ML; 2%	N012125 002	
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ISOCAINE HYDROCHLORIDE W/ LEVONORDEFRIN

SEPTODONT	0.05MG/ML; 2%	A084697 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVONORDEFRIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFRIN

BELMORA LLC	0.05MG/ML;2%	A084850	002	Oct 21, 1983
POLOCAINE W/ LEVONORDEFRIN				
DENTSPLY PHARM	0.05MG/ML;2%	A089517	001	Apr 14, 1988
SCANDONEST L				
DEPROCO	0.05MG/ML;2%	A088388	001	Oct 10, 1984

LEVONORDEFRIN; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAINE AND NOVOCAIN W/ NEO-COBEFRIN

EASTMAN KODAK	0.05MG/ML;2%;0.4%	N008592	007	
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LEVONORGESTREL

IMPLANT; IMPLANTATION

JADELLE

+ POPULATION COUNCIL	75MG/IMPLANT **	N020544	001	Nov 01, 1996
LEVONORGESTREL				
WYETH PHARMS INC	75MG/IMPLANT	N020627	001	Aug 15, 1996
NORPLANT				
POPULATION COUNCIL	36MG/IMPLANT	N019897	001	Dec 10, 1990
NORPLANT SYSTEM IN PLASTIC CONTAINER				
WYETH PHARMS INC	36MG/IMPLANT	N020088	001	Dec 10, 1990

TABLET; ORAL

LEVONORGESTREL

ALVOGEN	1.5MG	A202246	001	Jun 05, 2015
FDN CONSUMER	0.75MG **	A078665	001	Aug 28, 2009
	1.5MG	A200670	001	Jul 12, 2012
L PERRIGO CO	0.75MG	A090740	001	Dec 30, 2010
LOTUS PHARM CO LTD	0.75MG	A202684	001	Sep 02, 2016
LUPIN LTD	0.75MG	A091328	001	Jan 23, 2013
MYLAN LABS LTD	0.75MG	A202740	001	Sep 02, 2016
NAARI PTE LTD	1.5MG	A207660	001	May 02, 2019
WATSON LABS	0.75MG	A078666	001	Jun 24, 2009
PLAN B				
+ FDN CONSUMER	0.75MG **	N021045	001	Jul 28, 1999
+	0.75MG **	N021045	002	Aug 24, 2006

LEVOPROPOXYPHENE NAPSYLATE ANHYDROUS

CAPSULE; ORAL

NOVRAD

LILLY	EQ 50MG BASE	N012928	006	
	EQ 100MG BASE	N012928	004	

SUSPENSION; ORAL

NOVRAD

LILLY	EQ 50MG BASE/5ML	N012928	002	
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LEVORPHANOL TARTRATE

INJECTABLE; INJECTION

LEVO-DROMORAN

VALEANT PHARM INTL	2MG/ML	N008719	001	Dec 19, 1991
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TABLET; ORAL

LEVO-DROMORAN

+ VALEANT PHARM INTL	2MG **	N008720	001	Dec 19, 1991
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LEVORPHANOL TARTRATE

HIKMA	1MG	A074278	002	Jun 18, 2018
	3MG	A074278	003	Jun 18, 2018

LEVOTHYROXINE SODIUM

CAPSULE; ORAL

LEVOTHYROXINE SODIUM

TEVA PHARMS USA INC	0.075MG	A211369	001	Oct 28, 2020
	0.088MG	A213256	001	Jan 06, 2021
	0.1MG	A213256	002	Jan 06, 2021
	0.112MG	A211369	003	Apr 16, 2021
	0.125MG	A213256	003	Jan 06, 2021
	0.137MG	A211369	005	May 02, 2023
	0.15MG	A211369	002	Oct 28, 2020
	0.175MG	A211369	006	May 02, 2023
	0.2MG	A211369	004	Nov 09, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOTHYROXINE SODIUM

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

DR REDDYS 100MCG/VIAL A208837 001 Mar 27, 2020

PAR STERILE PRODUCTS 200MCG/VIAL A205366 001 Dec 07, 2015

TABLET; ORAL

EUTHYROX

PROVELL 0.3MG N021292 012 May 31, 2002

LEVOTHYROXINE SODIUM

AMNEAL 0.025MG A210831 001 Feb 19, 2019

0.05MG A210831 002 Feb 19, 2019

0.075MG A210831 003 Feb 19, 2019

0.088MG A210831 004 Feb 19, 2019

0.1MG A210831 005 Feb 19, 2019

0.112MG A210831 006 Feb 19, 2019

0.125MG A210831 007 Feb 19, 2019

0.137MG A210831 008 Feb 19, 2019

0.15MG A210831 009 Feb 19, 2019

0.175MG A210831 010 Feb 19, 2019

0.2MG A210831 011 Feb 19, 2019

0.3MG A210831 012 Feb 19, 2019

MERCK KGAA 0.025MG A076752 001 Jun 16, 2005

0.05MG A076752 002 Jun 16, 2005

0.075MG A076752 003 Jun 16, 2005

0.088MG A076752 004 Jun 16, 2005

0.1MG A076752 005 Jun 16, 2005

0.112MG A076752 006 Jun 16, 2005

0.125MG A076752 007 Jun 16, 2005

0.15MG A076752 008 Jun 16, 2005

0.175MG A076752 009 Jun 16, 2005

0.2MG A076752 010 Jun 16, 2005

0.3MG A076752 011 Jun 16, 2005

LEVOXYL

+ KING PHARMS 0.3MG ** N021301 012 May 25, 2001

LIDOCAINE

AEROSOL; ORAL

XYLOCAINE

ASTRAZENECA 10% N014394 001

FILM, EXTENDED RELEASE; BUCCAL

DENTIPATCH

NOVEN 23MG/PATCH N020575 001 May 21, 1996

OINTMENT; TOPICAL

ALPHACAINE

CARLISLE 5% A084944 001

5% A084946 001

5% A084947 001

LIDOCAINE

AILEX PHARMS LLC 5% A212486 001 Oct 17, 2019

BELMORA LLC 5% A080210 001

RISING 5% A208604 001 Sep 20, 2017

TEVA PHARMS USA 5% A210256 001 Jan 16, 2018

VERTICE 5% A205318 001 Feb 01, 2016

VITRUVIAS THERAP 5% A208822 001 Sep 25, 2017

XYLOCAINE

+ ASTRAZENECA 5% ** N008048 001

PATCH; TOPICAL

DENTIPATCH

NOVEN 46.1MG/PATCH N020575 002 May 21, 1996

LIDOCAINE

NOVEN PHARMS INC 5% A203265 001 Dec 01, 2020

SOLUTION; TOPICAL

XYLOCAINE

ASTRAZENECA 5% N014127 001

SUPPOSITORY; RECTAL

XYLOCAINE

ASTRAZENECA 100MG N013077 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ALPHACAINE HYDROCHLORIDE

CARLISLE 2% A084721 001

LIDOCAINE HYDROCHLORIDE

ABBOTT 10% A087980 001 Feb 02, 1983

20% A089362 001 May 25, 1988

ABRAXIS PHARM 1% A080420 001

1% A086761 001

1.5% A080420 005

2% A080420 002

2% A080420 004

2% A086761 002

2% N017508 001

4% N017508 002

20% N017508 004

AKORN 1% A085037 001

2% A085037 002

AM REGENT 1% A080850 001

1% A091564 001 Aug 14, 2015

BEL MAR 1% A080710 001

2% A080760 001

BELMORA LLC 2% A080504 001

DELL LABS 1% A083387 001

2% A083388 001

ELKINS SINN 0.5% A085131 001

4% A084626 001

GD SEARLE LLC 1% A083135 001

2% A083135 002

HOSPIRA 1% A040013 001 Jun 23, 1995

1.5% A088330 001 May 17, 1984

2% A088331 001 May 17, 1984

20% A083158 003

INTL MEDICATION 1% N017701 002

2% N017701 001

1GM/VIAL N018543 001

2GM/VIAL N018543 002

LUITPOLD 2% A083198 001

LYPHOMED 1% A080390 001

2% A080390 002

MILES 1% A080414 001

2% A080414 002

RISING 0.5% A091056 001 Dec 08, 2010

0.5% A091058 001 Sep 30, 2010

1% A091056 002 Dec 08, 2010

1% A091058 002 Sep 30, 2010

2% A202242 001 Apr 11, 2014

WATSON LABS 1% A080377 001

1% A083627 001

2% A080377 002

2% A083627 002

WYETH AYERST 1% A083083 001

2% A083083 002

LIDOCAINE HYDROCHLORIDE 0.1% AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 100MG/100ML N018461 001

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 200MG/100ML N018967 001 Mar 30, 1984

LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5%

HOSPIRA 200MG/100ML A083158 005

LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT 200MG/100ML N018954 001 Jul 09, 1985

HOSPIRA 200MG/100ML N018388 001

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 400MG/100ML N018967 002 Mar 30, 1984

LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5%

HOSPIRA 400MG/100ML A083158 006

LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA 400MG/100ML N018388 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	800MG/100ML	N018967	003	Mar 30, 1984
LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER				
HOSPIRA	800MG/100ML	N018388	003	Nov 05, 1982
LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER				
HOSPIRA	1.5%	A088326	001	Jul 31, 1984
	10%	A088367	001	Jul 31, 1984
	20%	A088368	001	Jul 31, 1984
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE				
INTL MEDICATION	4%	N017702	002	
	20%	N017702	001	
MYLAN LABS LTD	2%	A090665	001	Sep 27, 2010
LIDOCATON				
PHARMATON	2%	A084727	001	Aug 17, 1983
LIDOPEN				
MERIDIAN MEDCL TECHN	10%	N017549	001	
XYLOCAINE				
ASTRAZENECA	1%	N010418	005	
	1.5%	N010418	009	
	2%	N010418	007	
XYLOCAINE 4% PRESERVATIVE FREE				
+ FRESENIUS KABI USA	4%	N010417	001	
XYLOCAINE DENTAL				
DENTSPLY PHARM	2%	N021380	001	
XYLOCAINE PRESERVATIVE FREE				
+ FRESENIUS KABI USA	1% **	N016801	005	Jan 19, 1988
+	2% **	N016801	001	
+	4% **	N016801	002	
	10% **	N016801	003	
+	20% **	N016801	004	
INJECTABLE; SPINAL				
XYLOCAINE 1.5% W/ DEXTROSE 7.5%				
FRESENIUS KABI USA	1.5%	N016297	001	
XYLOCAINE 5% W/ GLUCOSE 7.5%				
ASTRAZENECA	5%	N010496	002	Jul 07, 1982
JELLY; TOPICAL				
ANESTACON				
BIONPHARMA	2%	A080429	001	
LIDOCAINE HYDROCHLORIDE				
COSETTE	2%	A081318	001	Apr 29, 1993
WATSON LABS INC	2%	A040837	001	Mar 23, 2011
XYLOCAINE				
+ EPIC PHARMA LLC	2% **	N008816	001	
SOLUTION; ORAL				
LIDOCAINE HYDROCHLORIDE				
HIKMA	2%	A040014	001	Jul 10, 1995
LIDOCAINE HYDROCHLORIDE VISCOUS				
ACTAVIS MID ATLANTIC	2%	A086578	001	
INTL MEDICATION	2%	A086389	001	Feb 02, 1982
XYLOCAINE VISCOUS				
+ FRESENIUS KABI USA	2% **	N009470	001	
SOLUTION; TOPICAL				
LARYNGOTRACHEAL ANESTHESIA KIT				
KENDALL IL	4%	A087931	001	Jun 10, 1983
LIDOCAINE HYDROCHLORIDE				
PACO	4%	A089688	001	Jun 30, 1989
LTA II KIT				
HOSPIRA	4%	A080409	001	
	4%	A088542	001	Jul 31, 1984
PEDIATRIC LTA KIT				
ABBOTT	2%	A088572	001	Jul 31, 1984
HOSPIRA	2%	A085995	001	
XYLOCAINE 4% PRESERVATIVE FREE				
+ FRESENIUS KABI USA	4% **	N010417	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE; OXYTETRACYCLINE

INJECTABLE; INJECTION

TERRAMYCIN

PFIZER

2%; 50MG/ML
2%; 125MG/MLA060567 001
A060567 002LIDOCAINE; PRILOCAINE

CREAM; TOPICAL

EMLA

+ TEVA BRANDED PHARM 2.5%; 2.5%
LIDOCAINE AND PRILOCAINE
PAI HOLDINGS PHARM 2.5%; 2.5%
RHODES PHARMS 2.5%; 2.5%N019941 001 Dec 30, 1992
A205887 001 Jun 29, 2018
A213253 001 Sep 21, 2020

DISC; TOPICAL

EMLA

ASTRAZENECA 2.5%; 2.5%

N020962 001 Feb 04, 1998

LIDOCAINE; TETRACAINE

CREAM; TOPICAL

PLIAGLIS

+ TARO 7%; 7%

N021717 001 Jun 29, 2006

PATCH; TOPICAL

SYNERA

+ GALEN SPECIALTY 70MG; 70MG

N021623 001 Jun 23, 2005

LINACLOTIDE

CAPSULE; ORAL

LINACLOTIDE

AUROBINDO PHARMA 145MCG
290MCG
MYLAN 145MCG
290MCGA209611 001 Feb 07, 2023
A209611 002 Feb 07, 2023
A209564 001 Feb 09, 2021
A209564 002 Feb 09, 2021LINAGLIPTIN

TABLET; ORAL

LINAGLIPTIN

ZYDUS PHARMS 5MG

A208448 001 Mar 30, 2023

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

LINAGLIPTIN AND METFORMIN HYDROCHLORIDE

ZYDUS PHARMS 2.5MG; 500MG
2.5MG; 850MG
2.5MG; 1GMA208449 001 Mar 30, 2023
A208449 002 Mar 30, 2023
A208449 003 Mar 30, 2023LINCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

LINCOCIN

PHARMACIA AND UPJOHN EQ 250MG BASE
EQ 500MG BASEN050316 001
N050316 002

INJECTABLE; INJECTION

LINCOMYCIN HYDROCHLORIDE

GENEYORK PHARMS EQ 300MG BASE/ML
SLATE RUN PHARMA EQ 300MG BASE/ML
WATSON LABS EQ 300MG BASE/MLA212770 001 Mar 12, 2021
A216662 001 Dec 21, 2022
A063180 001 Apr 16, 1991LINDANE

CREAM; TOPICAL

KWELL

REED AND CARNRICK 1%
1%A084218 001
N006309 001

LOTION; TOPICAL

GAMENE

SOLA BARNES HIND 1%

A084989 001

KWELL

REED AND CARNRICK 1%
1%A084218 002
N006309 003

LINDANE

OLTA PHARMS 1%
WOCKHARDT BIO AG 1%A087313 001
A088190 001 Aug 16, 1984

SCABENE

STIEFEL 1%

A086769 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LINDANESHAMPOO; TOPICAL
GAMENE

SOLA BARNES HIND 1%

A084988 001

KWELL

REED AND CARNRICK 1%

A084219 001

1%

N010718 001

LINDANE

OLTA PHARMS 1%

A087266 001

WOCKHARDT BIO AG 1%

A088191 001 Sep 18, 1984

SCABENE

STIEFEL 1%

A087940 001 Apr 08, 1983

LINEZOLID

SOLUTION; INTRAVENOUS

LINEZOLID

HOSPIRA 600MG/300ML (2MG/ML)

A205442 001 Jul 07, 2015

TEVA PHARMS 600MG/300ML (2MG/ML)

A200222 001 Jun 27, 2012

ZYVOX

+ PFIZER 400MG/200ML (2MG/ML) **

N021131 002 Apr 18, 2000

TABLET; ORAL

LINEZOLID

AMNEAL PHARMS 600MG

A204536 001 Dec 21, 2015

GATE PHARMS 600MG

A091210 001 Feb 05, 2016

TEVA PHARMS USA 600MG

A078061 001 May 18, 2015

ZYVOX

+ PFIZER 400MG **

N021130 001 Apr 18, 2000

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

TRIOSTAT

+ PAR STERILE PRODUCTS EQ 0.01MG BASE/ML

N020105 001 Dec 31, 1991

TABLET; ORAL

LIOTHYRONINE SODIUM

MYLAN EQ 0.005MG BASE

A090326 001 Jul 14, 2009

EQ 0.025MG BASE

A090326 002 Jul 14, 2009

EQ 0.05MG BASE

A090326 003 Jul 14, 2009

WATSON LABS EQ 0.025MG BASE

A085755 001 Jan 25, 1982

EQ 0.05MG BASE

A085753 001 Feb 03, 1982

LIOTRIX (T4;T3)

TABLET; ORAL

EUTHROID-0.5

PARKE DAVIS 0.03MG;0.0075MG

N016680 001

EUTHROID-1

PARKE DAVIS 0.06MG;0.015MG

N016680 002

EUTHROID-2

PARKE DAVIS 0.12MG;0.03MG

N016680 003

EUTHROID-3

PARKE DAVIS 0.18MG;0.045MG

N016680 004

THYROLAR-0.25

+ ALLERGAN 0.0125MG;0.0031MG

N016807 001

THYROLAR-0.5

+ ALLERGAN 0.025MG;0.0063MG

N016807 005

THYROLAR-1

+ ALLERGAN 0.05MG;0.0125MG

N016807 004

THYROLAR-2

+ ALLERGAN 0.1MG;0.025MG

N016807 002

THYROLAR-3

+ ALLERGAN 0.15MG;0.0375MG

N016807 003

THYROLAR-5

ALLERGAN 0.25MG;0.0625MG

N016807 006

LISINOPRIL

TABLET; ORAL

LISINOPRIL

HERITAGE PHARMA 2.5MG

A075752 001 Jul 01, 2002

5MG

A075752 002 Jul 01, 2002

10MG

A075752 003 Jul 01, 2002

20MG

A075752 004 Jul 01, 2002

30MG

A075752 005 Jul 01, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LISINAPRILTABLET;ORAL
LISINAPRIL

	40MG	A075752 006	Jul 01, 2002
HIKMA INTL PHARMS	2.5MG	A076063 001	Jul 01, 2002
	5MG	A076063 002	Jul 01, 2002
	10MG	A076063 003	Jul 01, 2002
	20MG	A076063 004	Jul 01, 2002
	30MG	A076063 006	Jun 27, 2003
	40MG	A076063 005	Jul 01, 2002
RISING	2.5MG	A075999 001	Jul 01, 2002
	5MG	A075999 002	Jul 01, 2002
	10MG	A075999 003	Jul 01, 2002
	20MG	A075999 004	Jul 01, 2002
	30MG	A075999 005	Jul 01, 2002
	40MG	A075999 006	Jul 01, 2002
STRIDES PHARMA	2.5MG	A076071 001	Jul 01, 2002
	5MG	A076071 002	Jul 01, 2002
	10MG	A076071 003	Jul 01, 2002
	20MG	A076071 004	Jul 01, 2002
	30MG	A076071 005	Jul 01, 2002
	40MG	A076071 006	Jul 01, 2002
TEVA	2.5MG	A075783 001	Jul 01, 2002
	5MG	A075783 002	Jul 01, 2002
	10MG	A075783 003	Jul 01, 2002
	20MG	A075783 004	Jul 01, 2002
	30MG	A075783 005	Jul 01, 2002
	40MG	A075783 006	Jul 01, 2002
PRINIVIL			
MERCK	2.5MG	N019558 006	Jan 28, 1994
	5MG	N019558 001	Dec 29, 1987
	10MG	N019558 002	Dec 29, 1987
	20MG	N019558 003	Dec 29, 1987
	40MG	N019558 004	Oct 25, 1988

LITHIUM CARBONATE

CAPSULE;ORAL

ESKALITH

NOVEN THERAP

LITHIUM CARBONATE

ABLE

APOTEX INC

MYLAN

USL PHARMA

WATSON LABS

LITHONATE

SOLVAY

TABLET;ORAL

ESKALITH

JDS PHARMS

LITHANE

BAYER PHARMS

LITHIUM CARBONATE

HIKMA INTL PHARMS

PFIZER

LITHOTABS

SOLVAY

TABLET, EXTENDED RELEASE;ORAL

ESKALITH CR

JDS PHARMS

LITHIUM CARBONATE

ABLE

ALEMBIC

HERITAGE PHARMA

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LITHIUM CARBONATE

TABLET, EXTENDED RELEASE;ORAL

LITHIUM CARBONATE

HIKMA INTL PHARMS

450MG

A076490 001 Jun 17, 2003

LITHIUM CITRATE

SYRUP;ORAL

LITHIUM CITRATE

+ HIKMA

EQ 300MG CARBONATE/5ML **

N018421 001

LITHONATE

SOLVAY

EQ 300MG CARBONATE/5ML

N017672 001

LOMEFLOXACIN HYDROCHLORIDE

TABLET;ORAL

MAXAQUIN

PHARMACIA

EQ 400MG BASE

N020013 001 Feb 21, 1992

LOMITAPIDE MESYLATE

CAPSULE;ORAL

JUXTAPID

+ AMRYT

EQ 40MG BASE

N203858 005 Apr 23, 2015

+

EQ 60MG BASE

N203858 006 Apr 23, 2015

LOMUSTINE

CAPSULE;ORAL

GLEOSTINE

+ CORDEN PHARMA

5MG

N017588 004 Dec 19, 2014

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

IMODIUM

J AND J CONSUMER INC

2MG **

N017690 001

+

2MG **

N017694 001

LOPERAMIDE HYDROCHLORIDE

CHARTWELL RX

2MG

A072993 001 Aug 28, 1992

ROXANE

2MG

A073080 001 Nov 27, 1991

TEVA

2MG

A073122 001 Aug 30, 1991

SOLUTION;ORAL

IMODIUM

JANSSEN PHARMS

1MG/5ML

N019037 001 Jul 31, 1984

LOPERAMIDE HYDROCHLORIDE

AKORN

1MG/5ML

A074352 001 Nov 17, 1995

ALLIED

1MG/5ML

A073079 001 Apr 30, 1992

ALPHARMA US PHARMS

1MG/5ML

A073187 001 Sep 15, 1992

DURAMED PHARMS BARR

1MG/5ML

A074991 001 Dec 29, 1997

PERRIGO

1MG/5ML

A073243 001 Jan 21, 1992

TEVA

1MG/5ML

A073478 001 Jun 23, 1995

WATSON LABS

1MG/5ML

A073062 001 May 28, 1993

TABLET;ORAL

LOPERAMIDE HYDROCHLORIDE

ABLE

2MG

A073528 001 Nov 30, 1993

CONTRACT PHARMACAL

2MG

A073254 001 Jul 30, 1993

PERRIGO

2MG

A074194 001 Oct 30, 1992

TABLET, CHEWABLE;ORAL

IMODIUM A-D EZ CHEWS

+ J AND J CONSUMER INC

2MG

N020448 001 Jul 24, 1997

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET;ORAL

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

GRANULES

2MG;125MG

A215981 001 Aug 29, 2022

TABLET, CHEWABLE;ORAL

IMODIUM MULTI-SYMPOM RELIEF

+ J AND J CONSUMER INC

2MG;125MG

N020606 001 Jun 26, 1996

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

PERRIGO

2MG;125MG

A076029 001 Aug 30, 2002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LOPINAVIR; RITONAVIR

CAPSULE; ORAL

KALETRA

ABBVIE	133.3MG;33.3MG	N021226	001	Sep 15, 2000
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LORACARBEF

CAPSULE; ORAL

LORABID

KING PHARMS	200MG	N050668	001	Dec 31, 1991
	400MG	N050668	002	Apr 05, 1996

FOR SUSPENSION; ORAL

LORABID

KING PHARMS	100MG/5ML	N050667	001	Dec 31, 1991
	200MG/5ML	N050667	002	Dec 31, 1991

LORATADINE

CAPSULE; ORAL

LORATADINE

AUROBINDO PHARMA	10MG	A211900	001	Mar 24, 2023
STRIDES PHARMA	10MG	A211926	001	Jan 15, 2020

SYRUP; ORAL

CLARITIN HIVES RELIEF

+ BAYER HEALTHCARE LLC	1MG/ML **	N020641	003	Nov 19, 2003
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LORATADINE

PHARM ASSOC	1MG/ML	A075565	001	Oct 05, 2004
RANBAXY LABS LTD	1MG/ML	A076529	001	Aug 20, 2004
TEVA	1MG/ML	A075505	001	Nov 07, 2003

TABLET; ORAL

LORATADINE

MYLAN	10MG	A075790	001	Nov 07, 2008
	10MG	A078447	001	Aug 12, 2011
FERRIGO	10MG	N021512	001	Jun 24, 2004

TABLET, ORALLY DISINTEGRATING; ORAL

LORATADINE

ACTAVIS LABS FL INC	10MG	A075990	001	Nov 03, 2003
GLAXOSMITHKLINE	10MG	A075822	001	Feb 10, 2003

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

LORATADINE AND PSEUDOEPHEDRINE SULFATE

HERITAGE PHARMA	5MG;120MG	A076208	001	Jan 28, 2004
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LORAZEPAM

CONCENTRATE; ORAL

LORAZEPAM

CHARTWELL MOLECULAR	2MG/ML	A079244	001	Apr 28, 2009
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INJECTABLE; INJECTION

LORAZEPAM

AKORN	2MG/ML	A074974	001	Jul 23, 1998
BEDFORD	2MG/ML	A077076	001	Jul 13, 2005
	4MG/ML	A077076	002	Jul 13, 2005
DAVA PHARMS INC	2MG/ML	A074793	001	Mar 16, 2000
	4MG/ML	A074793	002	Mar 16, 2000
DR REDDYS	1MG/0.5ML	A074551	003	Sep 12, 1996
	2MG/ML	A074535	001	Sep 12, 1996
	2MG/ML	A074551	001	Sep 12, 1996
	4MG/ML	A074535	002	Sep 12, 1996
	4MG/ML	A074551	002	Sep 12, 1996
HIKMA	2MG/ML	A074496	001	Sep 28, 1998
	4MG/ML	A074496	002	Sep 28, 1998
HOSPIRA	2MG/ML	A074280	001	May 27, 1994
	2MG/ML	A074300	001	Apr 12, 1994
	4MG/ML	A074280	002	May 27, 1994
	4MG/ML	A074300	003	Mar 19, 1997
RISING	2MG/ML	A200217	001	Apr 04, 2017
	2MG/ML	A200542	001	Apr 28, 2017
	4MG/ML	A200217	002	Apr 04, 2017
	4MG/ML	A200542	002	Apr 28, 2017
WATSON LABS	2MG/ML	A074276	001	Apr 15, 1994
	4MG/ML	A074276	002	Apr 15, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LORAZEPAM

INJECTABLE; INJECTION

LORAZEPAM PRESERVATIVE FREE

BEDFORD LABS

2MG/ML

A077074 001 Jul 13, 2005

4MG/ML

A077074 002 Jul 13, 2005

SOLUTION; ORAL

LORAZEPAM

ROXANE

0.5MG/5ML

A074648 001 Mar 18, 1997

TABLET; ORAL

LORAZ

QUANTUM PHARMICS

0.5MG

A070200 001 Aug 09, 1985

1MG

A070201 001 Aug 09, 1985

2MG

A070202 001 Aug 09, 1985

LORAZEPAM

AM THERAP

0.5MG

A070727 001 Mar 07, 1986

1MG

A070728 001 Mar 07, 1986

2MG

A070729 001 Mar 07, 1986

AMNEAL PHARMS

0.5MG

A078826 001 Jun 23, 2010

1MG

A078826 002 Jun 23, 2010

2MG

A078826 003 Jun 23, 2010

ANDA REPOSITORY

0.5MG

A072555 002 Mar 29, 1991

1MG

A072555 003 Mar 29, 1991

2MG

A072555 001 Mar 29, 1991

ANI PHARMS

0.5MG

A077396 001 Dec 13, 2006

1MG

A077396 002 Dec 13, 2006

2MG

A077396 003 Dec 13, 2006

CHARTWELL RX

0.5MG

A071591 002 Oct 13, 1987

1MG

A071591 003 Oct 13, 1987

2MG

A071591 001 Oct 13, 1987

HALSEY

0.5MG

A071434 001 Sep 01, 1987

1MG

A071435 001 Sep 01, 1987

2MG

A071436 001 Sep 01, 1987

MUTUAL PHARM

0.5MG

A070472 001 Dec 10, 1985

1MG

A070473 001 Dec 10, 1985

2MG

A070474 001 Dec 10, 1985

PAR PHARM

0.5MG

A070675 001 Dec 01, 1986

1MG

A070676 001 Dec 01, 1986

2MG

A070677 001 Dec 01, 1986

RISING

0.5MG

A077657 001 Mar 16, 2006

1MG

A077657 002 Mar 16, 2006

2MG

A077657 003 Mar 16, 2006

SANDOZ

0.5MG

A071193 001 Apr 15, 1988

1MG

A071194 001 Apr 15, 1988

2MG

A071195 001 Apr 15, 1988

SUPERPHARM

0.5MG

A071245 001 Feb 09, 1987

1MG

A071246 001 Feb 09, 1987

2MG

A071247 001 Feb 09, 1987

USL PHARMA

1MG

A070539 001 Dec 22, 1986

2MG

A070540 001 Dec 22, 1986

WARNER CHILCOTT

1MG

A071038 001 Jan 12, 1988

2MG

A071039 001 Jan 12, 1988

WATSON LABS

0.5MG

A071086 001 Mar 23, 1987

0.5MG

A071117 001 Jul 24, 1986

1MG

A071087 001 Mar 23, 1987

1MG

A071118 001 Jul 24, 1986

2MG

A071088 001 Mar 23, 1987

2MG

A071110 001 Jul 24, 1986

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

APOTEX CORP

25MG

A090790 001 Oct 06, 2010

50MG

A090790 002 Oct 06, 2010

100MG

A090790 003 Oct 06, 2010

HIKMA

25MG

A077459 001 Oct 06, 2010

50MG

A077459 002 Oct 06, 2010

100MG

A077459 003 Oct 06, 2010

HISUN PHARM HANGZHOU

25MG

A204795 001 Apr 04, 2019

50MG

A204795 002 Apr 04, 2019

100MG

A204795 003 Apr 04, 2019

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

MYLAN	25MG	A091590 001	Oct 06, 2010
	50MG	A091590 002	Oct 06, 2010
	100MG	A091590 003	Oct 06, 2010
TEVA	25MG	A076958 001	Apr 06, 2010
	50MG	A076958 002	Apr 06, 2010
	100MG	A076958 003	Apr 06, 2010
TORRENT PHARMS	25MG	A090467 001	Oct 06, 2010
	50MG	A090467 002	Oct 06, 2010
	100MG	A090467 003	Oct 06, 2010
UPSHER SMITH LABS	25MG	A090544 001	Oct 06, 2010
	50MG	A090544 002	Oct 06, 2010
	100MG	A090544 003	Oct 06, 2010

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS; OPHTHALMIC

LOTEMAX

PHARMOS	0.5%	N020841 001	Mar 09, 1998
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LOVASTATIN

TABLET; ORAL

LOVASTATIN

AUROBINDO PHARMA USA	10MG	A075451 001	Dec 17, 2001
	10MG	A075935 001	Dec 17, 2001
	20MG	A075451 002	Dec 17, 2001
	20MG	A075935 002	Dec 17, 2001
	40MG	A075451 003	Dec 17, 2001
	40MG	A075935 003	Dec 17, 2001
SUN PHARM INDUSTRIES	10MG	A077520 001	Apr 14, 2006
	20MG	A077520 002	Apr 14, 2006
	40MG	A077520 003	Apr 14, 2006
MEVACOR			
+ MERCK	10MG **	N019643 002	Mar 28, 1991
+	20MG **	N019643 003	Aug 31, 1987
+	40MG **	N019643 004	Dec 14, 1988

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

COVIS	10MG	N021316 001	Jun 26, 2002
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LOXAPINE HYDROCHLORIDE

CONCENTRATE; ORAL

LOXITANE C

TEVA BRANDED PHARM	EQ 25MG BASE/ML	N017658 001	
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INJECTABLE; INJECTION

LOXITANE IM

ACTAVIS LABS UT INC	EQ 50MG BASE/ML	N018039 001	
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LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

RISING	EQ 5MG BASE	A076762 001	Nov 01, 2004
	EQ 10MG BASE	A076762 002	Nov 01, 2004
	EQ 25MG BASE	A076762 003	Nov 01, 2004
	EQ 50MG BASE	A076762 004	Nov 01, 2004

LOXITANE

+ TEVA BRANDED PHARM	EQ 5MG BASE **	N017525 001	
+	EQ 10MG BASE **	N017525 002	
+	EQ 25MG BASE **	N017525 003	
+	EQ 50MG BASE **	N017525 004	

TABLET; ORAL

LOXITANE

+ TEVA BRANDED PHARM	EQ 10MG BASE **	N017525 006	
+	EQ 25MG BASE **	N017525 007	
+	EQ 50MG BASE **	N017525 008	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LUBIPROSTONE

CAPSULE; ORAL

LUBIPROSTONE

ANCHEN PHARMS

8MCG

A201442 001 Jun 27, 2022

24MCG

A201442 002 Jun 27, 2022

LUCINACTANT

SUSPENSION; INTRATRACHEAL

SURFAXIN

LEES PHARM HK

8.5ML

N021746 001 Mar 06, 2012

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

TEVA PHARMS USA

20MG

A208060 001 May 17, 2019

40MG

A208060 002 May 17, 2019

60MG

A208060 003 May 17, 2019

80MG

A208060 004 May 17, 2019

120MG

A208060 005 May 17, 2019

WATSON LABS TEVA

20MG

A208016 001 Feb 02, 2021

40MG

A208016 002 Feb 02, 2021

60MG

A208016 003 Feb 02, 2021

80MG

A208016 004 Feb 02, 2021

120MG

A208016 005 Feb 02, 2021

LYPRESSIN

SOLUTION; NASAL

DIAPID

NOVARTIS

0.185MG/ML

N016755 001

MACIMORELIN ACETATE

FOR SOLUTION; ORAL

MACRILEN

+ NOVO

EQ 60MG BASE/POUCH

N205598 001 Dec 20, 2017

MACITENTAN

TABLET; ORAL

MACITENTAN

AUROBINDO PHARMA LTD

10MG

A211198 001 Apr 18, 2023

ZYDUS

10MG

A211224 001 Apr 06, 2021

MAFENIDE ACETATE

FOR SOLUTION; TOPICAL

MAFENIDE ACETATE

NOVAST LABS

5%

A206716 001 Jul 31, 2017

PAR FORM

5%

A201511 001 Feb 12, 2013

SULFAMYLON

+ MYLAN INSTITUTIONAL

5%

N019832 003 Jun 05, 1998

MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PLASMA-LYTE 56 IN PLASTIC CONTAINER

BAXTER HLTHCARE

32MG/100ML; 128MG/100ML; 234MG/100ML

N019047 001 Jun 15, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 0.82MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML; 12MG/100ML

N019006 001 Apr 04, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML

N018252 001

SOLUTION; IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER

HOSPIRA INC

14MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML

N018406 001

PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

HOSPIRA INC

30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML

N018406 002 Jul 08, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

SOLUTION;IRRIGATION

SYNOVALYTE IN PLASTIC CONTAINER

BAXTER HLTHCARE	30MG/100ML;37MG/100ML;368MG/100ML;526MG/100ML;502MG/100ML	N019326 001	Jan 25, 1985
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MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE

TABLET;ORAL

MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE

SANTARUS	343MG;20MG;750MG	N022456 001	Dec 04, 2009
	343MG;40MG;750MG	N022456 002	Dec 04, 2009

TABLET, CHEWABLE;ORAL

ZEGERID

SANTARUS	700MG;20MG;600MG	N021850 001	Mar 24, 2006
	700MG;40MG;600MG	N021850 002	Mar 24, 2006

MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; POTASSIUM SULFATE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

SOLUTION;ORAL

SUCLEAR

+ BRAINTREE LABS	1.6GM/BOT,3.13GM/BOT,17.5GM/BOT,N/A,N/A,N/A,N/A;N/A,N/A,N/A,N/A,210GM,0.74GM,2.86GM,5.6GM **	N203595 001	Jan 18, 2013
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MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

POWDER;ORAL

COLPREP KIT

+ GATOR PHARMS	1.6GM/BOT;3.13GM/BOT;17.5GM/BOT	N204553 001	Dec 27, 2016
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MALATHION

LOTION;TOPICAL

MALATHION

MYLAN PHARMS INC	0.5%	A078743 001	Mar 06, 2009
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OVIDE

+ TARO	0.5% **	N018613 001	Aug 02, 1982
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MANGAFODIPIR TRISODIUM

INJECTABLE;INJECTION

TESLASCAN

IC TARGETS	37.9MG/ML	N020652 001	Nov 26, 1997
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MANGANESE CHLORIDE TETRAHYDRATE

FOR SOLUTION;ORAL

LUMENHANCE

BRACCO	3.49MG/GM	N020686 001	Dec 19, 1997
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MANGANESE SULFATE

INJECTABLE;INJECTION

MANGANESE SULFATE

+ ABRAXIS PHARM	EQ 0.1MG MANGANESE/ML **	N019228 001	May 05, 1987
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MANNITOL

INJECTABLE;INJECTION

MANNITOL 10%

B BRAUN	10GM/100ML	N016080 002	
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HOSPIRA	10GM/100ML	N016269 002	
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MILES	10GM/100ML	N016472 002	
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MANNITOL 10% IN PLASTIC CONTAINER

ICU MEDICAL INC	10GM/100ML	N019603 002	Jan 08, 1987
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MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER

B BRAUN	10GM/100ML	N016080 006	
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MANNITOL 15%

B BRAUN	15GM/100ML	N016080 003	
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HOSPIRA	15GM/100ML	N016269 003	
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MILES	15GM/100ML	N016472 005	
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MANNITOL 15% IN PLASTIC CONTAINER

ICU MEDICAL INC	15GM/100ML	N019603 003	Jan 08, 1990
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MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%

B BRAUN	15GM/100ML	N016080 005	
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MANNITOL 20%

B BRAUN	20GM/100ML	N014738 001	
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	20GM/100ML	N016080 004	
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HOSPIRA	20GM/100ML	N016269 004	
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MILES	20GM/100ML	N016472 004	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MANNITOL

INJECTABLE; INJECTION

MANNITOL 25%

ABRAXIS PHARM	12.5GM/50ML	A086754	001	
HOSPIRA	12.5GM/50ML	N016269	005	
IGI LABS INC	12.5GM/50ML	A089239	001	May 06, 1987
	12.5GM/50ML	A089240	001	May 06, 1987
INTL MEDICATION	12.5GM/50ML	A083051	001	
LUITPOLD	12.5GM/50ML	A087409	001	Jan 21, 1982
MERCK	12.5GM/50ML	N005620	001	
WATSON LABS	12.5GM/50ML	A087460	001	Jun 27, 1983

MANNITOL 5%

B BRAUN	5GM/100ML	N016080	001	
HOSPIRA	5GM/100ML	N016269	001	

MANNITOL 5% IN PLASTIC CONTAINER

ICU MEDICAL INC	5GM/100ML	N019603	001	Jan 08, 1987
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MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%

B BRAUN	5GM/100ML	N016080	007	
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SOLUTION; IRRIGATION

RESECTISOL

B BRAUN	5GM/100ML	N016704	002	
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RESECTISOL IN PLASTIC CONTAINER

B BRAUN	5GM/100ML	N016772	002	
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MANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL

HOSPIRA	540MG/100ML; 2.7GM/100ML	A080224	001	
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SORBITOL-MANNITOL IN PLASTIC CONTAINER

HOSPIRA	540MG/100ML; 2.7GM/100ML	N017636	001	
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MAPROTILINE HYDROCHLORIDE

TABLET; ORAL

LUDIOMIL

NOVARTIS	25MG	N017543	001	
	50MG	N017543	002	
	75MG	N017543	003	Sep 30, 1982

MAPROTILINE HYDROCHLORIDE

AM THERAP	25MG	A072129	001	Jan 14, 1988
	50MG	A072130	001	Jan 14, 1988
	75MG	A072131	001	Jan 14, 1988
HERITAGE PHARMA	25MG	A072162	001	Jun 01, 1988
	50MG	A072163	001	Jun 01, 1988
RISING	25MG	A072285	002	Oct 03, 1988
	50MG	A072285	001	Oct 03, 1988
	75MG	A072285	003	Oct 03, 1988
WATSON LABS	25MG	A071943	001	Dec 30, 1987
	50MG	A071944	001	Dec 30, 1987
	75MG	A071945	001	Dec 30, 1987
	75MG	A072164	001	Jun 01, 1988

MASOPROCOL

CREAM; TOPICAL

ACTINEX

UNIV AZ CANCER CTR	10%	N019940	001	Sep 04, 1992
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MAZINDOL

TABLET; ORAL

MAZANOR

WYETH AYERST	1MG	N017980	002	
	2MG	N017980	001	

SANOREX

+ NOVARTIS	1MG **	N017247	001	
+	2MG **	N017247	002	

MEBENDAZOLE

TABLET, CHEWABLE; ORAL

VERMOX

+ JANSSEN PHARMS	100MG **	N017481	001	
+	500MG	N208398	001	Oct 19, 2016

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEBUTAMATE

TABLET; ORAL

DORMATE

MEDPOINTE PHARM HLC 600MG N017374 001

MECAMYLAMINE HYDROCHLORIDE

TABLET; ORAL

INVERSINE

+ TARGACEPT 2.5MG ** N010251 001

MECHLORETHAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

MUSTARGEN

+ RECORDATI RARE 10MG/VIAL N006695 001

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

MECLIZINE HYDROCHLORIDE

ABC HOLDING 12.5MG A085253 001

25MG A085252 001

AMNEAL PHARMS 50MG A201451 003 Feb 23, 2011

ANABOLIC 25MG A085891 001

ANI PHARMS 12.5MG A084657 002

12.5MG A085269 001

12.5MG A088732 001 Dec 11, 1985

25MG A084657 001

25MG A085740 001

BUNDY 12.5MG A084382 001

25MG A084872 001

IVAX SUB TEVA PHARMS 12.5MG A083784 001

KV PHARM 12.5MG A085524 001

25MG A085523 001

PLIVA 25MG A088734 001 Dec 11, 1985

RISING 12.5MG A040179 001 Jan 30, 1997

25MG A040179 002 Jan 30, 1997

STRIDES PHARMA 50MG A089674 001 Mar 31, 1988

SUPERPHARM 12.5MG A089113 001 Aug 20, 1985

25MG A089114 001 Aug 20, 1985

UDL 12.5MG A088256 001 Jun 13, 1983

25MG A088257 001 Jun 13, 1983

VANGARD 12.5MG A087877 001 Apr 20, 1982

25MG A087620 001 Jan 04, 1982

WATSON LABS 12.5MG A085195 001

TABLET, CHEWABLE; ORAL

MECLIZINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS 25MG A084976 001

NEXGEN PHARMA INC 25MG A086392 001

PLIVA 25MG A088733 001 Dec 11, 1985

MECLOCYCLINE SULFOSALICYLATE

CREAM; TOPICAL

MECLAN

JOHNSON AND JOHNSON 1% N050518 001

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLODIUM

QUANTUM PHARMICS EQ 50MG BASE A071380 001 Jul 14, 1987

EQ 100MG BASE A071381 001 Jul 14, 1987

MECLOFENAMATE SODIUM

AM THERAP EQ 50MG BASE A071362 001 Feb 10, 1987

EQ 100MG BASE A071363 001 Feb 10, 1987

ANI PHARMS EQ 50MG BASE A071469 002 Apr 15, 1987

EQ 100MG BASE A071469 001 Apr 15, 1987

BARR EQ 50MG BASE A072848 001 Mar 20, 1989

EQ 100MG BASE A072809 001 Mar 20, 1989

CHARTWELL RX EQ 50MG BASE A072262 001 Nov 29, 1988

EQ 100MG BASE A072263 001 Nov 29, 1988

PAR PHARM EQ 50MG BASE A072077 001 Mar 10, 1988

EQ 100MG BASE A072078 001 Mar 10, 1988

USL PHARMA EQ 50MG BASE A071007 001 Mar 25, 1988

EQ 100MG BASE A071008 001 Mar 25, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLOFENAMATE SODIUM

VITARINE	EQ 50MG BASE	A071710 001	Jun 15, 1988
	EQ 100MG BASE	A071684 001	Jun 15, 1988
WATSON LABS	EQ 50MG BASE	A070400 001	Nov 25, 1986
	EQ 50MG BASE	A071640 001	Aug 11, 1987
	EQ 100MG BASE	A070401 001	Nov 25, 1986
	EQ 100MG BASE	A071641 001	Aug 11, 1987
MECLOMEN			
PARKE DAVIS	EQ 50MG BASE	N018006 001	
	EQ 100MG BASE	N018006 002	

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

+	PFIZER	100MG/ML **	N012541 002
+		400MG/ML	N012541 003

MEDROXYPROGESTERONE ACETATE

CIPLA	150MG/ML	A210335 001	Jan 25, 2019
HIKMA	150MG/ML	A214309 001	Jan 05, 2023
SANDOZ	150MG/ML	A078711 001	May 20, 2009
SUN PHARM	150MG/ML	A210760 001	May 01, 2019
	150MG/ML	A210761 001	Apr 24, 2019
TEVA PHARMS USA	150MG/ML	A076552 001	Oct 27, 2004

TABLET; ORAL

AMEN

AMARIN PHARMS	10MG	A083242 001	
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CURRETAB

SOLVAY	10MG	A085686 001	
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CYCRIN

ESI	2.5MG	A081239 001	Oct 30, 1992
	5MG	A081240 001	Oct 30, 1992
	10MG	A089386 001	Sep 09, 1987

MEDROXYPROGESTERONE ACETATE

DURAMED PHARMS BARR	2.5MG	A040311 001	Dec 01, 1999
	5MG	A040311 002	Dec 01, 1999
	10MG	A040311 003	Dec 01, 1999
UPSHER SMITH LABS	10MG	A088484 001	Jul 26, 1984

MEDRYSONE

SUSPENSION; OPHTHALMIC

HMS

ALLERGAN	1%	N016624 003	
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MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

CHARTWELL RX	250MG	A209209 001	Sep 18, 2020
NOSTRUM LABS INC	250MG	A090359 001	Feb 05, 2013

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

LARIAM

+	ROCHE	250MG **	N019591 001	May 02, 1989
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MEFLOQUINE HYDROCHLORIDE

HIKMA INTL PHARMS	250MG	A077699 001	Apr 21, 2010
SANDOZ	250MG	A076175 001	Feb 20, 2002
US ARMY WALTER REED	250MG **	N019578 001	May 02, 1989

MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE

+	BRISTOL MYERS SQUIBB	40MG/ML **	N020264 001	Sep 10, 1993
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MEGESTROL ACETATE

BIONPHARMA	125MG/ML	A204688 001	Dec 01, 2017
HIKMA	40MG/ML	A075997 001	Feb 15, 2002
NOVITIUM PHARMA	40MG/ML	A077404 001	Feb 16, 2006
TEVA PHARMS	40MG/ML	A075681 001	May 05, 2003
XTRIUM LABS INC	40MG/ML	A076721 001	Nov 01, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEGESTROL ACETATE

TABLET; ORAL

MEGACE

+ BRISTOL MYERS SQUIBB	20MG **	N016979	001	
+	40MG **	N016979	002	

MEGESTROL ACETATE

HIKMA	20MG	A074458	001	Sep 29, 1995
	40MG	A074458	002	Sep 29, 1995
TEVA	40MG	A074745	001	Feb 27, 1998
USL PHARMA	20MG	A070646	001	Oct 02, 1987
	40MG	A070647	001	Oct 02, 1987

MELOXICAM

CAPSULE; ORAL

VIVLODEX

+ ICEUTICA OPERATIONS	5MG **	N207233	001	Oct 22, 2015
+	10MG **	N207233	002	Oct 22, 2015

SOLUTION; INTRAVENOUS

ANJESO

+ BAUDAX	30MG/ML (30MG/ML) **	N210583	001	Feb 20, 2020
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TABLET; ORAL

MELOXICAM

ANDA REPOSITORY	7.5MG	A077935	001	Jul 19, 2006
	15MG	A077935	002	Jul 19, 2006
CHARTWELL RX	7.5MG	A077936	001	Jul 19, 2006
	15MG	A077936	002	Jul 19, 2006
CR DOUBLE CRANE	7.5MG	A078039	001	Dec 14, 2006
	15MG	A078039	002	Dec 14, 2006
IMPAX LABS INC	7.5MG	A077930	001	Jul 19, 2006
	15MG	A077930	002	Jul 19, 2006
MYLAN	7.5MG	A077923	001	Jul 19, 2006
	7.5MG	A077934	001	Jul 20, 2006
	15MG	A077923	002	Jul 19, 2006
	15MG	A077934	002	Jul 20, 2006
ROXANE	7.5MG	A077925	001	Jul 19, 2006
	15MG	A077925	002	Jul 19, 2006
SCIEGEN PHARMS INC	7.5MG	A077920	001	Jul 19, 2006
	15MG	A077920	002	Jul 19, 2006
SUN PHARM INDS INC	7.5MG	A077937	001	Jul 19, 2006
	15MG	A077937	002	Jul 19, 2006
YABAO PHARM	7.5MG	A077933	001	Jul 19, 2006
	15MG	A077933	002	Jul 19, 2006

TABLET, ORALLY DISINTEGRATING; ORAL

QMIIZ ODT

+ TERSERA	7.5MG	N211210	001	Oct 19, 2018
+	15MG	N211210	002	Oct 19, 2018

MELPHALAN

TABLET; ORAL

ALKERAN

+ APOTEX	2MG **	N014691	002	
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MELPHALAN FLUFENAMIDE HYDROCHLORIDE

POWDER; INTRAVENOUS

PEPAXTO

+ ONCOPEPTIDES AB	EQ 20MG BASE/VIAL	N214383	001	Feb 26, 2021
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MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

ALKERAN

+ APOTEX	EQ 50MG BASE/VIAL **	N020207	001	Nov 18, 1992
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MELPHALAN HYDROCHLORIDE

MYLAN INSTITUTIONAL	EQ 50MG BASE/VIAL	A090299	001	Oct 27, 2009
PAR STERILE PRODUCTS	EQ 50MG BASE/VIAL	A204773	001	Aug 22, 2016
USWM	EQ 50MG BASE/VIAL	A207032	001	May 03, 2019

POWDER; INTRAVENOUS

MELPHALAN HYDROCHLORIDE

ACTAVIS LLC	EQ 50MG BASE/VIAL	A209323	001	Mar 06, 2020
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

MEMANTINE HYDROCHLORIDE

ANCHEN PHARMS	7MG	A205784 001	Jun 09, 2017
	14MG	A205784 002	Jun 09, 2017
	21MG	A205784 003	Jun 09, 2017
	28MG	A205784 004	Jun 09, 2017
ANI PHARMS	7MG	A205365 001	Feb 28, 2020
	14MG	A205365 002	Feb 28, 2020
	21MG	A205365 003	Feb 28, 2020
	28MG	A205365 004	Feb 28, 2020
RISING	7MG	A206032 001	Sep 28, 2016
	14MG	A206032 002	Sep 28, 2016
	21MG	A206032 003	Sep 28, 2016
	28MG	A206032 004	Sep 28, 2016
SUN PHARM	7MG	A205905 001	Sep 28, 2016
	14MG	A205905 002	Sep 28, 2016
	21MG	A205905 003	Sep 28, 2016
	28MG	A205905 004	Sep 28, 2016

NAMENDA XR

+ ABBVIE 7MG

N022525 001 Jun 21, 2010

SOLUTION;ORAL

MEMANTINE HYDROCHLORIDE

APOZEAL PHARMS 2MG/ML

A205446 001 Dec 07, 2015

NAMENDA

+ ALLERGAN 2MG/ML

N021627 001 Apr 18, 2005

TABLET;ORAL

MEMANTINE HYDROCHLORIDE

CHARTWELL	5MG	A090244 001	Jul 11, 2018
	10MG	A090244 002	Jul 11, 2018
HIKMA	5MG	A208173 001	Feb 28, 2020
	10MG	A208173 002	Feb 28, 2020
JUBILANT GENERICS	5MG	A091585 001	Oct 13, 2015
	10MG	A091585 002	Oct 13, 2015
LANNETT CO INC	5MG	A207236 001	Nov 10, 2016
	10MG	A207236 002	Nov 10, 2016
ORBION PHARMS	5MG	A090044 001	Mar 12, 2012
	10MG	A090044 002	Mar 12, 2012
RISING	5MG	A079225 001	Jan 30, 2015
	10MG	A079225 002	Jan 30, 2015
TEVA PHARMS	5MG	A090052 001	Oct 25, 2011
	10MG	A090052 002	Oct 25, 2011
TORRENT	5MG	A200155 001	Oct 13, 2015
	10MG	A200155 002	Oct 13, 2015
YILING	5MG	A212947 001	Apr 03, 2020
	10MG	A212947 002	Apr 03, 2020

MENADIOL SODIUM DIPHOSPHATE

INJECTABLE; INJECTION

KAPPADIONE

LILLY 10MG/ML

N005725 001

SYNKAYVITE

ROCHE	5MG/ML	N003718 004
	10MG/ML	N003718 006
	37.5MG/ML	N003718 008

TABLET;ORAL

SYNKAYVITE

ROCHE 5MG N003718 010

MENADIONE

TABLET;ORAL

MENADIONE

LILLY 5MG N002139 003

MEPENZOLATE BROMIDE

SOLUTION;ORAL

CANTIL

SANOFI AVENTIS US 25MG/5ML N010679 004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPENZOLATE BROMIDE

TABLET; ORAL

CANTIL

+ SANOFI AVENTIS US 25MG N010679 003

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

VALIDUS PHARMS	25MG/ML	N005010	007
	50MG/ML	N005010	002
	75MG/ML	N005010	009
	100MG/ML	N005010	003

MEPERIDINE HYDROCHLORIDE

ABBOTT

25MG/ML	A080388	001
50MG/ML	A080385	001
50MG/ML	A080387	001
75MG/ML	A080389	001
100MG/ML	A080386	001

BAXTER HLTHCARE

25MG/ML	A088279	001	Jun 15, 1984
50MG/ML	A088280	001	Jun 15, 1984
75MG/ML	A088281	001	Jun 15, 1984
100MG/ML	A088282	001	Jun 15, 1984

IGI LABS INC

25MG/ML	A089781	001	Mar 31, 1989
50MG/ML	A089782	001	Mar 31, 1989
50MG/ML	A089783	001	Mar 31, 1989
50MG/ML	A089784	001	Mar 31, 1989
75MG/ML	A089785	001	Mar 31, 1989
100MG/ML	A089786	001	Mar 31, 1989
100MG/ML	A089787	001	Mar 31, 1989
100MG/ML	A089788	001	Mar 31, 1989

INTL MEDICATION

10MG/ML A086332 001

PARKE DAVIS

50MG/ML A080364 002

75MG/ML A080364 003

100MG/ML A080364 001

WATSON LABS

50MG/ML A073444 001 Mar 17, 1992

100MG/ML A073445 001 Mar 17, 1992

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

HOSPIRA

10MG/ML A040305 001 Mar 10, 1999

+ ICU MEDICAL INC

10MG/ML A088432 001 Aug 16, 1984

INTL MEDICATION

10MG/ML A081309 001 Aug 30, 1993

SPECGX LLC

10MG/ML A040163 001 May 12, 1997

WATSON LABS

10MG/ML A073443 001 Mar 17, 1992

SYRUP; ORAL

DEMEROL

VALIDUS PHARMS 50MG/5ML ** N005010 005

TABLET; ORAL

DEMEROL

+ VALIDUS PHARMS 50MG ** N005010 001

+ 100MG N005010 004

MEPERIDINE HYDROCHLORIDE

ANDA REPOSITORY

50MG A040893 001 Jun 24, 2009

75MG A040893 002 Jun 24, 2009

100MG A040893 003 Jun 24, 2009

150MG A040893 004 Jun 24, 2009

BARR

50MG A088639 001 Jul 02, 1984

100MG A088640 001 Sep 19, 1984

DURAMED PHARMS BARR

50MG A040318 001 Oct 05, 1999

100MG A040318 002 Oct 05, 1999

HIKMA

50MG A040110 001 Mar 12, 1997

100MG A040110 002 Mar 12, 1997

SPECGX LLC

50MG A040352 001 Jun 13, 2000

100MG A040352 002 Jun 13, 2000

STRIDES PHARMA

50MG A040191 001 Dec 17, 1998

100MG A040191 002 Dec 17, 1998

SUN PHARM INDS INC

50MG A040446 001 Aug 08, 2002

100MG A040446 002 Aug 08, 2002

SUN PHARM INDUSTRIES

50MG A080448 001

100MG A080448 002

WATSON LABS

50MG A040186 001 Jun 30, 1997

100MG A040186 002 Jun 30, 1997

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPERIDINE HYDROCHLORIDE

TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

WYETH AYERST 50MG A080454 001

MEPERIDINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERGAN

HIKMA 25MG/ML; 25MG/ML N011730 001

MEPHENTERMINE SULFATE

INJECTABLE; INJECTION

WYAMINE SULFATE

BAXTER HLTHCARE CORP EQ 15MG BASE/ML N008248 002

EQ 30MG BASE/ML N008248 001

MEPHENYTOIN

TABLET; ORAL

MESANTOIN

+ NOVARTIS 100MG ** N006008 001

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE

SOLVAY 3% A084777 002 Apr 18, 1982

CARBOCAINE

+ EASTMAN KODAK 3% ** N012125 003

ISOCAINE HYDROCHLORIDE

SEPTODONT 3% A080925 001

MEPIVACAINE HYDROCHLORIDE

BELMORA LLC 3% A083559 001

HOSPIRA INC 3% A040806 001 Apr 28, 2008

INTL MEDICATION SYS 1% A087509 001 Oct 05, 1982

WATSON LABS 1% A088769 001 Nov 20, 1984

2% A088770 001 Nov 20, 1984

POLOCAINE

DENTSPLY PHARM 3% A088653 001 Aug 21, 1984

MEPREDNISONE

TABLET; ORAL

BETAPAR

SCHERING 4MG N016053 002

MEPROBAMATE

CAPSULE; ORAL

EQUANIL

WYETH AYERST 400MG N012455 002

CAPSULE, EXTENDED RELEASE; ORAL

MEPROSPAN

MEDPOINTE PHARM HLC 200MG N011284 001

400MG N011284 002

TABLET; ORAL

AMOSENE

FERNDALE LABS 400MG A084030 001

BAMATE

ALRA 200MG A080380 001

400MG A080380 002

EQUANIL

WYETH AYERST 200MG N010028 005

400MG N010028 004

MEPRIAM

TEVA 400MG N016069 001

MEPROBAMATE

ACELLA 400MG A084153 001

BARR 600MG A084230 001

CHARTWELL MOLECULAR 200MG N014882 002

400MG N014882 001

ELKINS SINN 200MG N015426 002

400MG N015426 001

HEATHER 400MG N016928 003

600MG A084329 001

IMPAX LABS 200MG N014322 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPROBAMATE

TABLET;ORAL

MEPROBAMATE

	400MG		N014322	001	
IVAX SUB TEVA PHARMS	200MG		N015438	001	
	400MG		N015438	002	
	600MG		A084181	001	
LEDERLE	400MG		A086299	001	
LEE KM	400MG		A089538	001	Nov 25, 1987
MALLARD	400MG		N015072	002	
MK LABS	200MG		N014368	004	
	400MG		N014368	002	
MYLAN	400MG		A083618	001	
NEXGEN PHARMA INC	200MG		A084220	001	
	400MG		A084589	001	
PARKE DAVIS	200MG		A084744	001	
	400MG		A084744	002	
PERRIGO	200MG		A084546	001	
	400MG		A084547	001	
PHARMAVITE	400MG		A084438	001	
PUREPAC PHARM	200MG		A084804	001	
	400MG		A084804	002	
PVT FORM	400MG		N014601	001	
RISING	400MG		A080655	001	
ROXANE	600MG		A084332	001	
SANDOZ	200MG		N014547	002	
	400MG		N014547	001	
SCHERER LABS	400MG		A083343	001	
SOLVAY	200MG		A084435	001	
STANLABS PHARM	200MG		N014474	002	
	400MG		N014474	004	
SUN PHARM INDUSTRIES	200MG		A080699	001	
	400MG		A080699	002	
TABLICAPS	400MG		A083494	001	
TARO	200MG		A200998	001	May 23, 2011
	400MG		A200998	002	May 23, 2011
USL PHARMA	200MG		A087825	001	Mar 18, 1982
	400MG		A087826	001	Mar 18, 1982
VALEANT PHARM INTL	200MG		N015139	006	
	400MG		N015139	005	
VANGARD	400MG		A088011	001	Jul 14, 1982
+ WATSON LABS	200MG		A083304	001	
	200MG		A085720	001	
	400MG		A083308	001	
	400MG		A085721	001	
	600MG		A084274	001	
	600MG		A085719	001	
WEST WARD	200MG		N015417	003	
	400MG		N015417	002	
WHITEWORTH TOWN PLSN	200MG		A083830	001	
	400MG		A083442	001	
MILTOWN					
+ MEDPOINTE PHARM HLC	200MG **		N009698	004	
	400MG **		N009698	002	
	600MG		A083919	001	
NEURAMATE					
HALSEY	200MG		N014359	002	
	400MG		N014359	001	
TRANMEP					
SOLVAY	400MG		A084369	001	
	400MG		N016249	001	

MEQUINOL; TRETINOIN

SOLUTION;TOPICAL

SOLAGE

ALMIRALL	2%;0.01%		N020922	001	Dec 10, 1999
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

HOSPIRA INC	500MG/VIAL	A090940 001	Jun 22, 2010
	1GM/VIAL	A090940 002	Jun 22, 2010
SANDOZ	500MG/VIAL	A091201 001	Mar 29, 2011
	1GM/VIAL	A091201 002	Mar 29, 2011

MERSALYL SODIUM; THEOPHYLLINE

INJECTABLE; INJECTION

MERSALYL-THEOPHYLLINE

WATSON LABS	100MG/ML; 50MG/ML	A084875 001	
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MESALAMINE

CAPSULE, EXTENDED RELEASE; ORAL

MESALAMINE

TEVA PHARMS USA	375MG	A209970 001	May 06, 2022
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ENEMA; RECTAL

MESALAMINE

G AND W LABS INC	4GM/60ML	A076841 001	Sep 30, 2004
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SUPPOSITORY; RECTAL

CANASA

ABBVIE	500MG	N021252 001	Jan 05, 2001
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MESALAMINE

AMNEAL	1GM	A210509 001	Jan 02, 2020
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ROWASA

+ MEDA PHARMS

500MG **	N019919 001	Dec 18, 1990
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TABLET, DELAYED RELEASE; ORAL

ASACOL

APIL	400MG	N019651 001	Jan 31, 1992
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ASACOL HD

+ ABBVIE

800MG **	N021830 001	May 29, 2008
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MESALAMINE

MYLAN	1.2GM	A203574 001	Nov 20, 2018
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MESNA

INJECTABLE; INTRAVENOUS

MESNA

MYLAN INSTITUTIONAL	100MG/ML	A076488 001	Mar 08, 2012
MYLAN LABS LTD	100MG/ML	A203364 001	Jul 18, 2014
TEVA PHARMS USA	100MG/ML	A075764 001	Apr 27, 2001

MESORIDAZINE BESYLATE

CONCENTRATE; ORAL

SERENTIL

NOVARTIS	EQ 25MG BASE/ML	N016997 001	
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INJECTABLE; INJECTION

SERENTIL

NOVARTIS	EQ 25MG BASE/ML	N016775 001	
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TABLET; ORAL

SERENTIL

NOVARTIS	EQ 10MG BASE **	N016774 001	
	EQ 25MG BASE **	N016774 002	
	EQ 50MG BASE **	N016774 003	
	EQ 100MG BASE **	N016774 004	

MESTRANOL; NORETHINDRONE

TABLET; ORAL-20

NORINYL

ACTAVIS LABS UT INC	0.1MG; 2MG	N013625 004	
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TABLET; ORAL-21

NORETHIN 1/50M-21

HERITAGE PHARMA	0.05MG; 1MG	A071539 001	Apr 12, 1988
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NORETHINDRONE AND MESTRANOL

WATSON LABS	0.05MG; 1MG	A070758 001	Jul 01, 1988
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NORINYL 1+50 21-DAY

ACTAVIS LABS UT INC	0.05MG; 1MG	N013625 002	
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NORINYL 1+80 21-DAY

GD SEARLE LLC	0.08MG; 1MG	N016724 001	
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ORTHO-NOVUM 1/50 21

ORTHO MCNEIL PHARM	0.05MG; 1MG	N012728 004	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MESTRANOL; NORETHINDRONE

TABLET; ORAL-21

ORTHO-NOVUM 1/80 21				
ORTHO MCNEIL PHARM	0.08MG;1MG		N016715	001
ORTHO-NOVUM 10-21				
ORTHO MCNEIL PHARM	0.06MG;10MG		N012728	001
ORTHO-NOVUM 2-21				
ORTHO MCNEIL PHARM	0.1MG;2MG		N012728	005

TABLET; ORAL-28

NORETHIN 1/50M-28				
HERITAGE PHARMA	0.05MG;1MG		A071540	001 Apr 12, 1988
NORETHINDRONE AND MESTRANOL				
WATSON LABS	0.05MG;1MG		A070759	001 Jul 01, 1988
NORINYL 1+50 28-DAY				
+ ACTAVIS LABS UT INC	0.05MG;1MG		N016659	001
NORINYL 1+80 28-DAY				
GD SEARLE LLC	0.08MG;1MG		N016725	001
ORTHO-NOVUM 1/50 28				
ORTHO MCNEIL JANSSEN	0.05MG;1MG		N016709	001
ORTHO-NOVUM 1/80 28				
ORTHO MCNEIL PHARM	0.08MG;1MG		N016715	002

MESTRANOL; NORETHYNODREL

TABLET; ORAL

ENOVID				
GD SEARLE LLC	0.075MG;5MG		N010976	008
	0.15MG;9.85MG		N010976	005

TABLET; ORAL-20

ENOVID				
GD SEARLE LLC	0.075MG;5MG		N010976	004
ENOVID-E				
GD SEARLE LLC	0.1MG;2.5MG		N010976	006

TABLET; ORAL-21

ENOVID-E 21				
GD SEARLE LLC	0.1MG;2.5MG		N010976	007

METAPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

ALUPENT				
BOEHRINGER INGELHEIM	0.65MG/INH		N016402	001

SOLUTION; INHALATION

ALUPENT				
BOEHRINGER INGELHEIM	0.4%		N018761	002 Oct 10, 1986
	0.6%		N018761	001 Jun 30, 1983
	5%		N017659	001

METAPROTERENOL SULFATE

APOTEX INC	0.4%		A075402	001 Feb 28, 2001
	0.6%		A075403	001 Feb 28, 2001
ASTRAZENECA	0.4%		A071275	001 Jul 27, 1988
	0.6%		A071018	001 Jul 27, 1988
DEY	0.33%		A071806	001 Aug 05, 1988
	0.5%		A071805	001 Aug 05, 1988
	5%		A070805	001 Aug 17, 1987
MYLAN SPECIALITY LP	0.4%		A071786	001 Aug 05, 1988
	0.6%		A070804	001 Aug 17, 1987
NEPHRON	0.4%		A071855	001 Jul 14, 1988
	0.6%		A071726	001 Jul 14, 1988
WOCKHARDT	0.4%		A075586	001 May 30, 2002
	0.6%		A075586	002 May 30, 2002
	5%		A072190	001 Jun 07, 1988
PROMETA				
MURO	5%		A073340	001 Mar 30, 1992

SYRUP; ORAL

ALUPENT				
BOEHRINGER INGELHEIM	10MG/5ML		N017571	001

METAPROTERENOL SULFATE

APOTEX INC	10MG/5ML		A075235	001 Jan 27, 2000
COSETTE	10MG/5ML		A072761	001 Feb 27, 1992
G AND W LABS INC	10MG/5ML		A073034	001 Aug 30, 1991
MORTON GROVE	10MG/5ML		A071656	001 Oct 13, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METAPROTERENOL SULFATE

SYRUP;ORAL

METAPROTERENOL SULFATE

WOCKHARDT 10MG/5ML

A074702 001 Mar 24, 1997

PROMETA

MURO 10MG/5ML

A072023 001 Sep 15, 1988

TABLET;ORAL

ALUPENT

BOEHRINGER INGELHEIM 10MG

N015874 002

20MG

N015874 001

METAPROTERENOL SULFATE

AM THERAP 10MG

A072054 001 Jun 23, 1988

20MG

A072055 001 Jun 23, 1988

HERITAGE PHARMA 10MG

A072519 001 Mar 30, 1990

20MG

A072520 001 Mar 30, 1990

STRIDES PHARMA 10MG

A072024 001 Jun 28, 1988

20MG

A072025 001 Jun 28, 1988

USL PHARMA 10MG

A071013 001 Jan 25, 1988

20MG

A071014 001 Jan 25, 1988

WATSON LABS 10MG

A073013 001 Jan 31, 1991

20MG

A072795 001 Jan 31, 1991

METARAMINOL BITARTRATE

INJECTABLE;INJECTION

ARAMINE

+ MERCK EQ 10MG BASE/ML **

N009509 002 Dec 22, 1987

METARAMINOL BITARTRATE

ABRAXIS PHARM EQ 10MG BASE/ML

A080431 001

ELKINS SINN EQ 10MG BASE/ML

A083363 001

FRESENIUS KABI USA EQ 10MG BASE/ML

A080722 001

GD SEARLE LLC EQ 10MG BASE/ML

A086418 001

EQ 20MG BASE/ML

A086418 002

METAXALONE

TABLET;ORAL

METAXALONE

INGENUS PHARMS LLC 800MG

A213836 001 Oct 21, 2020

+ PRIMUS PHARMS 640MG **

N022503 001 Jun 01, 2015

SKELAXIN

+ KING PHARMS 400MG **

N013217 001

+ 800MG

N013217 003 Aug 30, 2002

METFORMIN HYDROCHLORIDE

FOR SUSPENSION, EXTENDED RELEASE;ORAL

RIOMET ER

+ SUN PHARM 500MG/5ML

N212595 001 Aug 29, 2019

SOLUTION;ORAL

RIOMET

+ RANBAXY 500MG/5ML

N021591 001 Sep 11, 2003

TABLET;ORAL

GLUCOPHAGE

+ EMD SERONO INC 500MG **

N020357 001 Mar 03, 1995

+ 625MG **

N020357 003 Nov 05, 1998

+ 750MG **

N020357 004 Nov 05, 1998

+ 850MG **

N020357 002 Mar 03, 1995

+ 1GM **

N020357 005 Nov 05, 1998

METFORMIN HYDROCHLORIDE

BARR 500MG

A075971 001 Jan 25, 2002

850MG

A075971 002 Jan 25, 2002

1GM

A075971 003 Jan 25, 2002

HERITAGE PHARMA 500MG

A075978 001 Jan 25, 2002

850MG

A075978 002 Jan 25, 2002

1GM

A075978 003 Nov 05, 2002

INDICUS PHARMA 500MG

A079148 001 Nov 25, 2008

850MG

A079148 002 Nov 25, 2008

1GM

A079148 003 Nov 25, 2008

IPCA LABS LTD 500MG

A078422 001 Aug 06, 2007

850MG

A078422 002 Aug 06, 2007

1GM

A078422 003 Aug 06, 2007

IVAX SUB TEVA PHARMS 500MG

A075975 001 Jan 24, 2002

625MG

A075975 004 Jan 24, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METFORMIN HYDROCHLORIDE

TABLET;ORAL

METFORMIN HYDROCHLORIDE

	750MG	A075975 005	Jan 24, 2002
	850MG	A075975 002	Jan 24, 2002
	1GM	A075975 003	Jan 24, 2002
MACLEODS PHARMS LTD	500MG	A205330 001	Oct 31, 2017
	850MG	A205330 002	Oct 31, 2017
	1GM	A205330 003	Oct 31, 2017
MYLAN	500MG	A075976 001	Jan 24, 2002
	850MG	A075976 002	Jan 24, 2002
	1GM	A075976 003	Jan 24, 2002
MYLAN PHARMS INC	500MG	A075969 001	Jan 29, 2002
	850MG	A075969 002	Jan 29, 2002
	1GM	A075969 003	Jan 29, 2002
PROVIDENT PHARM	500MG	A077853 001	Jul 28, 2006
	850MG	A077853 002	Jul 28, 2006
	1GM	A077853 003	Jul 28, 2006
SANDOZ	500MG	A075985 001	Jan 25, 2002
	850MG	A075985 002	Jan 25, 2002
	1GM	A075985 003	Jan 25, 2002
SUN PHARM INDS INC	500MG	A075967 001	Jan 29, 2002
	850MG	A075967 002	Jan 29, 2002
	1GM	A075967 003	Jan 29, 2002
SUN PHARM INDUSTRIES	500MG	A076038 001	Feb 21, 2002
	850MG	A076038 002	Feb 21, 2002
	1GM	A076038 003	Feb 21, 2002
SUNSHINE	500MG	A208999 001	Oct 12, 2018
	850MG	A208999 002	Oct 12, 2018
	1GM	A208999 003	Oct 12, 2018
TEVA	500MG	A076328 001	Dec 16, 2002
	850MG	A076328 002	Dec 16, 2002
	1GM	A076328 003	Dec 16, 2002
TORRENT PHARMS	500MG	A077711 001	Jan 24, 2007
	850MG	A077711 002	Jan 24, 2007
	1GM	A077711 003	Jan 24, 2007
WATSON LABS	500MG	A075979 001	Jan 24, 2002
	850MG	A075979 002	Jan 24, 2002
	1GM	A075979 003	Jan 24, 2002
WATSON LABS FLORIDA	500MG	A075961 001	Jan 25, 2002
	850MG	A075961 002	Jan 25, 2002
	1GM	A075961 003	Jan 25, 2002
TABLET, EXTENDED RELEASE;ORAL			
FORTAMET			
+ ANDRX LABS LLC	500MG **	N021574 001	Apr 27, 2004
GLUCOPHAGE XR			
+ EMD SERONO INC	500MG **	N021202 001	Oct 13, 2000
+	750MG **	N021202 004	Apr 11, 2003
METFORMIN HYDROCHLORIDE			
ACTAVIS ELIZABETH	500MG	A076450 001	Oct 01, 2004
	750MG	A076878 001	Apr 13, 2005
ACTAVIS LABS FL INC	500MG	A076172 001	Jun 16, 2004
AMTA	500MG	A213394 001	Aug 03, 2021
	1GM	A213394 002	Aug 03, 2021
APOTEX	500MG	A076706 001	Dec 14, 2004
	750MG	A076706 002	Dec 29, 2005
BARR	500MG	A076496 001	Nov 25, 2005
	750MG	A076863 001	Oct 14, 2004
IMPAX LABS	500MG	A076249 001	Jul 30, 2004
	750MG	A076985 001	Sep 13, 2005
IVAX SUB TEVA PHARMS	500MG	A076545 001	Dec 01, 2003
MYLAN	500MG	A076650 001	Sep 13, 2005
	750MG	A077113 001	Sep 08, 2005
RANBAXY LABS LTD	500MG	A076413 001	Jun 18, 2004
	750MG	A077211 001	Jun 29, 2005
SANDOZ	500MG	A076223 001	Dec 14, 2004
SUN PHARM INDUSTRIES	500MG	A077124 001	Dec 21, 2005
TORRENT	500MG	A090014 001	Dec 30, 2009
TORRENT PHARMS LTD	750MG	A079226 001	Feb 18, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

WATSON LABS INC 500MG

A076818 001 Dec 14, 2004

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOPLUS MET

+ TAKEDA PHARMS USA 500MG;EQ 15MG BASE **

N021842 001 Aug 29, 2005

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

MYLAN 500MG;EQ 15MG BASE

A090406 001 Feb 25, 2011

850MG;EQ 15MG BASE

A090406 002 Feb 25, 2011

SANDOZ 500MG;EQ 15MG BASE

A091273 001 Apr 16, 2013

850MG;EQ 15MG BASE

A091273 002 Apr 16, 2013

TORRENT PHARMS LTD 500MG;EQ 15MG BASE

A202001 001 Feb 13, 2013

850MG;EQ 15MG BASE

A202001 002 Feb 13, 2013

TABLET, EXTENDED RELEASE;ORAL

ACTOPLUS MET XR

+ TAKEDA PHARMS USA 1GM;EQ 15MG BASE

N022024 001 May 12, 2009

+ 1GM;EQ 30MG BASE

N022024 002 May 12, 2009

METFORMIN HYDROCHLORIDE; REPAGLINIDE

TABLET;ORAL

PRANDIMET

+ NOVO NORDISK INC 500MG;1MG

N022386 001 Jun 23, 2008

+ 500MG;2MG

N022386 002 Jun 23, 2008

REPAGLINIDE AND METFORMIN HYDROCHLORIDE

LUPIN LTD 500MG;1MG

A200624 001 Jul 15, 2015

500MG;2MG

A200624 002 Jul 15, 2015

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET;ORAL

AVANDAMET

+ SB PHARMCO 500MG;EQ 1MG BASE **

N021410 001 Oct 10, 2002

+ 500MG;EQ 2MG BASE **

N021410 002 Oct 10, 2002

+ 500MG;EQ 4MG BASE **

N021410 003 Oct 10, 2002

+ 1GM;EQ 2MG BASE **

N021410 004 Aug 25, 2003

+ 1GM;EQ 4MG BASE **

N021410 005 Aug 25, 2003

ROSIGLITAZONE MALEATE AND METFORMIN HYDROCHLORIDE

TEVA 500MG;EQ 2MG BASE

A077337 001 May 07, 2014

500MG;EQ 1MG BASE

A077337 005 May 19, 2017

500MG;EQ 4MG BASE

A077337 002 May 07, 2014

1GM;EQ 4MG BASE

A077337 004 May 07, 2014

1GM;EQ 2MG BASE

A077337 003 May 07, 2014

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

KOMBIGLYZE XR

+ ASTRAZENECA AB 500MG;EQ 5MG BASE

N200678 001 Nov 05, 2010

+ 1GM;EQ 2.5MG BASE

N200678 003 Nov 05, 2010

+ 1GM;EQ 5MG BASE

N200678 002 Nov 05, 2010

METHACHOLINE CHLORIDE

FOR SOLUTION;INHALATION

PROVOCHOLINE

+ METHAPHARM 1600MG/VIAL

N019193 002 Aug 29, 2016

METHACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

RONDONMYCIN

MEDPOINTE PHARM HLC EQ 140MG BASE

A060641 001

EQ 280MG BASE

A060641 002

SYRUP;ORAL

RONDONMYCIN

MEDPOINTE PHARM HLC EQ 70MG BASE/5ML

A060641 003

METHADONE HYDROCHLORIDE

CONCENTRATE;ORAL

METHADONE HYDROCHLORIDE

LANNETT CO INC 10MG/ML

A212094 001 Mar 03, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHADONE HYDROCHLORIDE

POWDER; FOR RX COMPOUNDING

METHADONE HYDROCHLORIDE

MALLINCKRODT INC

50GM/BOT

N006383 002

100GM/BOT

N006383 003

500GM/BOT

N006383 004

SYRUP; ORAL

DOLOPHINE HYDROCHLORIDE

HIKMA

10MG/30ML

N006134 004

TABLET; ORAL

DOLOPHINE HYDROCHLORIDE

+ HIKMA

5MG **

N006134 002

+

10MG **

N006134 010

METHADONE HYDROCHLORIDE

NOSTRUM LABS INC

5MG

A210484 001 Aug 02, 2018

10MG

A210484 002 Aug 02, 2018

ROXANE

40MG

A074081 001 Apr 28, 1995

SUN PHARM INDUSTRIES

5MG

A208305 001 Mar 30, 2018

10MG

A208305 002 Mar 30, 2018

VISTAPHARM

5MG

A040241 001 May 29, 1998

METHADOSE

SPECGX LLC

5MG

A040050 001 Apr 15, 1993

10MG

A040050 002 Apr 15, 1993

TABLET, DISPERSIBLE; ORAL

WESTADONE

SANDOZ

2.5MG

N017108 001

TABLET, EFFERVESCENT; ORAL

WESTADONE

SANDOZ

5MG

N017108 002

10MG

N017108 003

40MG

N017108 004

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

METHAMPEX

TEVA

10MG

A083889 001

METHAMPHETAMINE HYDROCHLORIDE

ABLE

5MG

A040529 001 Feb 25, 2004

REXAR

5MG

A084931 001

10MG

A084931 002

TEVA

5MG

A086359 001

TABLET, EXTENDED RELEASE; ORAL

DESOXYN

AJENAT PHARMS

5MG

N005378 004

10MG

N005378 003

15MG

N005378 005

METHANTHELIN BROMIDE

TABLET; ORAL

BANTHINE

SHIRE

50MG

N007390 001

METHARBITAL

TABLET; ORAL

GEMONIL

ABBVIE

100MG

N008322 001

METHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

APPLIED ANAL

25MG

A040011 001 Jul 17, 1997

50MG

A040011 002 Jul 17, 1997

ATHEM

25MG

A040102 001 Aug 28, 1996

50MG

A040102 002 Aug 28, 1996

NEPTAZANE

+ LEDERLE

25MG **

N011721 002 Nov 25, 1991

+

50MG **

N011721 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHDILAZINETABLET, CHEWABLE;ORAL
TACARYL

WESTWOOD SQUIBB 3.6MG N011950 009

METHDILAZINE HYDROCHLORIDE

SYRUP;ORAL

METHDILAZINE HYDROCHLORIDE

ALPHARMA US PHARMS 4MG/5ML A087122 001

TACARYL

WESTWOOD SQUIBB 4MG/5ML N011950 007

TABLET;ORAL

TACARYL

WESTWOOD SQUIBB 8MG N011950 006

METHENAMINE HIPPURATE

TABLET;ORAL

METHENAMINE HIPPURATE

IMPAX LABS INC 1GM A076411 001 Jun 20, 2003

METHICILLIN SODIUM

INJECTABLE; INJECTION

STAPHCILLIN

APOTHECON EQ 900MG BASE/VIAL A061449 001

EQ 900MG BASE/VIAL N050117 001

EQ 3.6GM BASE/VIAL A061449 002

EQ 3.6GM BASE/VIAL N050117 002

EQ 5.4GM BASE/VIAL A061449 003

EQ 5.4GM BASE/VIAL N050117 003

METHIMAZOLE

TABLET;ORAL

METHIMAZOLE

DISCOVERY THERAP 15MG A040619 003 Jul 12, 2005

MYLAN 20MG A040350 003 Jun 07, 2001

QINGDAO BAHEAL PHARM 20MG A040547 004 Feb 18, 2005

SUN PHARM INDS INC 5MG A040870 001 Sep 25, 2007

10MG A040870 002 Sep 25, 2007

TAPAZOLE

+ KING PHARMS 5MG ** N007517 002

+ 10MG ** N007517 004

KING PHARMS LLC 5MG A040320 001 Mar 31, 2000

10MG A040320 002 Mar 31, 2000

METHIXENE HYDROCHLORIDE

TABLET;ORAL

TREST

NOVARTIS 1MG N013420 001

METHOCARBAMOL

INJECTABLE; INJECTION

METHOCARBAMOL

DR REDDYS 100MG/ML A086459 001

MARSAM PHARMS LLC 100MG/ML A089849 001 Dec 27, 1991

SOLUTION;IM-IV

METHOCARBAMOL

BAXTER HLTHCARE CORP 1GM/10ML (100MG/ML) A215065 001 Jul 14, 2022

MYLAN INSTITUTIONAL 1GM/10ML (100MG/ML) A204404 001 Dec 05, 2014

TABLET;ORAL

DELAXIN

FERNDAL LABS 500MG A085454 001

FORBAXIN

FOREST LABS 750MG A085136 001

METHOCARBAMOL

ABLE 500MG A040413 001 Mar 17, 2003

750MG A040413 002 Mar 17, 2003

AM THERAP 500MG A089417 001 Feb 11, 1987

750MG A089418 001 Feb 11, 1987

ANI PHARMS 500MG A084277 001

750MG A084276 002

ASCOT 500MG A087660 001 Oct 27, 1982

750MG A087661 001 Oct 27, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

CHARTWELL MOLECULAR	500MG	A084756 002	Mar 31, 2003
	750MG	A084756 001	
CLONMEL HLTHCARE	500MG	A085961 001	
	750MG	A085963 001	
FOSUN PHARMA	500MG	A084616 001	
	750MG	A084615 001	
HEATHER	500MG	A084675 001	
	750MG	A084924 001	
HIKMA INTL PHARMS	500MG	A085159 001	
	750MG	A085123 001	
IMPAX LABS	500MG	A084927 001	
	750MG	A084928 001	
INWOOD LABS	500MG	A085137 001	
IVAX SUB TEVA PHARMS	500MG	A084648 001	
	750MG	A084649 001	
KV PHARM	500MG	A085660 001	
	750MG	A085658 001	
MYLAN	500MG	A084259 001	
	750MG	A084323 001	
NYLOS	750MG	A085033 001	
PIONEER PHARMS	500MG	A088731 001	Dec 13, 1985
	750MG	A089082 001	Dec 13, 1985
PURACAP PHARM	500MG	A084231 002	
	750MG	A084471 001	
PUREPAC PHARM	500MG	A085718 001	
	750MG	A085718 002	
ROXANE	500MG	A088646 001	Feb 29, 1984
	750MG	A088647 001	Feb 29, 1984
SANDOZ	500MG	A087283 001	
	750MG	A087282 001	
SOLVAY	500MG	A084448 001	
	750MG	A084449 001	
SUN PHARM INDUSTRIES	500MG	A084488 001	
	750MG	A084486 001	
SUPERPHARM	500MG	A087589 001	Jan 22, 1982
	750MG	A087590 001	Jan 22, 1982
TABLICAPS	500MG	A084846 001	
UPSHER SMITH	500MG	A087453 001	
	750MG	A087454 001	
WATSON LABS	500MG	A083605 001	
	500MG	A085180 001	
	750MG	A083605 002	
	750MG	A085192 001	
ROBAXIN			
+ AUXILIUM PHARMS LLC	500MG **	N011011 004	
ROBAXIN-750			
+ AUXILIUM PHARMS LLC	750MG **	N011011 006	

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

PAR STERILE PRODUCTS	200MG/VIAL	N011559 004	Dec 21, 2012
+	2.5GM/VIAL	N011559 002	
	5GM/VIAL	N011559 003	

METHOTREXATE

SOLUTION; INTRAVENOUS

METHOTREXATE

+ ACCORD HLTHCARE	5GM/50ML (100MG/ML)	N214121 001	Aug 24, 2020
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SOLUTION; SUBCUTANEOUS

OTREXUP

+ OTTER PHARMS	7.5MG/0.4ML (7.5MG/0.4ML)	N204824 005	Nov 07, 2014
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OTREXUP PFS

+ OTTER PHARMS	10MG/0.4ML (10MG/0.4ML)	N204824 009	May 31, 2017
+	15MG/0.6ML (15MG/0.6ML)	N204824 010	May 31, 2017
+	17.5MG/0.7ML (17.5MG/0.7ML)	N204824 011	May 31, 2017
+	20MG/0.8ML (20MG/0.8ML)	N204824 012	May 31, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP PFS

+		22.5MG/0.9ML (22.5MG/0.9ML)	N204824	013	May 31, 2017
+		25MG/ML (25MG/ML)	N204824	014	May 31, 2017

RASUVO

+	MEDEXUS	27.5MG/0.55ML (27.5MG/0.55ML)	N205776	009	Jul 10, 2014
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REDITREX

+	NORDIC GRP	7.5MG/0.3ML (7.5MG/0.3ML)	N210737	001	Nov 27, 2019
+		10MG/0.4ML (10MG/0.4ML)	N210737	002	Nov 27, 2019
+		12.5MG/0.5ML (12.5MG/0.5ML)	N210737	003	Nov 27, 2019
+		15MG/0.6ML (15MG/0.6ML)	N210737	004	Nov 27, 2019
+		17.5MG/0.7ML (17.5MG/0.7ML)	N210737	005	Nov 27, 2019
+		20MG/0.8ML (20MG/0.8ML)	N210737	006	Nov 27, 2019
+		22.5MG/ML (22.5MG/ML)	N210737	007	Nov 27, 2019
+		25MG/1ML (25MG/1ML)	N210737	008	Nov 27, 2019

METHOTREXATE SODIUM

INJECTABLE; INJECTION

ABITREXATE

ABIC

		EQ 25MG BASE/ML	A089161	001	Mar 10, 1987
		EQ 50MG BASE/VIAL	A089354	001	Jul 17, 1987
		EQ 100MG BASE/VIAL	A089355	001	Jul 17, 1987
		EQ 250MG BASE/VIAL	A089356	001	Jul 17, 1987

FOLEX

+	PHARMACIA AND UPJOHN	EQ 25MG BASE/VIAL	A087695	001	Apr 08, 1983
		EQ 50MG BASE/VIAL	A087695	002	Apr 08, 1983
+		EQ 100MG BASE/VIAL	A087695	003	Apr 08, 1983
		EQ 250MG BASE/VIAL	A088954	001	Oct 24, 1985

FOLEX PFS

	PHARMACIA AND UPJOHN	EQ 25MG BASE/ML	A081242	001	Aug 23, 1991
		EQ 25MG BASE/ML	A089180	001	Jan 03, 1986

METHOTREXATE LPF

+	HOSPIRA	EQ 25MG BASE/ML	N011719	007	Mar 31, 1982
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METHOTREXATE PRESERVATIVE FREE

	FRESENIUS KABI USA	EQ 25MG BASE/ML	A040265	001	Feb 26, 1999
+	HOSPIRA	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N011719	014	Apr 13, 2005
+		EQ 500MG BASE/20ML (EQ 25MG BASE/ML) **	N011719	013	Apr 13, 2005
+		EQ 2.5GM BASE/100ML (EQ 25MG BASE/ML)	N011719	011	Apr 13, 2005

METHOTREXATE SODIUM

ABRAXIS PHARM

		EQ 2.5MG BASE/ML	A089323	001	Jun 13, 1986
		EQ 20MG BASE/VIAL	A088935	001	Oct 11, 1985
		EQ 25MG BASE/ML	A089263	001	Jun 13, 1986
		EQ 25MG BASE/ML	A089322	001	Jun 13, 1986
		EQ 50MG BASE/VIAL	A088936	001	Oct 11, 1985
		EQ 100MG BASE/VIAL	A088937	001	Oct 11, 1985

	FRESENIUS KABI USA	EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	A040263	002	Feb 26, 1999
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+	HOSPIRA	EQ 2.5MG BASE/ML	N011719	004	
+		EQ 20MG BASE/VIAL	N011719	001	
+		EQ 25MG BASE/ML	N011719	005	
+		EQ 50MG BASE/VIAL	N011719	003	
+		EQ 100MG BASE/VIAL	N011719	006	
	NORBROOK	EQ 25MG BASE/ML	A088648	001	May 09, 1986
	PHARMACHEMIE USA	EQ 25MG BASE/ML	A089158	001	Jul 08, 1988

METHOTREXATE SODIUM PRESERVATIVE FREE

	EUGIA PHARMA SPECLTS	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	A201529	001	Mar 29, 2012
		EQ 100MG BASE/4ML (EQ 25MG BASE/ML)	A201529	002	Mar 29, 2012
		EQ 200MG BASE/8ML (EQ 25MG BASE/ML)	A201529	003	Mar 29, 2012
		EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	A201529	004	Mar 29, 2012
+	HOSPIRA	EQ 1GM BASE/VIAL	N011719	009	Apr 07, 1988
	MYLAN	EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	A201530	001	Mar 29, 2012

MEXATE

+	BRISTOL	EQ 20MG BASE/VIAL	A086358	001	
+		EQ 50MG BASE/VIAL	A086358	002	
+		EQ 100MG BASE/VIAL	A086358	003	
+		EQ 250MG BASE/VIAL	A086358	004	

MEXATE-AQ

+	BRISTOL MYERS	EQ 25MG BASE/ML	A088760	001	Feb 14, 1985
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOTREXATE SODIUM

INJECTABLE; INJECTION

MEXATE-AQ PRESERVED

BRISTOL MYERS SQUIBB EQ 25MG BASE/ML

A089887 001 Apr 14, 1989

TABLET; ORAL

METHOTREXATE SODIUM

DURAMED PHARMS BARR EQ 2.5MG BASE

A040233 001 Jun 17, 1999

+ STRIDES PHARMA

EQ 2.5MG BASE **

N008085 002

METHOXAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

VASOXYL

GLAXOSMITHKLINE 10MG/ML

N006772 002

20MG/ML

N006772 001

METHOXSALLEN

CAPSULE; ORAL

8-MOP

+ VALEANT PHARM INTL 10MG

N009048 001

METHOXSALLEN

ACTAVIS INC 10MG

A202603 001 Jun 09, 2015

ANI PHARMS 10MG

A087781 001 Jun 08, 1982

LOTION; TOPICAL

OXSORALEN

+ VALEANT PHARM INTL 1%

N009048 002

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

CHARTWELL RX 2.5MG

A040624 001 Dec 28, 2006

5MG

A040624 002 Dec 28, 2006

PVT FORM 2.5MG

A080970 001

PAMINE

FOUGERA PHARMS 2.5MG **

N008848 001

PAMINE FORTE

FOUGERA PHARMS 5MG **

N008848 002 Mar 25, 2003

METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

+ PARKE DAVIS 150MG

N010596 007

METHYCLOTHIAZIDE

TABLET; ORAL

AQUATENSEN

MEDPOINTE PHARM HLC 5MG

N017364 001

ENDURON

+ ABBVIE 2.5MG **

N012524 001

+ 5MG **

N012524 004

METHYCLOTHIAZIDE

CHARTWELL RX 2.5MG

A089835 001 Aug 18, 1988

5MG

A089837 001 Aug 18, 1988

IVAX PHARMS 2.5MG

A087913 001 Jun 03, 1982

5MG

A087786 001 May 18, 1982

MYLAN 2.5MG

A087671 001 Aug 17, 1982

MYLAN PHARMS INC 5MG

A087672 001 Aug 17, 1982

PAR PHARM 2.5MG

A089135 001 Feb 12, 1986

5MG

A089136 001 Feb 12, 1986

USL PHARMA 5MG

A088745 001 Mar 21, 1985

WATSON LABS 2.5MG

A085487 001 Mar 11, 1982

2.5MG

A088750 001 Sep 06, 1984

5MG

A085476 001 Mar 11, 1982

5MG

A088724 001 Sep 06, 1984

METHYCLOTHIAZIDE; PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTRON

ABBOTT 5MG; 25MG

N016047 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYCLOTHIAZIDE; RESERPINE

TABLET; ORAL

DIUTENSEN-R

MEDPOINTE PHARM HLC 2.5MG;0.1MG

N012708 005

METHYL AMINOLEVULINATE HYDROCHLORIDE

CREAM; TOPICAL

METVIXIA

GALDERMA LABS LP EQ 16.8% BASE

N021415 001 Jul 27, 2004

METHYLDOPA

SUSPENSION; ORAL

ALDOMET

MERCK

250MG/5ML

N018389 001

TABLET; ORAL

ALDOMET

+ MERCK

125MG **

N013400 003

+

250MG **

N013400 001

+

500MG **

N013400 002

METHYLDOPA

ACCORD HLTHCARE

125MG

A070070 003 Oct 15, 1985

250MG

A070084 001 Oct 15, 1985

500MG

A070085 001 Oct 15, 1985

CHARTWELL RX

125MG

A071700 001 Mar 02, 1988

250MG

N018934 001 Jun 29, 1984

500MG

N018934 002 Jun 29, 1984

DURAMED PHARMS BARR

250MG

A071006 001 Dec 16, 1986

500MG

A071009 001 Dec 16, 1986

HALSEY

125MG

A071751 001 Mar 28, 1988

250MG

A071752 001 Mar 28, 1988

500MG

A071753 001 Mar 28, 1988

HERITAGE PHARMA

250MG

A070098 001 Feb 20, 1986

500MG

A070343 001 Feb 20, 1986

PARKE DAVIS

125MG

A070331 001 Apr 15, 1986

250MG

A070332 001 Apr 15, 1986

500MG

A070333 001 Apr 15, 1986

PLIVA

125MG

A072126 001 Jul 07, 1988

250MG

A072127 001 Jul 07, 1988

500MG

A072128 001 Jul 07, 1988

PUREPAC PHARM

125MG

A070749 001 Feb 07, 1986

250MG

A070750 001 Feb 07, 1986

500MG

A070452 001 Feb 07, 1986

RISING

250MG

A070076 002 Apr 18, 1985

500MG

A070076 001 Apr 18, 1985

ROXANE

125MG

A070192 001 Apr 25, 1986

250MG

A070193 001 Apr 25, 1986

500MG

A070194 001 Apr 25, 1986

STRIDES PHARMA

125MG

A070535 001 Jan 02, 1987

250MG

A070536 001 Jan 02, 1987

500MG

A070537 001 Jan 02, 1987

SUN PHARM INDUSTRIES

125MG

A070073 001 Oct 09, 1986

250MG

A070060 001 Oct 09, 1986

500MG

A070074 001 Oct 09, 1986

SUPERPHARM

250MG

A070669 001 Jun 23, 1989

500MG

A070670 001 Jun 23, 1989

TEVA

125MG

A071105 001 Dec 05, 1986

250MG

A071106 001 Dec 05, 1986

500MG

A071067 001 Dec 05, 1986

WATSON LABS

125MG

A070245 001 Feb 25, 1986

125MG

A070260 001 Jun 24, 1985

250MG

A070246 001 Feb 25, 1986

250MG

A070261 001 Jun 24, 1985

250MG

A070703 001 Jun 06, 1986

500MG

A070247 001 Feb 25, 1986

500MG

A070262 001 Jun 24, 1985

500MG

A070625 001 Jun 06, 1986

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

ALDOMET

+ MERCK 50MG/ML **

N013401 001

METHYLDOPATE HYDROCHLORIDE

ABRAXIS PHARM 50MG/ML

A070652 001 Jun 03, 1986

AM REGENT 50MG/ML

A071279 001 Oct 02, 1987

BAXTER HLTHCARE 50MG/ML

A070291 001 Jul 01, 1986

HOSPIRA 50MG/ML

A070691 001 Jun 19, 1987

50MG/ML

A070698 001 Jun 15, 1987

50MG/ML

A070699 001 Jun 15, 1987

50MG/ML

A070849 001 Jun 19, 1987

MARSAM PHARMS LLC 50MG/ML

A071812 001 Dec 22, 1987

SMITH AND NEPHEW 50MG/ML

A070841 001 Jan 02, 1987

TEVA PARENTERAL 50MG/ML

A072974 001 Nov 22, 1991

METHYLERGONOVINE MALEATE

TABLET; ORAL

METHERGINE

+ EDISON THERAPS LLC 0.2MG **

N006035 003

METHYLPHENIDATE

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE; ORAL

METHYLPHENIDATE

ACTAVIS ELIZABETH 8.6MG

A210924 001 Jun 19, 2020

17.3MG

A210924 002 Jun 19, 2020

25.9MG

A210924 003 Jun 19, 2020

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

ADHANSIA XR

+ PURDUE PHARMA LP 25MG

N212038 001 Feb 27, 2019

+ 35MG

N212038 002 Feb 27, 2019

+ 45MG

N212038 003 Feb 27, 2019

+ 55MG

N212038 004 Feb 27, 2019

+ 70MG

N212038 005 Feb 27, 2019

+ 85MG

N212038 006 Feb 27, 2019

METHYLPHENIDATE HYDROCHLORIDE

BARR LABS INC 10MG

A079031 004 Oct 15, 2014

20MG

A079031 001 Jul 13, 2012

30MG

A079031 002 Jul 13, 2012

40MG

A079031 003 Jul 13, 2012

RITALIN LA

+ SANDOZ 60MG **

N021284 005 Oct 27, 2014

SOLUTION; ORAL

METHYLPHENIDATE HYDROCHLORIDE

CHARTWELL MOLECULAR 5MG/5ML

A207414 001 Dec 16, 2020

10MG/5ML

A207414 002 Dec 16, 2020

TABLET; ORAL

METHYLPHENIDATE HYDROCHLORIDE

ABLE 5MG

A040404 001 Mar 29, 2001

10MG

A040404 002 Mar 29, 2001

20MG

A040404 003 Mar 29, 2001

ACTAVIS ELIZABETH 5MG

A040321 001 Feb 05, 2002

10MG

A040321 002 Feb 05, 2002

20MG

A040321 003 Feb 05, 2002

ALVOGEN 5MG

A206840 001 Sep 15, 2016

10MG

A206840 002 Sep 15, 2016

20MG

A206840 003 Sep 15, 2016

AUROLIFE PHARMA LLC 5MG

A209276 001 Oct 25, 2018

10MG

A209276 002 Oct 25, 2018

20MG

A209276 003 Oct 25, 2018

CEDIPROF INC 5MG

A208737 001 Feb 01, 2019

10MG

A208737 002 Feb 01, 2019

20MG

A208737 003 Feb 01, 2019

LANNETT CO INC 5MG

A086429 001

10MG

A085799 001

20MG

A086428 001

NOSTRUM LABS INC 5MG

A207587 001 Mar 03, 2017

10MG

A207587 002 Mar 03, 2017

20MG

A207587 003 Mar 03, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPHENIDATE HYDROCHLORIDE

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

WATSON LABS	5MG	A040220 001	Aug 29, 1997
	10MG	A040220 002	Aug 29, 1997
	20MG	A040220 003	Aug 29, 1997

TABLET, CHEWABLE;ORAL

METHYLIN

+ SPECGX LLC	2.5MG **	N021475 001	Apr 15, 2003
+	5MG **	N021475 002	Apr 15, 2003
+	10MG **	N021475 003	Apr 15, 2003

METHYLPHENIDATE HYDROCHLORIDE

NOSTRUM LABS INC	2.5MG	A204954 001	Jan 26, 2017
	5MG	A204954 002	Jan 26, 2017
	10MG	A204954 003	Jan 26, 2017
NOVEL LABS INC	2.5MG	A204115 001	Feb 25, 2015
	5MG	A204115 002	Feb 25, 2015
	10MG	A204115 003	Feb 25, 2015

TABLET, EXTENDED RELEASE;ORAL

METADATE ER

LANNETT CO INC	10MG	A040306 001	Oct 20, 1999
	20MG	A089601 001	Jun 01, 1988

METHYLPHENIDATE HYDROCHLORIDE

ABLE	20MG	A076032 001	May 09, 2001
ALVOGEN	10MG	A204772 001	Feb 29, 2016
	18MG	A210818 001	Nov 30, 2018
	20MG	A204772 002	Feb 29, 2016
	27MG	A210818 002	Nov 30, 2018
	36MG	A210818 003	Nov 30, 2018
	54MG	A210818 004	Nov 30, 2018
AMNEAL PHARMS	18MG	A207515 001	Feb 01, 2018
	27MG	A207515 002	Feb 01, 2018
	36MG	A207515 003	Feb 01, 2018
	54MG	A207515 004	Feb 01, 2018
ANI PHARMS	18MG	A208607 001	Jul 14, 2017
	27MG	A208607 002	Jul 14, 2017
	36MG	A208607 003	Jul 14, 2017
	54MG	A208607 004	Jul 14, 2017
HERITAGE PHARMA	20MG	A075450 001	Dec 21, 2001
RHODES PHARMS	18MG	A214111 001	May 31, 2022
	27MG	A214111 002	May 31, 2022
	36MG	A214111 003	May 31, 2022
	54MG	A214111 004	May 31, 2022
STRIDES PHARMA	36MG	A204659 001	Jul 15, 2019
	54MG	A204659 002	Jul 15, 2019
WATSON LABS	20MG	A040410 001	Feb 09, 2001

RITALIN-SR

+ NOVARTIS	20MG **	N018029 001	Mar 30, 1982
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METHYLPREDNISOLONE

TABLET;ORAL

MEDROL

PFIZER	24MG	N011153 005	
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METHYLPREDNISOLONE

AMNEAL	4MG	A207481 001	Sep 21, 2021
	8MG	A207481 002	Sep 21, 2021
	16MG	A207481 003	Sep 21, 2021
	32MG	A207481 004	Sep 21, 2021
CHARTWELL RX	2MG	A209097 001	Feb 22, 2019
	4MG	A209097 002	Feb 22, 2019
	8MG	A209097 003	Feb 22, 2019
	16MG	A209097 004	Feb 22, 2019
	32MG	A209097 005	Feb 22, 2019
DURAMED PHARMS BARR	4MG	A088497 001	Feb 21, 1984
HEATHER	4MG	A085650 001	
INVATECH	4MG	A087341 001	
NOVAST LABS	4MG	A210985 001	Jan 09, 2019
PAR PHARM	16MG	A089207 001	Apr 25, 1988
	24MG	A089208 001	Apr 25, 1988
	32MG	A089209 001	Apr 25, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPREDNISOLONE

TABLET; ORAL

METHYLPREDNISOLONE

WATSON LABS

4MG

A086161 001 Feb 09, 1982

16MG

A086159 001 Feb 09, 1982

METHYLPREDNISOLONE ACETATE

ENEMA; RECTAL

MEDROL

PHARMACIA AND UPJOHN

40MG/BOT

N018102 001

INJECTABLE; INJECTION

M-PREDROL

BEL MAR

40MG/ML

A086666 001

80MG/ML

A087135 001

METHYLPREDNISOLONE ACETATE

AKORN

40MG/ML

A086903 001 Oct 20, 1982

80MG/ML

A086903 002 Oct 20, 1982

SAGENT PHARMS INC

20MG/ML

A201835 001 Jun 27, 2018

WATSON LABS

20MG/ML

A085597 001

20MG/ML

A087248 001

40MG/ML

A085374 001

40MG/ML

A085600 001

80MG/ML

A085595 001

80MG/ML

A086507 001

OINTMENT; TOPICAL

MEDROL ACETATE

PHARMACIA AND UPJOHN

0.25%

N012421 001

1%

N012421 002

METHYLPREDNISOLONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-MEDROL ACETATE

PHARMACIA AND UPJOHN

0.25%;EQ 3.5MG BASE/GM

A060611 002

1%;EQ 3.5MG BASE/GM

A060611 001

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

ABBOTT

EQ 40MG BASE/VIAL

A089573 001 Feb 22, 1991

EQ 125MG BASE/VIAL

A089574 001 Feb 22, 1991

EQ 500MG BASE/VIAL

A089575 001 Feb 22, 1991

EQ 1GM BASE/VIAL

A089576 001 Feb 22, 1991

HOSPIRA

EQ 40MG BASE/VIAL

A040664 001 Dec 20, 2005

EQ 40MG BASE/VIAL

A085853 001

EQ 125MG BASE/VIAL

A040665 001 Dec 20, 2005

EQ 125MG BASE/VIAL

A085855 001

EQ 500MG BASE/VIAL

A085854 001

EQ 500MG BASE/VIAL

A089173 001 Aug 18, 1987

EQ 1GM BASE/VIAL

A085852 001

EQ 1GM BASE/VIAL

A089174 001 Aug 18, 1987

HOSPIRA INC

EQ 40MG BASE/VIAL

A040793 001 Nov 25, 2008

EQ 125MG BASE/VIAL

A040827 001 Nov 25, 2008

METHYLPREDNISOLONE

ELKINS SINN

EQ 125MG BASE/VIAL

A086906 002

EQ 500MG BASE/VIAL

A086906 003

EQ 1GM BASE/VIAL

A086906 004

ORGANON USA INC

EQ 500MG BASE/VIAL

A087535 001 Jun 25, 1982

EQ 1GM BASE/VIAL

A087535 002 Jun 25, 1982

METHYLPREDNISOLONE SODIUM SUCCINATE

ABRAXIS PHARM

EQ 40MG BASE/VIAL

A088676 001 Jun 08, 1984

EQ 40MG BASE/VIAL

A089143 001 Mar 28, 1986

EQ 125MG BASE/VIAL

A088677 001 Jun 08, 1984

EQ 125MG BASE/VIAL

A089144 001 Mar 28, 1986

EQ 500MG BASE/VIAL

A088678 001 Jun 08, 1984

EQ 500MG BASE/VIAL

A089186 001 Mar 28, 1986

EQ 500MG BASE/VIAL

A089187 001 Mar 28, 1986

EQ 1GM BASE/VIAL

A088679 001 Jun 08, 1984

EQ 1GM BASE/VIAL

A089188 001 Mar 28, 1986

EQ 1GM BASE/VIAL

A089189 001 Mar 28, 1986

BEDFORD LABS

EQ 40MG BASE/VIAL

A040662 001 Feb 21, 2007

EQ 125MG BASE/VIAL

A040641 002 Feb 21, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

	EQ 500MG BASE/VIAL	A040641 003	Feb 21, 2007
	EQ 500MG BASE/VIAL	A040709 001	Feb 21, 2007
	EQ 1GM BASE/VIAL	A040641 004	Feb 21, 2007
	EQ 1GM BASE/VIAL	A040709 002	Feb 21, 2007
ELKINS SINN	EQ 40MG BASE/VIAL	A086906 001	
EUGIA PHARMA	EQ 40MG BASE/VIAL	A207667 001	Dec 15, 2015
	EQ 125MG BASE/VIAL	A207667 002	Dec 15, 2015
	EQ 500MG BASE/VIAL	A207667 003	Dec 15, 2015
	EQ 2GM BASE/VIAL	A207667 004	Dec 15, 2015
INTL MEDICATION	EQ 40MG BASE/VIAL	A087812 001	Feb 09, 1983
	EQ 125MG BASE/VIAL	A087813 001	Feb 09, 1983
	EQ 500MG BASE/VIAL	A087851 001	Feb 09, 1983
	EQ 1GM BASE/VIAL	A087852 001	Feb 09, 1983
TEVA PARENTERAL	EQ 125MG BASE/VIAL	A081266 001	Nov 30, 1992
	EQ 500MG BASE/VIAL	A081267 001	Nov 30, 1992
	EQ 1GM BASE/VIAL	A081268 001	Nov 30, 1992
WATSON LABS	EQ 40MG BASE/VIAL	A086953 001	Jul 22, 1982
	EQ 125MG BASE/VIAL	A087030 001	Jul 22, 1982
	EQ 500MG BASE/VIAL	A088523 001	Jul 24, 1984
	EQ 1GM BASE/VIAL	A088524 001	Jul 24, 1984

METHYLPREDNISOLONE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC

NEO-MEDROL

PHARMACIA AND UPJOHN	0.1%;EQ 3.5MG BASE/GM	A060645 001	
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METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

HEATHER	10MG	A084967 001	
TESTRED			
+ BAUSCH	10MG	A083976 001	
VIRILON			
CHARTWELL	10MG	A087750 001	Nov 24, 1982
TABLET; BUCCAL			
ANDROID 5			
VALEANT PHARM INTL	5MG	A087222 001	
ORETON			
SCHERING	10MG	A080281 001	
TABLET; BUCCAL, SUBLINGUAL			
METANDREN			
NOVARTIS	5MG	N003240 004	
	10MG	N003240 001	
	10MG	N003240 005	
	25MG	N003240 003	
METHYLTESTOSTERONE			
IMPAX LABS	10MG	A084287 001	
LILLY	10MG	A080256 001	
	25MG	A080256 002	
PUREPAC PHARM	10MG	A080308 001	
	10MG	A080475 001	
	10MG	A080475 002	
	25MG	A080475 003	
PVT FORM	5MG	A083836 001	
TABLICAPS	10MG	A085125 001	
USL PHARMA	10MG	A080271 001	
TABLET; ORAL			
ANDROID 10			
VALEANT PHARMS NORTH	10MG	A086450 001	
METHYLTESTOSTERONE			
IMPAX LABS	25MG	A084310 001	
INWOOD LABS	10MG	A080839 001	
	25MG	A080973 001	
KV PHARM	10MG	A084312 001	
LANNETT	10MG	A087092 001	Nov 05, 1982
	25MG	A087111 001	Jan 27, 1983
PARKE DAVIS	10MG	A084244 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLTESTOSTERONE

TABLET; ORAL

METHYLTESTOSTERONE

	25MG	A084241 001
PUREPAC PHARM	10MG	A080309 001
	25MG	A080310 001
PVT FORM	5MG	A080214 001
	10MG	A080214 002
	25MG	A080214 003
TABLICAPS	10MG	A080313 001
	25MG	A085270 001
WATSON LABS	10MG	A080933 001
	25MG	A080931 001
WEST WARD	10MG	A084331 001
	25MG	A084331 002
	25MG	A084642 001
ORETON METHYL		
SCHERING	10MG	N003158 001
	25MG	N003158 002

METHYPRYLON

CAPSULE; ORAL

NOLUDAR

ROCHE	300MG	N009660 008
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ELIXIR; ORAL

NOLUDAR

ROCHE	50MG/5ML	N009660 007
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TABLET; ORAL

NOLUDAR

ROCHE	50MG	N009660 002
	200MG	N009660 004

METHYSERGIDE MALEATE

TABLET; ORAL

SANSERT

NOVARTIS	2MG	N012516 001
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METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

METIPRANOLOL

SANDOZ	0.3%	A075720 001	Aug 06, 2001
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OPTIPRANOLOL

+ BAUSCH AND LOMB	0.3% **	N019907 001	Dec 29, 1989
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METOCLOPRAMIDE HYDROCHLORIDE

CONCENTRATE; ORAL

METOCLOPRAMIDE INTENSOL

ROXANE	EQ 10MG BASE/ML	A072995 001	Jan 30, 1992
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INJECTABLE; INJECTION

METOCLOPRAMIDE HYDROCHLORIDE

BEDFORD	EQ 5MG BASE/ML	A072155 001	Mar 30, 1992
	EQ 5MG BASE/ML	A072244 001	Mar 30, 1992
	EQ 5MG BASE/ML	A072247 001	May 18, 1992
HOSPIRA	EQ 5MG BASE/ML	A070505 001	Jun 23, 1989
	EQ 5MG BASE/ML	A070506 001	Jun 22, 1989
	EQ 5MG BASE/ML	A070847 001	Nov 07, 1988
	EQ 5MG BASE/ML	A071291 001	Mar 03, 1989
	EQ 5MG BASE/ML	A071990 001	Jan 18, 1989
	EQ 5MG BASE/ML	A073117 001	Jan 17, 1991
	EQ 5MG BASE/ML	A074147 001	Aug 02, 1996
LYPHOMED	EQ 10MG BASE/2ML	A070293 001	Jan 24, 1986
NORBROOK	EQ 10MG BASE/2ML	A070892 001	Aug 26, 1988
SMITH AND NEPHEW	EQ 5MG BASE/ML	A070623 001	Mar 02, 1987
	EQ 10MG BASE/2ML	A070622 001	Mar 02, 1987

REGLAN

+ HIKMA	EQ 5MG BASE/ML **	N017862 001
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	EQ 10MG BASE/ML **	N017862 004	May 28, 1987
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SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 5MG BASE/5ML	A071340 001	Aug 18, 1988
CHARTWELL MOLECULAR	EQ 5MG BASE/5ML	A073680 001	Oct 27, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOCLOPRAMIDE HYDROCHLORIDE

SOLUTION;ORAL

METOCLOPRAMIDE HYDROCHLORIDE

MORTON GROVE	EQ 5MG BASE/5ML	A070949 001	Mar 06, 1987
PACO	EQ 5MG BASE/5ML	A071665 001	Dec 05, 1988
PHARMOBEDIENT CNSLTG	EQ 5MG BASE/5ML	A074703 001	Oct 31, 1997
ROXANE	EQ 5MG BASE/5ML	A072038 001	Dec 05, 1988
TEVA	EQ 5MG BASE/5ML	A070819 001	Jul 10, 1987
	EQ 5MG BASE/5ML	A071315 001	Jun 30, 1993
VISTAPHARM	EQ 5MG BASE/5ML	A075051 001	Jan 26, 2001

REGLAN

+ ROBINS AH	EQ 5MG BASE/5ML **	N018821 001	Mar 25, 1983
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TABLET;ORAL

CLOPRA

QUANTUM PHARMICS	EQ 5MG BASE	A072384 001	Jun 02, 1988
	EQ 10MG BASE	A070294 001	Jul 29, 1985

CLOPRA-"YELLOW"

QUANTUM PHARMICS	EQ 10MG BASE	A070632 001	Oct 28, 1985
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MAXOLON

KING PHARMS	EQ 10MG BASE	A070106 001	Mar 04, 1986
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METOCLOPRAMIDE HYDROCHLORIDE

AIPING PHARM INC	EQ 10MG BASE	A072215 001	Jan 30, 1990
CHARTWELL RX	EQ 5MG BASE	A074478 001	Oct 05, 1995
	EQ 10MG BASE	A074478 002	Oct 05, 1995
CLONMEL	EQ 10MG BASE	A072639 001	May 09, 1991
HALSEY	EQ 10MG BASE	A070906 001	Oct 28, 1986
INTERPHARM	EQ 10MG BASE	A071213 001	Sep 24, 1986
MUTUAL PHARM	EQ 10MG BASE	A070660 001	Feb 10, 1987
NORTHSTAR HLTHCARE	EQ 5MG BASE	A078374 001	Nov 30, 2007
	EQ 10MG BASE	A078374 002	Nov 30, 2007
PAR PHARM	EQ 10MG BASE	A070342 001	Mar 25, 1986
SANDOZ	EQ 5MG BASE	A072436 001	Jun 22, 1989
	EQ 10MG BASE	A070850 001	Feb 03, 1987
SCHERING	EQ 10MG BASE	A070598 001	Feb 02, 1987
SUN PHARM INDUSTRIES	EQ 5MG BASE	A071536 002	Jan 16, 1997
	EQ 10MG BASE	A071536 001	Apr 28, 1993
SUPERPHARM	EQ 10MG BASE	A070926 001	Jun 26, 1987
USL PHARMA	EQ 10MG BASE	A070339 001	Jul 29, 1985
WATSON LABS	EQ 10MG BASE	A070363 001	Mar 02, 1987
	EQ 10MG BASE	A070453 001	Jun 06, 1986
	EQ 10MG BASE	A070511 001	Jan 22, 1986
	EQ 10MG BASE	A070645 001	May 11, 1987

TABLET, ORALLY DISINTEGRATING;ORAL

METOZOLV ODT

+ SALIX PHARMS	EQ 5MG BASE **	N022246 001	Sep 04, 2009
+	EQ 10MG BASE **	N022246 002	Sep 04, 2009

REGLAN ODT

MEDA PHARMS	EQ 5MG BASE	N021793 001	Jun 10, 2005
	EQ 10MG BASE	N021793 002	Jun 10, 2005

METOCURINE IODIDE

INJECTABLE;INJECTION

METUBINE IODIDE

LILLY	2MG/ML	N006632 003	
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METOLAZONE

TABLET;ORAL

DIULO

GD SEARLE LLC	2.5MG	N018535 001	
	5MG	N018535 002	
	10MG	N018535 003	

METOLAZONE

ANI PHARMS	2.5MG	A075543 001	Jan 06, 2004
	5MG	A075543 002	Mar 01, 2004
	10MG	A075543 003	Dec 24, 2003
ROXANE	10MG	A076482 002	Apr 29, 2004
WATSON LABS	10MG	A076891 001	Jul 21, 2004

MYKROX

CHARTWELL MOLECULAR	0.5MG	N019532 001	Oct 30, 1987
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOLAZONE

TABLET; ORAL

ZAROXOLYN

+	I3 PHARMS	2.5MG **	N017386 001
+		5MG **	N017386 002
+		10MG **	N017386 003

METOPROLOL FUMARATE

TABLET, EXTENDED RELEASE; ORAL

LOPRESSOR

NOVARTIS	EQ 100MG TARTRATE	N019786 001	Dec 27, 1989
	EQ 200MG TARTRATE	N019786 002	Dec 27, 1989
	EQ 300MG TARTRATE	N019786 003	Dec 27, 1989
	EQ 400MG TARTRATE	N019786 004	Dec 27, 1989

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

ACCORD HLTHCARE	EQ 25MG TARTRATE	A215637 001	Oct 18, 2022
	EQ 50MG TARTRATE	A215637 002	Oct 18, 2022
	EQ 100MG TARTRATE	A215637 003	Oct 18, 2022
	EQ 200MG TARTRATE	A215637 004	Oct 18, 2022
ACTAVIS LABS FL INC	EQ 25MG TARTRATE	A076862 002	Aug 03, 2009
	EQ 100MG TARTRATE	A077298 001	Apr 15, 2010
	EQ 200MG TARTRATE	A077298 002	Apr 15, 2010
LUPIN	EQ 25MG TARTRATE	A209272 001	Aug 15, 2023
	EQ 50MG TARTRATE	A209272 002	Aug 15, 2023
	EQ 100MG TARTRATE	A209272 003	Aug 15, 2023
	EQ 200MG TARTRATE	A209272 004	Aug 15, 2023
NESHER PHARMS	EQ 25MG TARTRATE	A077779 001	Mar 20, 2008
	EQ 50MG TARTRATE	A077176 001	May 14, 2008
	EQ 100MG TARTRATE	A076640 002	May 18, 2007
	EQ 200MG TARTRATE	A076640 001	May 18, 2007
PRINSTON INC	EQ 100MG TARTRATE	A210597 001	Jan 04, 2022
	EQ 200MG TARTRATE	A210597 002	Jan 04, 2022
SANDOZ	EQ 25MG TARTRATE	A076969 001	Jul 31, 2006
	EQ 50MG TARTRATE	A076969 002	May 18, 2007
	EQ 100MG TARTRATE	A076969 003	Mar 20, 2008
	EQ 200MG TARTRATE	A076969 004	Mar 20, 2008

METOPROLOL TARTRATE

INJECTABLE; INJECTION

LOPRESSOR

+	NOVARTIS	1MG/ML **	N018704 001	Mar 30, 1984
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METOPROLOL TARTRATE

AM REGENT	1MG/ML	A090386 001	Sep 30, 2009
HOSPIRA	1MG/ML	A075160 001	Jul 06, 1998
LUITPOLD	1MG/ML	A091307 001	Dec 29, 2010
STERISCIENCE	1MG/ML	A090317 001	Apr 19, 2010
WATSON LABS	1MG/ML	A074032 001	Dec 21, 1993

TABLET; ORAL

METOPROLOL TARTRATE

APOTHECON	50MG	A074258 001	Jan 27, 1994
	100MG	A074258 002	Jan 27, 1994
CHARTWELL RX	50MG	A073288 001	Mar 25, 1994
	100MG	A073289 001	Mar 25, 1994
HERITAGE PHARMA	50MG	A074141 001	Jan 31, 1995
	100MG	A074141 002	Jan 31, 1995
MYLAN	50MG	A073666 001	Dec 21, 1993
	100MG	A073666 002	Dec 21, 1993
PUREPAC PHARM	50MG	A074380 001	Jul 29, 1994
	100MG	A074380 002	Jul 29, 1994
SUN PHARM INDUSTRIES	25MG	A073654 002	Jul 15, 2009
	50MG	A073654 003	Dec 21, 1993
	100MG	A073654 001	Dec 21, 1993
TEVA	50MG	A074143 001	Sep 30, 1994
	100MG	A074143 002	Sep 30, 1994
TEVA PHARMS	50MG	A074333 001	Jan 27, 1994
	100MG	A074333 002	Jan 27, 1994
WATSON LABS	50MG	A074217 001	May 27, 1994
	100MG	A074217 002	May 27, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE

ZYDUS PHARMS

25MG

A212402 001 Apr 19, 2023

50MG

A212402 002 Apr 19, 2023

100MG

A212402 003 Apr 19, 2023

METRIZAMIDE

INJECTABLE; INJECTION

AMIPAQUE

GE HEALTHCARE

2.5GM/VIAL

N017982 003 Sep 12, 1983

3.75GM/VIAL

N017982 001

6.75GM/VIAL

N017982 002

13.5GM/VIAL

N017982 004 Sep 12, 1983

METRONIDAZOLE

CAPSULE; ORAL

METRONIDAZOLE

ABLE

375MG

A076505 001 Nov 13, 2003

GEL; TOPICAL

METRONIDAZOLE

CHARTWELL RX

1%

A090903 001 Jul 22, 2011

GEL; VAGINAL

METRONIDAZOLE

ENCUBE

0.75%

A077264 001 Oct 31, 2006

INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

PFIZER

500MG/100ML

N018353 002

METRO I.V.

B BRAUN

500MG/100ML

N018674 001 Aug 31, 1982

METRONIDAZOLE

ABBOTT

500MG/100ML

N018889 001 Nov 18, 1983

ABRAXIS PHARM

500MG/100ML

A070071 001 Dec 03, 1984

HIKMA

500MG/100ML

N018907 001 Mar 30, 1984

INTL MEDICATION

500MG/100ML

A070004 001 May 08, 1985

WATSON LABS

500MG/100ML

A070042 001 Dec 20, 1984

500MG/100ML

A070170 001 Apr 01, 1986

METRONIDAZOLE IN PLASTIC CONTAINER

RISING

500MG/100ML

A205531 001 May 09, 2017

TABLET; ORAL

FLAGYL

+ PFIZER

250MG

N012623 001

+

500MG

N012623 003

METROMIDOL

LABS AF

250MG

A074523 001 Oct 24, 1996

500MG

A074523 002 Oct 24, 1996

METRONIDAZOLE

ABLE

250MG

A076519 001 Jun 27, 2003

500MG

A076519 002 Jun 27, 2003

CHARTWELL MOLECULES

250MG

N018845 001 Aug 18, 1983

500MG

N018930 001 Aug 18, 1983

FOSUN PHARMA

250MG

N018620 001 Mar 04, 1982

250MG

N018740 001 Oct 22, 1982

500MG

N018620 002 Jun 02, 1983

500MG

N018740 002 Oct 22, 1982

HALSEY

250MG

A070021 001 Apr 02, 1985

500MG

A070593 001 Feb 27, 1986

IVAX SUB TEVA PHARMS

250MG

N018517 001

500MG

N018517 002 May 05, 1982

LNK

250MG

N019029 001 Apr 10, 1984

MUTUAL PHARM

250MG

N018818 001 Feb 16, 1983

500MG

N018818 002 Feb 16, 1983

SUPERPHARM

250MG

A070008 001 Dec 11, 1984

500MG

A070009 001 Dec 11, 1984

WATSON LABS

250MG

N018599 001 Sep 17, 1982

250MG

N018764 001 Sep 17, 1982

500MG

N018599 002 Feb 13, 1984

500MG

N018764 002 Dec 20, 1982

PROTOSTAT

ORTHO MCNEIL PHARM

250MG

N018871 001 Mar 02, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METRONIDAZOLETABLET; ORAL
PROTOSTAT

	500MG	N018871 002	Mar 02, 1983
SATRIC			
SAVAGE LABS	250MG	A070029 001	Mar 19, 1985
	500MG	A070731 001	Jun 08, 1987

TABLET, EXTENDED RELEASE; ORAL

FLAGYL ER

+ PFIZER	750MG	N020868 001	Nov 26, 1997
METRONIDAZOLE			
ABLE	750MG	A076462 001	Jun 25, 2003
ALEMBIC	750MG	A090222 001	May 05, 2010

METRONIDAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

FLAGYL I.V.

PFIZER	EQ 500MG BASE/VIAL **	N018353 001	
METRONIDAZOLE HYDROCHLORIDE			
ABRAXIS PHARM	EQ 500MG BASE/VIAL	A070295 001	Oct 15, 1985

METYRAPONE

TABLET; ORAL

METOPIRONE

HRA PHARMA	250MG	N012911 001	
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MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

WATSON LABS	150MG	A074711 001	Feb 26, 1997
	150MG	A074865 001	Apr 13, 1998
	200MG	A074711 002	Feb 26, 1997
	200MG	A074865 002	Apr 13, 1998
	250MG	A074711 003	Feb 26, 1997
	250MG	A074865 003	Apr 13, 1998

MEXITIL

+ BOEHRINGER INGELHEIM	150MG	N018873 002	Dec 30, 1985
	200MG	N018873 003	Dec 30, 1985
	250MG	N018873 004	Dec 30, 1985

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION

MEZLIN

BAYER PHARMS	EQ 1GM BASE/VIAL	A062333 001	
	EQ 1GM BASE/VIAL	A062372 005	Jan 13, 1983
	EQ 1GM BASE/VIAL	N050549 001	
	EQ 2GM BASE/VIAL	A062333 002	
	EQ 2GM BASE/VIAL	A062372 001	May 13, 1982
	EQ 2GM BASE/VIAL	N050549 002	
	EQ 3GM BASE/VIAL	A062333 003	
	EQ 3GM BASE/VIAL	A062372 002	May 13, 1982
	EQ 3GM BASE/VIAL	A062697 001	Jan 22, 1987
	EQ 3GM BASE/VIAL	N050549 003	
	EQ 4GM BASE/VIAL	A062333 004	
	EQ 4GM BASE/VIAL	A062372 003	May 13, 1982
	EQ 4GM BASE/VIAL	A062697 002	Jan 22, 1987
	EQ 4GM BASE/VIAL	N050549 004	
	EQ 20GM BASE/VIAL	A062372 004	Mar 02, 1988
	EQ 20GM BASE/VIAL	N050549 005	Mar 02, 1988

MICAFUNGIN SODIUM

POWDER; INTRAVENOUS

MICAFUNGIN

+ TEVA PHARMS USA INC	EQ 50MG BASE/VIAL	N212125 001	Jul 30, 2021
	EQ 100MG BASE/VIAL	N212125 002	Jul 30, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MICONAZOLE

INJECTABLE; INJECTION

MONISTAT

JANSSEN PHARMA 10MG/ML N018040 001

MICONAZOLE NITRATE

CREAM; TOPICAL

MONISTAT-DERM

INSIGHT PHARMS 2% N017494 001

CREAM; VAGINAL

MICONAZOLE NITRATE

TEVA 2% A074136 001 Jan 04, 1995

TEVA PHARMS 2% A074030 001 Oct 30, 1992

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

M-ZOLE 7 DUAL PACK

ACTAVIS MID ATLANTIC 2%, 100MG A074586 001 Jul 17, 1997

MICONAZOLE 7 COMBINATION PACK

COSETTE 2%, 100MG A076585 001 Mar 26, 2004

LOTION; TOPICAL

MONISTAT-DERM

INSIGHT PHARMS 2% N017739 001

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

PERRIGO 100MG A074395 001 Mar 20, 1997

TAMPON; VAGINAL

MONISTAT 5

PERSONAL PRODS 100MG N018592 001 Oct 27, 1989

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

APOTHECON EQ 1MG BASE/ML A075620 001 Nov 01, 2000

EQ 5MG BASE/ML A075620 002 Nov 01, 2000

EQ 5MG BASE/ML A075641 001 Oct 19, 2000

BAXTER HLTHCARE CORP EQ 1MG BASE/ML A075637 001 Oct 31, 2000

EQ 5MG BASE/ML A075637 002 Oct 31, 2000

BEDFORD EQ 5MG BASE/ML A075249 001 Jun 23, 2000

BEN VENUE EQ 5MG BASE/ML A075455 001 Jun 20, 2000

HOSPIRA EQ 1MG BASE/ML A075396 001 Jun 20, 2000

EQ 1MG BASE/ML A075856 001 Jun 13, 2002

EQ 5MG BASE/ML A075396 002 Jun 20, 2000

EQ 5MG BASE/ML A075484 001 Jun 20, 2000

EQ 5MG BASE/ML A075856 002 Jun 13, 2002

HOSPIRA INC EQ 1MG BASE/ML A075409 002 Jun 20, 2000

EQ 5MG BASE/ML A075409 001 Jun 20, 2000

IGI LABS INC EQ 5MG BASE/ML A075263 001 Jun 26, 2000

INTL MEDICATED EQ 1MG BASE/ML A076144 001 Jan 26, 2005

EQ 5MG BASE/ML A076144 002 Jan 26, 2005

INTL MEDICATION EQ 1MG BASE/ML A076020 001 Jul 16, 2004

EQ 5MG BASE/ML A076020 002 Jul 16, 2004

PAI HOLDINGS EQ 1MG BASE/ML A078141 001 May 30, 2008

EQ 1MG BASE/ML A078511 001 Nov 10, 2008

EQ 5MG BASE/ML A078141 002 May 30, 2008

EQ 5MG BASE/ML A078511 002 Nov 10, 2008

RISING EQ 5MG BASE/ML A075481 001 Jun 30, 2000

VERSED

+ HLR EQ 1MG BASE/ML ** N018654 002 May 26, 1987

+ EQ 5MG BASE/ML ** N018654 001 Dec 20, 1985

SOLUTION; INTRAMUSCULAR

MIDAZOLAM HYDROCHLORIDE (AUTOINJECTOR)

+ RAFA LABS LTD EQ 10MG BASE/0.7ML (EQ 10MG BASE/0.7ML) N216359 001 Aug 08, 2022

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

AKORN EQ 2MG BASE/ML A075958 001 Sep 04, 2003

PHARM ASSOC EQ 2MG BASE/ML A077115 001 Sep 09, 2005

SUN PHARM INDS LTD EQ 2MG BASE/ML A076058 001 Mar 15, 2002

VERSED

+ ROCHE EQ 2MG BASE/ML ** N020942 001 Oct 15, 1998

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

CHARTWELL RX	2.5MG	A076514 001	Sep 11, 2003
	5MG	A076514 002	Sep 11, 2003
	10MG	A076514 003	Jul 02, 2004

PROAMATINE

+ TAKEDA PHARMS USA	2.5MG **	N019815 001	Sep 06, 1996
+	5MG **	N019815 002	Sep 06, 1996
+	10MG **	N019815 003	Mar 20, 2002

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

MILNACIPRAN HYDROCHLORIDE

AMNEAL PHARMS	12.5MG	A205081 001	Apr 22, 2016
	25MG	A205081 002	Apr 22, 2016
	50MG	A205081 003	Apr 22, 2016
	100MG	A205081 004	Apr 22, 2016
USPHARMA WINDLAS	12.5MG	A205071 001	Jan 27, 2016
	25MG	A205071 002	Jan 27, 2016
	50MG	A205071 003	Jan 27, 2016
	100MG	A205071 004	Jan 27, 2016

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

BAXTER HLTHCARE CORP	EQ 1MG BASE/ML	A076427 001	Sep 21, 2004
GLAND PHARMA LTD	EQ 1MG BASE/ML	A077190 001	Oct 31, 2006
HIKMA	EQ 1MG BASE/ML	A075852 001	May 28, 2002
HOSPIRA	EQ 1MG BASE/ML	A075830 001	May 28, 2002
	EQ 1MG BASE/ML	A075884 001	May 28, 2002
INTL MEDICATED	EQ 1MG BASE/ML	A076013 001	Aug 02, 2002
MYLAN INSTITUTIONAL	EQ 1MG BASE/ML	A076428 001	Jun 16, 2003

MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A076414 001	Aug 18, 2004
BAXTER HLTHCARE	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A076259 001	Aug 08, 2002
HIKMA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075510 001	May 28, 2002
HOSPIRA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075885 001	May 28, 2002
	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A075885 002	May 28, 2002
WOODWARD	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A077151 001	Jul 20, 2005

PRIMACOR

+ SANOFI AVENTIS US	EQ 1MG BASE/ML **	N019436 001	Dec 31, 1987
PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER			
+ SANOFI AVENTIS US	EQ 10MG BASE/100ML **	N020343 001	Aug 09, 1994
+	EQ 15MG BASE/100ML **	N020343 002	Aug 09, 1994
+	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	N020343 003	Aug 09, 1994
	**		
+	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	N020343 004	Aug 09, 1994
	**		

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DYNACIN

ALVOGEN	EQ 50MG BASE	A063067 003	Aug 14, 1990
	EQ 75MG BASE	A063067 002	Sep 15, 1999
	EQ 100MG BASE	A063067 001	Jul 31, 1990

MINOCIN

+ BAUSCH	EQ 75MG BASE **	N050649 003	Feb 12, 2001
TRIAx PHARMS	EQ 50MG BASE	N050315 002	
	EQ 100MG BASE	N050315 001	

CAPSULE, EXTENDED RELEASE; ORAL

XIMINO

JOURNEY	EQ 45MG BASE	N201922 001	Jul 11, 2012
	EQ 67.5MG BASE	N201922 002	Jul 11, 2012
	EQ 90MG BASE	N201922 003	Jul 11, 2012
	EQ 112.5MG BASE	N201922 004	Jul 11, 2012
	EQ 135MG BASE	N201922 005	Jul 11, 2012

INJECTABLE; INJECTION

MINOCIN

LEDERLE	EQ 100MG BASE/VIAL	A062139 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MINOCYCLINE HYDROCHLORIDE

SUSPENSION;ORAL

MINOCIN

BAUSCH EQ 50MG BASE/5ML N050445 001

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDE

+ TRIAX PHARMS EQ 50MG BASE ** N050451 003 Aug 10, 1982

+ EQ 100MG BASE ** N050451 002 Aug 10, 1982

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDE

BARR LABS INC EQ 45MG BASE A065485 001 Mar 17, 2009

EQ 65MG BASE A065485 004 May 18, 2012

EQ 80MG BASE A065485 007 Apr 26, 2017

EQ 90MG BASE A065485 002 Mar 17, 2009

EQ 105MG BASE A065485 008 Apr 26, 2017

EQ 115MG BASE A065485 005 May 18, 2012

EQ 135MG BASE A065485 003 Mar 17, 2009

IMPAX LABS INC EQ 45MG BASE A090024 001 Feb 03, 2009

EQ 90MG BASE A090024 002 Feb 03, 2009

EQ 135MG BASE A090024 003 Feb 03, 2009

MYLAN PHARMS INC EQ 45MG BASE A090911 001 Jul 20, 2010

EQ 90MG BASE A090911 002 Jul 20, 2010

EQ 135MG BASE A090911 003 Jul 20, 2010

RISING EQ 55MG BASE A203443 001 Aug 21, 2019

EQ 65MG BASE A201467 001 Jul 30, 2019

EQ 80MG BASE A203443 002 Aug 22, 2014

EQ 105MG BASE A203443 003 Aug 22, 2014

EQ 115MG BASE A201467 002 Jul 30, 2019

SOLODYN

+ BAUSCH EQ 45MG BASE ** N050808 001 May 08, 2006

+ EQ 90MG BASE ** N050808 002 May 08, 2006

+ EQ 135MG BASE ** N050808 003 May 08, 2006

MINOXIDIL

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

APOTEX INC 2% A074924 001 Apr 29, 1998

BAUSCH AND LOMB 2% A074643 001 Apr 09, 1996

COPLY PHARM 2% A074500 001 May 23, 1996

HIKMA 2% A074731 001 Dec 24, 1996

SIGHT PHARMS 2% A074743 002 Oct 18, 1996

TEVA 2% A074589 001 Apr 05, 1996

MINOXIDIL (FOR WOMEN)

APOTEX INC 2% A074924 002 Apr 29, 1998

HIKMA 2% A074731 002 May 11, 2005

SIGHT PHARMS 2% A074743 001 Oct 18, 1996

MINOXIDIL EXTRA STRENGTH (FOR MEN)

APOTEX INC 5% A075839 001 Oct 01, 2001

AVACOR PRODS 5% A075619 001 Nov 17, 2000

PERRIGO NEW YORK 5% A075737 001 Mar 15, 2002

TABLET;ORAL

LONITEN

+ PFIZER 2.5MG ** N018154 001

+ 10MG ** N018154 003

MINODYL

QUANTUM PHARMICS 2.5MG A072153 001 Jul 13, 1988

10MG A071534 001 Mar 19, 1987

MINOXIDIL

ROYCE LABS 2.5MG A071799 001 Nov 10, 1987

10MG A071796 001 Nov 10, 1987

USL PHARMA 2.5MG A071537 001 Dec 16, 1988

MIPOMERSEN SODIUM

SOLUTION;SUBCUTANEOUS

KYNAMRO

+ KASTLE THERAPS LLC 200MG/ML (200MG/ML) N203568 001 Jan 29, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MIRABEGRON

TABLET, EXTENDED RELEASE;ORAL

MIRABEGRON

SAWAI USA	25MG	A209446	001	Dec 27, 2019
ZYDUS PHARMS	25MG	A209488	001	Sep 29, 2022
	50MG	A209488	002	Sep 29, 2022

MIRTAZAPINE

TABLET;ORAL

MIRTAZAPINE

ACTAVIS ELIZABETH	15MG	A076241	001	Jun 25, 2003
	15MG	A076308	001	Jun 20, 2003
	30MG	A076241	002	Jun 25, 2003
	30MG	A076308	002	Jun 20, 2003
	45MG	A076241	003	Jun 25, 2003
	45MG	A076308	003	Jun 20, 2003
ACTAVIS LABS FL INC	15MG	A076336	001	Jun 20, 2003
	30MG	A076336	002	Jun 20, 2003
	45MG	A076336	003	Jun 20, 2003
IVAX SUB TEVA PHARMS	15MG	A076244	001	Dec 22, 2003
	30MG	A076244	002	Dec 22, 2003
	45MG	A076244	003	Dec 22, 2003
MYLAN PHARMS INC	15MG	A076176	001	Jun 19, 2003
	30MG	A076176	002	Jun 19, 2003
	45MG	A076176	003	Jun 19, 2003
ROXANE	15MG	A076270	001	Jun 19, 2003
	30MG	A076270	002	Jun 19, 2003
	45MG	A076270	003	Jun 19, 2003
UPSHER SMITH LABS	15MG	A076189	001	Jun 19, 2003
	30MG	A076189	002	Jun 19, 2003
	45MG	A076189	003	Jun 19, 2003
WATSON LABS	15MG	A076312	001	Jun 19, 2003
	30MG	A076312	002	Jun 19, 2003
	45MG	A076312	003	Jun 19, 2003
REMERON				
+ ORGANON	45MG **	N020415	003	Mar 17, 1997

TABLET, ORALLY DISINTEGRATING;ORAL

MIRTAZAPINE

ACTAVIS ELIZABETH	15MG	A076689	001	Aug 31, 2005
	15MG	A077959	001	Feb 14, 2011
	30MG	A076689	002	Aug 31, 2005
	30MG	A077959	002	Feb 14, 2011
	45MG	A076689	003	Aug 31, 2005
	45MG	A077959	003	Feb 14, 2011
ACTAVIS LABS FL INC	15MG	A076307	001	Dec 17, 2003
	30MG	A076307	002	Dec 17, 2003
	45MG	A076307	003	Feb 28, 2006
IMPAX LABS INC	15MG	A076901	001	Jun 28, 2005
	30MG	A076901	002	Jun 28, 2005
	45MG	A076901	003	Jun 28, 2005

MISOPROSTOL

TABLET;ORAL

MISOPROSTOL

ACQ PHARMA	0.1MG	A210201	001	Jul 02, 2019
	0.2MG	A210201	002	Jul 02, 2019
NOVEL LABS INC	0.1MG	A091667	001	Jul 25, 2012
	0.2MG	A091667	002	Jul 25, 2012

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

HIKMA	5MG/VIAL	A064117	001	Apr 19, 1995
HOSPIRA	20MG/VIAL	A064106	001	Nov 29, 1995
MITOZYTREX				
+ SUPERGEN	5MG/VIAL **	N050763	001	Nov 14, 2002
MUTAMYCIN				
+ BRISTOL	5MG/VIAL **	N050450	001	
+ BRISTOL MYERS	20MG/VIAL **	N050450	002	
	5MG/VIAL	A062336	001	
	20MG/VIAL	A062336	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MITOMYCININJECTABLE; INJECTION
MUTAMYCIN

40MG/VIAL **

A062336 003 Mar 10, 1988

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

FRESENIUS KABI ONCOL EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

A078606 001 May 14, 2008

EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)

A078606 002 May 14, 2008

EQ 30MG BASE/15ML (EQ 2MG BASE/ML)

A078606 003 May 14, 2008

MYLAN LABS LTD EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

A201014 001 Dec 11, 2012

RISING EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

A078980 001 Apr 13, 2009

EQ 30MG BASE/15ML (EQ 2MG BASE/ML)

A078980 002 Apr 13, 2009

NOVANTRONE

+ EMD SERONO

EQ 20MG BASE/10ML (EQ 2MG BASE/ML) **

N019297 001 Dec 23, 1987

+ EMD SERONO

EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML) **

N019297 002 Dec 23, 1987

+ EMD SERONO

EQ 30MG BASE/15ML (EQ 2MG BASE/ML) **

N019297 003 Dec 23, 1987

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBVIE

EQ 0.5MG BASE/ML

N020098 002 Jan 22, 1992

EQ 50MG BASE/100ML

N020098 003 Jan 22, 1992

MIVACURIUM CHLORIDE

RISING

EQ 2MG BASE/ML

A078562 001 Apr 30, 2009

SOLUTION; INTRAVENOUS

MIVACRON

+ ABBVIE

EQ 2MG BASE/ML (EQ 2MG BASE/ML) **

N020098 001 Jan 22, 1992

+ ABBVIE

EQ 10MG BASE/5ML (EQ 2MG BASE/ML)

N020098 004 Jan 22, 1992

+ ABBVIE

EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

N020098 005 Jan 22, 1992

MODAFINIL

TABLET; ORAL

MODAFINIL

HIKMA PHARMS

100MG

A090543 001 Sep 26, 2012

200MG

A090543 002 Sep 26, 2012

MYLAN PHARMS INC

100MG

A076594 001 Jul 16, 2012

200MG

A076594 002 Jul 16, 2012

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

UNIVASC

UCB INC

7.5MG **

N020312 001 Apr 19, 1995

15MG **

N020312 002 Apr 19, 1995

MOLINDONE HYDROCHLORIDE

CAPSULE; ORAL

MOBAN

+ CHARTWELL RX

5MG **

N017111 001

+ CHARTWELL RX

10MG **

N017111 002

+ CHARTWELL RX

25MG **

N017111 003

CONCENTRATE; ORAL

MOBAN

ENDO PHARMS

20MG/ML

N017938 001

TABLET; ORAL

MOBAN

+ CHARTWELL RX

5MG **

N017111 004

+ CHARTWELL RX

10MG **

N017111 005

+ CHARTWELL RX

25MG **

N017111 006

+ CHARTWELL RX

50MG **

N017111 007

+ CHARTWELL RX

100MG **

N017111 008

MOMETASONE FUROATE

CREAM; TOPICAL

ELOCON

ORGANON

0.1% **

N019625 001 May 06, 1987

+ ORGANON

0.1% **

N019625 002 Apr 19, 2013

MOMETASONE FUROATE

ANDA REPOSITORY

0.1%

A076591 001 Apr 18, 2007

FOUGERA PHARMS

0.1%

A076171 001 Apr 08, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MOMETASONE FUROATE

LOTION; TOPICAL

ELOCON

+ ORGANON

0.1% **

N019796 001 Mar 30, 1989

MOMETASONE FUROATE

ENCUBE

0.1%

A076499 001 Nov 21, 2007

OINTMENT; TOPICAL

ELOCON

+ ORGANON

0.1%

N019543 001 Apr 30, 1987

MOMETASONE FUROATE

ENCUBE

0.1%

A076481 001 Nov 14, 2003

TARO

0.1%

A076624 001 Dec 03, 2004

TORRENT

0.1%

A207899 001 Jul 13, 2018

SPRAY, METERED; NASAL

MOMETASONE FUROATE

AMNEAL

0.05MG/SPRAY

A217460 001 Jun 14, 2023

NASONEX

+ ORGANON LLC

0.05MG/SPRAY **

N020762 001 Oct 01, 1997

MONOBENZONE

CREAM; TOPICAL

BENOQUIN

VALEANT PHARM INTL

20%

N008173 003

MONOCTANOIN

LIQUID; PERFUSION, BILIARY

MOCTANIN

ETHITEK

100%

N019368 001 Oct 29, 1985

MONTELUKAST SODIUM

GRANULE; ORAL

MONTELUKAST SODIUM

MYLAN PHARMS INC

EQ 4MG BASE/PACKET

A202776 001 Dec 18, 2012

TABLET; ORAL

MONTELUKAST SODIUM

AJANTA PHARMA LTD

EQ 10MG BASE

A203432 001 Jul 31, 2015

APOTEX CORP

EQ 10MG BASE

A201294 001 Aug 03, 2012

BRECKENRIDGE

EQ 10MG BASE

A205319 001 Oct 30, 2020

HIKMA

EQ 10MG BASE

A090655 001 Aug 03, 2012

L PERRIGO CO

EQ 10MG BASE

A206112 001 Apr 26, 2017

MYLAN PHARMS INC

EQ 10MG BASE

A079103 001 Aug 03, 2012

STRIDES PHARMA

EQ 10MG BASE

A091576 001 Aug 03, 2012

TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

AJANTA PHARMA LTD

EQ 4MG BASE

A203328 001 Jul 31, 2015

EQ 5MG BASE

A203328 002 Jul 31, 2015

APOTEX INC

EQ 4MG BASE

A201508 001 Aug 03, 2012

EQ 5MG BASE

A201508 002 Aug 03, 2012

HIKMA

EQ 4MG BASE

A091128 001 Aug 03, 2012

EQ 5MG BASE

A091128 002 Aug 03, 2012

JUBILANT GENERICS

EQ 4MG BASE

A203795 001 Feb 27, 2015

EQ 5MG BASE

A203795 002 Feb 27, 2015

MYLAN PHARMS INC

EQ 4MG BASE

A079142 001 Aug 03, 2012

EQ 5MG BASE

A079142 002 Aug 03, 2012

STRIDES PHARMA

EQ 4MG BASE

A091588 001 Aug 03, 2012

EQ 5MG BASE

A091588 002 Aug 03, 2012

UNICHEM

EQ 4MG BASE

A208621 001 Jul 02, 2018

EQ 5MG BASE

A208621 002 Jul 02, 2018

MORICIZINE HYDROCHLORIDE

TABLET; ORAL

ETHMOZINE

SHIRE

200MG

N019753 001 Jun 19, 1990

250MG

N019753 002 Jun 19, 1990

300MG

N019753 003 Jun 19, 1990

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

AVINZA

KING PHARMS LLC	30MG	N021260 001	Mar 20, 2002
	45MG	N021260 005	Dec 18, 2008
	60MG	N021260 002	Mar 20, 2002
	75MG	N021260 006	Dec 18, 2008
	90MG	N021260 003	Mar 20, 2002
	120MG	N021260 004	Mar 20, 2002

KADIAN

+ ALLERGAN	10MG	N020616 008	Apr 20, 2007
	20MG	N020616 001	Jul 03, 1996
	30MG	N020616 004	Mar 09, 2001
	40MG	N020616 009	Jul 09, 2012
	50MG	N020616 002	Jul 03, 1996
	60MG	N020616 005	Mar 09, 2001
	70MG	N020616 010	Jul 09, 2012
	80MG	N020616 006	Oct 27, 2006
	100MG	N020616 003	Jul 03, 1996
	130MG	N020616 011	Jul 09, 2012
	150MG	N020616 012	Jul 09, 2012
	200MG	N020616 007	Feb 27, 2007

MORPHINE SULFATE

IMPAX LABS INC	20MG	A200411 001	Apr 12, 2016
	30MG	A200411 002	Apr 12, 2016
	40MG	A200411 007	Jul 25, 2018
	50MG	A200411 003	Apr 12, 2016
	60MG	A200411 004	Apr 12, 2016
	80MG	A200411 005	Apr 12, 2016
	100MG	A200411 006	Apr 12, 2016
NORTEC DEV ASSOC	20MG	A203158 001	Aug 04, 2021
	30MG	A203158 002	Aug 04, 2021
	50MG	A203158 003	Aug 04, 2021
	60MG	A203158 004	Aug 04, 2021
	80MG	A203158 005	Aug 04, 2021
	100MG	A203158 006	Aug 04, 2021
STRIDES PHARMA	20MG	A200812 001	Nov 10, 2011
	30MG	A200812 002	Nov 10, 2011
	50MG	A200812 003	Nov 10, 2011
	60MG	A200812 004	Nov 10, 2011
	80MG	A200812 005	Nov 10, 2011
	100MG	A200812 006	Nov 10, 2011
TEVA PHARMS USA	20MG	A202718 001	Dec 29, 2014
	30MG	A202718 002	Dec 29, 2014
	40MG	A202718 007	Jun 03, 2015
	50MG	A202718 003	Dec 29, 2014
	60MG	A202718 004	Dec 29, 2014
	70MG	A202718 008	Jun 03, 2015
	80MG	A202718 005	Dec 29, 2014
	100MG	A202718 006	Dec 29, 2014

INJECTABLE; INJECTION

ASTRAMORPH PF

FRESENIUS KABI USA	0.5MG/ML	A071050 001	Oct 07, 1986
	0.5MG/ML	A071051 001	Oct 07, 1986
	1MG/ML	A071052 001	Oct 07, 1986
	1MG/ML	A071053 001	Oct 07, 1986

MORPHINE SULFATE

HOSPIRA	1MG/ML	A071850 001	May 11, 1988
+ HOSPIRA INC	15MG/ML	N202515 005	Nov 14, 2011
ICU MEDICAL INC	0.5MG/ML	N019917 001	Oct 30, 1992
	1MG/ML **	N019916 001	Oct 30, 1992
	5MG/ML **	N019916 002	Oct 27, 2006
SPECGX LLC	1MG/ML	N020631 001	Jul 03, 1996
	2MG/ML	N020631 002	Jul 03, 1996
WATSON LABS	0.5MG/ML	A073373 001	Sep 30, 1991
	0.5MG/ML	A073375 001	Sep 30, 1991
	1MG/ML	A073374 001	Sep 30, 1991
	1MG/ML	A073376 001	Sep 30, 1991

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORPHINE SULFATEINJECTABLE, LIPOSOMAL;EPIDURAL
DEPODUR

PACIRA PHARMS INC	10MG/ML (10MG/ML)	N021671 001	May 18, 2004
	15MG/1.5ML (10MG/ML)	N021671 002	May 18, 2004
	20MG/2ML (10MG/ML)	N021671 003	May 18, 2004

SOLUTION;INTRAMUSCULAR

MORPHINE SULFATE (AUTOINJECTOR)

+ MERIDIAN MEDCL TECHN	10MG/0.7ML (10MG/0.7ML)	N019999 001	Jul 12, 1990
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SOLUTION;ORAL

MORPHINE SULFATE

ANI PHARMS	10MG/5ML	A205509 001	Apr 17, 2018
	20MG/5ML	A205509 002	Apr 17, 2018
	100MG/5ML	A205509 003	Apr 17, 2018
CHARTWELL MOLECULAR	10MG/5ML	A202309 001	Nov 25, 2015
	20MG/5ML	A202310 001	Oct 30, 2015
	100MG/5ML	N201517 001	Jun 23, 2011
NOSTRUM LABS INC	10MG/5ML	A201011 001	Feb 05, 2014
	20MG/5ML	A201011 002	Feb 05, 2014
	100MG/5ML	A201011 003	Oct 06, 2016
TRIS PHARMA INC	20MG/5ML	A203519 001	May 18, 2016
VISTAPHARM	10MG/5ML	A201947 001	Jan 05, 2012
	20MG/5ML	A201947 002	Jan 05, 2012
WINDER LABS LLC	10MG/5ML	A211454 001	Feb 12, 2021
	20MG/5ML	A211454 002	Feb 12, 2021
	100MG/5ML	A211454 003	Feb 12, 2021

TABLET;ORAL

MORPHINE SULFATE

DR REDDYS LABS SA	15MG	A207270 001	Jan 12, 2022
	30MG	A207270 002	Jan 12, 2022
INGENUS PHARMS NJ	15MG	A215584 001	Feb 07, 2022
	30MG	A215584 002	Feb 07, 2022

TABLET, EXTENDED RELEASE;ORAL

ARYMO ER

+ ZYLA	15MG	N208603 001	Jan 09, 2017
	30MG	N208603 002	Jan 09, 2017
	60MG	N208603 003	Jan 09, 2017

MORPHABOND ER

+ OHEMO LIFE	15MG	N206544 001	Oct 02, 2015
	30MG	N206544 002	Oct 02, 2015
	60MG	N206544 003	Oct 02, 2015
	100MG	N206544 004	Oct 02, 2015

MORPHINE SULFATE

DR REDDYS LABS SA	15MG	A205386 001	Oct 28, 2016
	30MG	A205386 002	Oct 28, 2016
	60MG	A205386 003	Oct 28, 2016
	100MG	A205386 004	Oct 28, 2016
EPIC PHARMA LLC	15MG	A091357 001	Jun 23, 2016
	30MG	A091357 002	Jun 23, 2016
	60MG	A091357 003	Jun 23, 2016
	100MG	A091357 004	Jun 23, 2016
	200MG	A091357 005	Jun 23, 2016
NESHER PHARMS	15MG	A076733 001	May 19, 2004
	30MG	A076720 002	Dec 23, 2005
	60MG	A076720 001	May 19, 2004
	100MG	A077855 001	Sep 27, 2007
	200MG	A077855 002	Sep 27, 2007
RISING	15MG	A200824 001	Oct 18, 2011
	30MG	A200824 002	Oct 18, 2011
	60MG	A200824 003	Oct 18, 2011
	100MG	A200824 004	Oct 18, 2011
	200MG	A200824 005	Oct 18, 2011
SUN PHARM INDUSTRIES	15MG	A205634 001	Aug 25, 2016
	30MG	A205634 002	Aug 25, 2016
	60MG	A205634 003	Aug 25, 2016
	100MG	A205634 004	Aug 25, 2016
	200MG	A205634 005	Aug 25, 2016
WATSON LABS	100MG	A075656 001	Jan 30, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORPHINE SULFATETABLET, EXTENDED RELEASE;ORAL
ORAMORPH SR

XANODYNE PHARMS INC	15MG	N019977 004	Nov 23, 1994
	30MG	N019977 001	Aug 15, 1991
	60MG	N019977 002	Aug 15, 1991
	100MG	N019977 003	Aug 15, 1991

MORPHINE SULFATE; NALTREXONE HYDROCHLORIDECAPSULE, EXTENDED RELEASE;ORAL
EMBEDA

+	ALPHARMA PHARMS	20MG;0.8MG **	N022321 001	Aug 13, 2009
+		30MG;1.2MG **	N022321 002	Aug 13, 2009
+		50MG;2MG **	N022321 003	Aug 13, 2009
+		60MG;2.4MG **	N022321 004	Aug 13, 2009
+		80MG;3.2MG **	N022321 005	Aug 13, 2009
+		100MG;4MG **	N022321 006	Aug 13, 2009

MOXALACTAM DISODIUMINJECTABLE; INJECTION
MOXAM

LILLY	EQ 250MG BASE/VIAL	N050550 001
	EQ 500MG BASE/VIAL	N050550 002
	EQ 1GM BASE/VIAL	N050550 003
	EQ 2GM BASE/VIAL	N050550 004
	EQ 10GM BASE/VIAL	N050550 008

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION;INTRAVENOUS

AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

+	BAYER HLTHCARE	400MG/250ML (1.6MG/ML) **	N021277 001	Nov 30, 2001
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SOLUTION/DROPS;OPHTHALMIC

MOXEZA

+	HARROW EYE	EQ 0.5% BASE	N022428 001	Nov 19, 2010
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MOXIFLOXACIN HYDROCHLORIDE

AKORN

	EQ 0.5% BASE	A202916 001	Nov 09, 2017
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TABLET;ORAL

AVELOX

+	BAYER HLTHCARE	EQ 400MG BASE **	N021085 001	Dec 10, 1999
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MOXIFLOXACIN HYDROCHLORIDE

MYLAN

	EQ 400MG BASE	A204635 001	Aug 31, 2015
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SUNSHINE

	EQ 400MG BASE	A206295 001	Sep 28, 2018
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TORRENT

	EQ 400MG BASE	A200160 001	Apr 03, 2014
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MUPIROCIN

OINTMENT;TOPICAL

BACTROBAN

+	GLAXOSMITHKLINE	2% **	N050591 001	Dec 31, 1987
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MUPIROCIN CALCIUM

CREAM;TOPICAL

BACTROBAN

+	GLAXOSMITHKLINE	EQ 2% BASE **	N050746 001	Dec 11, 1997
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MUPIROCIN

PADAGIS ISRAEL

	EQ 2% BASE	A212465 001	Aug 03, 2022
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OINTMENT;NASAL

BACTROBAN

+	GLAXOSMITHKLINE	EQ 2% BASE	N050703 001	Sep 18, 1995
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MYCOPHENOLATE MOFETIL

CAPSULE;ORAL

MYCOPHENOLATE MOFETIL

APOTEX CORP

	250MG	A090419 001	Apr 22, 2009
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DR REDDYS LABS LTD

	250MG	A091315 001	Oct 27, 2011
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JUBILANT CADISTA

	250MG	A090762 001	Dec 15, 2014
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STRIDES PHARMA

	250MG	A090111 001	Dec 22, 2009
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ZYDUS PHARMS USA INC

	250MG	A065433 001	May 04, 2009
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TABLET;ORAL

MYCOPHENOLATE MOFETIL

APOTEX

	500MG	A090499 001	Apr 22, 2009
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DR REDDYS LABS LTD

	500MG	A090464 001	Sep 13, 2010
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JUBILANT CADISTA

	500MG	A090661 001	Dec 15, 2014
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MYCOPHENOLATE MOFETIL

TABLET; ORAL

MYCOPHENOLATE MOFETIL

TEVA PHARMS	500MG
ZYDUS PHARMS USA INC	500MG

A065457	001	May 04, 2009
A065477	001	May 04, 2009

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

AMNEAL	500MG/VIAL
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A211374	001	Mar 05, 2021
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MYCOPHENOLIC SODIUM

TABLET, DELAYED RELEASE; ORAL

MYCOPHENOLIC SODIUM

TEVA PHARMS USA	EQ 180MG BASE
	EQ 360MG BASE

A202720	001	Oct 30, 2014
A202720	002	Oct 30, 2014

NABUMETONE

TABLET; ORAL

NABUMETONE

AUROBINDO PHARMA USA	500MG
	750MG
COPLEY PHARM	750MG
EPIC PHARMA LLC	500MG
	750MG
IMPAX LABS INC	500MG
	750MG
NOSTRUM LABS INC	500MG
	750MG
OXFORD PHARMS	500MG
	750MG

A090516	001	Jul 12, 2010
A090516	002	Jul 12, 2010
A075179	001	Jun 06, 2000
A075590	001	Feb 25, 2002
A075590	002	Feb 25, 2002
A075189	001	May 26, 2000
A075189	002	Sep 24, 2001
A090427	001	Dec 30, 2011
A090427	002	Dec 30, 2011
A079093	001	Feb 27, 2009
A079093	002	Feb 27, 2009

RELAFEN

+	SMITHKLINE BEECHAM	500MG **
+		750MG **

N019583	001	Dec 24, 1991
N019583	002	Dec 24, 1991

NADOLOL

TABLET; ORAL

CORGARD

USWM	120MG
	160MG

N018063	003
N018063	004

NADOLOL

HERITAGE PHARMA	20MG
	40MG
	80MG
	120MG
	160MG
NOVAST LABS	20MG
	40MG
	80MG
TEVA PHARMS	80MG
	120MG
	160MG

A074229	001	Aug 30, 1996
A074229	002	Aug 30, 1996
A074255	001	Jan 24, 1996
A074255	002	Jan 24, 1996
A074255	003	Jan 24, 1996
A210786	001	Jun 01, 2018
A210786	002	Jun 01, 2018
A210786	003	Jun 01, 2018
A074368	001	Aug 31, 1994
A074368	002	Aug 31, 1994
A074368	003	Aug 31, 1994

NAFCILLIN SODIUM

CAPSULE; ORAL

UNIPEN

WYETH AYERST	EQ 250MG BASE
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N050111	001
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FOR SOLUTION; ORAL

UNIPEN

WYETH AYERST	EQ 250MG BASE/5ML
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N050199	001
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INJECTABLE; INJECTION

NAFCILLIN SODIUM

APOTHECON	EQ 500MG BASE/VIAL
	EQ 1GM BASE/VIAL
	EQ 2GM BASE/VIAL
	EQ 4GM BASE/VIAL
FRESENIUS	EQ 1GM BASE/VIAL
	EQ 2GM BASE/VIAL
SANDOZ	EQ 500MG BASE/VIAL
	EQ 1GM BASE/VIAL
	EQ 1GM BASE/VIAL
	EQ 2GM BASE/VIAL

A061984	001	
A061984	002	
A061984	003	
A061984	005	
A206682	001	Dec 10, 2019
A206682	002	Dec 10, 2019
A062527	001	Aug 02, 1984
A062527	002	Aug 02, 1984
A062732	001	Dec 23, 1986
A062527	003	Aug 02, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

	EQ 2GM BASE/VIAL	A062732 002	Dec 23, 1986
	EQ 10GM BASE/VIAL	A062527 004	Aug 02, 1984
WATSON LABS INC	EQ 500MG BASE/VIAL	A062844 001	Oct 26, 1988
	EQ 1GM BASE/VIAL	A062844 002	Oct 26, 1988
	EQ 1.5GM BASE/VIAL	A062844 003	Oct 26, 1988
	EQ 2GM BASE/VIAL	A062844 004	Oct 26, 1988
	EQ 4GM BASE/VIAL	A062844 005	Oct 26, 1988
	EQ 10GM BASE/VIAL	A063008 001	Sep 29, 1988

NALLPEN

GLAXOSMITHKLINE

	EQ 500MG BASE/VIAL	A061999 001	
	EQ 1GM BASE/VIAL	A061999 002	
	EQ 1GM BASE/VIAL	A062755 001	Dec 19, 1986
	EQ 2GM BASE/VIAL	A061999 003	
	EQ 2GM BASE/VIAL	A062755 002	Dec 19, 1986
	EQ 10GM BASE/VIAL	A061999 004	

UNIPEN

WYETH AYERST

	EQ 500MG BASE/VIAL **	A062717 001	Dec 16, 1986
+	EQ 500MG BASE/VIAL **	N050320 001	
	EQ 1GM BASE/VIAL **	A062717 002	Dec 16, 1986
	EQ 2GM BASE/VIAL **	A062717 004	Dec 16, 1986
+	EQ 2GM BASE/VIAL **	N050320 003	
+	EQ 4GM BASE/VIAL **	N050320 004	
+	EQ 10GM BASE/VIAL **	N050320 005	
+	EQ 20GM BASE/VIAL **	N050320 006	

UNIPEN IN PLASTIC CONTAINER

+ WYETH AYERST

EQ 1GM BASE/VIAL ** N050320 002

TABLET; ORAL

UNIPEN

WYETH AYERST

EQ 500MG BASE N050462 001

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

+ SEBELA IRELAND LTD 1% ** N019599 001 Feb 29, 1988

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE

ABRAXIS PHARM

	10MG/ML	A070751 001	Jul 02, 1986
	20MG/ML	A070752 001	Sep 24, 1986

NALBUPHINE HYDROCHLORIDE

ABBOTT

20MG/ML A070917 001 Feb 03, 1989

ABBVIE

1.5MG/ML N020200 001 Mar 12, 1993

DR REDDYS

10MG/ML A074471 001 Mar 19, 1998

20MG/ML A074471 002 Mar 19, 1998

IGI LABS INC

10MG/ML A072070 001 Apr 10, 1989

10MG/ML A072071 001 Apr 10, 1989

10MG/ML A072072 001 Apr 10, 1989

20MG/ML A072073 001 Apr 10, 1989

20MG/ML A072074 001 Apr 10, 1989

20MG/ML A072075 001 Apr 10, 1989

RISING

10MG/ML A206506 001 Feb 06, 2019

10MG/ML A207595 001 Jan 11, 2019

20MG/ML A206506 002 Feb 06, 2019

20MG/ML A207595 002 Jan 11, 2019

NUBAIN

+ PAR PHARM INC

10MG/ML ** N018024 001

+ 20MG/ML ** N018024 002 May 27, 1982

NALIDIXIC ACID

SUSPENSION; ORAL

NEGGRAM

SANOFI AVENTIS US

250MG/5ML N017430 001

TABLET; ORAL

NALIDIXIC ACID

SUN PHARM INDUSTRIES

250MG A070270 001 Jun 29, 1988

500MG A070271 001 Jun 29, 1988

1GM A070272 001 Jun 29, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NALIDIXIC ACID

TABLET; ORAL

NALIDIXIC ACID

WATSON LABS

250MG

A071936 001 Jun 29, 1988

500MG

A072061 001 Jun 29, 1988

1GM

A071919 001 Jun 29, 1988

NEGGRAM

SANOFI AVENTIS US

250MG

N014214 002

500MG

N014214 004

1GM

N014214 005

NALMEFENE HYDROCHLORIDE

SOLUTION; INTRAMUSCULAR, INTRAVENOUS, SUBCUTANEOUS

REVEX

+ HIKMA

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **

N020459 001 Apr 17, 1995

+

EQ 2MG BASE/2ML (EQ 1MG BASE/ML) **

N020459 002 Apr 17, 1995

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

HIKMA

0.4MG/ML

A070298 001 Sep 24, 1986

0.4MG/ML

A070496 001 Sep 24, 1986

WYETH AYERST

0.02MG/ML

A070188 001 Sep 24, 1986

0.02MG/ML

A070189 001 Sep 24, 1986

0.4MG/ML

A070190 001 Sep 24, 1986

0.4MG/ML

A070191 001 Sep 24, 1986

NALOXONE HYDROCHLORIDE

ABRAXIS PHARM

0.02MG/ML

A070648 001 Nov 17, 1986

0.02MG/ML

A070661 001 Nov 17, 1986

0.4MG/ML

A070649 001 Nov 17, 1986

1MG/ML

A071604 001 Dec 16, 1988

ASTRAZENECA

0.02MG/ML

A072081 001 Apr 11, 1989

CHARTWELL RX

0.4MG/ML

A207846 001 Dec 17, 2018

HIKMA

0.02MG/ML

A071272 001 May 24, 1988

1MG/ML

A071273 001 May 24, 1988

1MG/ML

A071274 001 May 24, 1988

1MG/ML

A071287 001 May 24, 1988

HOSPIRA

0.02MG/ML

A070171 001 Sep 24, 1986

0.02MG/ML

A070252 001 Jan 16, 1987

0.02MG/ML

A070253 001 Jan 16, 1987

0.4MG/ML

A070254 001 Jan 07, 1987

0.4MG/ML

A070255 001 Jan 07, 1987

IGI LABS INC

0.02MG/ML

A072082 001 Apr 11, 1989

0.02MG/ML

A072083 001 Apr 11, 1989

0.02MG/ML

A072084 001 Apr 11, 1989

0.02MG/ML

A072085 001 Apr 11, 1989

0.4MG/ML

A072086 001 Apr 11, 1989

0.4MG/ML

A072087 001 Apr 11, 1989

0.4MG/ML

A072088 001 Apr 11, 1989

0.4MG/ML

A072089 001 Apr 11, 1989

0.4MG/ML

A072090 001 Apr 11, 1989

1MG/ML

A072091 001 Apr 11, 1989

1MG/ML

A072092 001 Apr 11, 1989

1MG/ML

A072093 001 Apr 11, 1989

INTL MEDICATION

0.4MG/ML

A070417 001 Sep 24, 1986

0.4MG/ML

A070639 001 Sep 24, 1986

1MG/ML

A072115 001 Apr 27, 1988

MARSAM PHARMS LLC

0.4MG/ML

A071811 001 Jul 19, 1988

PAR STERILE PRODUCTS

0.4MG/ML

A211286 001 Jan 17, 2020

1MG/ML

A215964 001 Jul 29, 2022

SMITH AND NEPHEW

0.02MG/ML

A071671 001 Nov 17, 1987

0.4MG/ML

A071681 001 Nov 17, 1987

0.4MG/ML

A071682 001 Nov 17, 1987

SOLOPAK

0.02MG/ML

A071672 001 Nov 17, 1987

0.4MG/ML

A071683 001 Nov 17, 1987

WATSON LABS

0.4MG/ML

A071339 001 Nov 18, 1987

NARCAN

+ ADAPT

0.02MG/ML **

N016636 002

+

0.4MG/ML **

N016636 001

+

1MG/ML **

N016636 003 Jun 14, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NARCAN

BRISTOL MYERS SQUIBB	0.4MG/ML	A071083	001	Jul 28, 1988
	1MG/ML	A071084	001	Jul 28, 1988
	1MG/ML	A071311	001	Jul 28, 1988

SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

EVZIO

+ KALEO INC	0.4MG/0.4ML (0.4MG/0.4ML)	N205787	001	Apr 03, 2014
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EVZIO (AUTOINJECTOR)

+ KALEO INC	2MG/0.4ML (2MG/0.4ML)	N209862	001	Oct 19, 2016
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NALOXONE HYDROCHLORIDE (AUTOINJECTOR)

+ KALEO INC	10MG/0.4ML (10MG/0.4ML)	N215457	001	Feb 28, 2022
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SPRAY, METERED; NASAL

NARCAN

+ EMERGENT	2MG/SPRAY **	N208411	002	Jan 24, 2017
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REXTOVY

+ AMPHASTAR PHARMS INC	4MG/SPRAY	N208969	001	Mar 07, 2023
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NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARGINIQ

+ PURDUE PHARMA LP	5MG; 10MG	N205777	001	Jul 23, 2014
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	10MG; 20MG	N205777	002	Jul 23, 2014
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	20MG; 40MG	N205777	003	Jul 23, 2014
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NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN NX

SANOFI AVENTIS US	EQ 0.5MG BASE; EQ 50MG BASE **	N018733	001	Dec 16, 1982
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NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

FOSUN PHARMA	50MG	A075434	001	Mar 08, 2000
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REVIA

+ TEVA WOMENS	50MG	N018932	001	Nov 20, 1984
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NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

TROXYCA ER

PFIZER	1.2MG; 10MG	N207621	001	Aug 19, 2016
	2.4MG; 20MG	N207621	002	Aug 19, 2016
	3.6MG; 30MG	N207621	003	Aug 19, 2016
	4.8MG; 40MG	N207621	004	Aug 19, 2016
	7.2MG; 60MG	N207621	005	Aug 19, 2016
	9.6MG; 80MG	N207621	006	Aug 19, 2016

NANDROLONE DECANOATE

INJECTABLE; INJECTION

DECA-DURABOLIN

WOODWARD SPECL	50MG/ML	N013132	001	Jun 12, 1986
	100MG/ML	N013132	002	Jun 12, 1986
	200MG/ML **	N013132	003	Jun 12, 1986

NANDROLONE DECANOATE

ABRAXIS PHARM	100MG/ML	A088290	001	Oct 03, 1983
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	200MG/ML	A088317	001	Oct 14, 1983
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AKORN	100MG/ML	A087519	001	Sep 28, 1983
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AM REGENT	200MG/ML	A091252	001	Aug 30, 2010
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WATSON LABS	50MG/ML	A086385	001	Jan 13, 1984
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	50MG/ML	A087598	001	Oct 06, 1983
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	50MG/ML	A088554	001	Feb 10, 1986
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	100MG/ML	A086598	001	Jan 13, 1984
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	100MG/ML	A087599	001	Oct 06, 1983
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	200MG/ML	A088128	001	Dec 05, 1983
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NANDROLONE PHENPROPIONATE

INJECTABLE; INJECTION

DURABOLIN

ORGANON USA INC 25MG/ML
50MG/MLN011891 001
N011891 002

NANDROLONE PHENPROPIONATE

WATSON LABS 25MG/ML
50MG/MLA086386 001 Jun 17, 1983
A087488 001 Jun 17, 1983NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ALBALON

ALLERGAN 0.1% **

A080248 001

NAFAZAIR

BAUSCH AND LOMB 0.1%

A040073 001 May 25, 1994

PHARMAFAIR 0.1%

A088101 001 Apr 15, 1983

NAPHAZOLINE HYDROCHLORIDE

RISING 0.1%

A083590 001

NAPHCN FORTE

+ ALCON 0.1%

A080229 001

OPCON

BAUSCH AND LOMB 0.1%

A087506 001

VASOCON

NOVARTIS 0.1%

A080235 002 Mar 24, 1983

NAPROXEN

TABLET; ORAL

NAPROSYN

+ ATNAHS PHARMA US 250MG

N017581 002

+ 375MG

N017581 003

NAPROXEN

CHARTWELL MOLECULES 250MG

A074410 001 Apr 28, 1995

375MG

A074410 002 Apr 28, 1995

500MG

A074410 003 Apr 28, 1995

DAVA PHARMS INC 250MG

A074105 001 Dec 21, 1993

375MG

A074105 002 Dec 21, 1993

500MG

A074105 003 Dec 21, 1993

HAMILTON PHARMS 250MG

A074110 001 Oct 30, 1992

375MG

A074110 002 Oct 30, 1992

500MG

A074110 003 Oct 30, 1992

HIKMA INTL PHARMS 250MG

A076494 001 Jan 14, 2004

375MG

A076494 002 Jan 14, 2004

500MG

A076494 003 Jan 14, 2004

IVAX SUB TEVA PHARMS 250MG

A074111 001 Feb 28, 1995

375MG

A074111 002 Feb 28, 1995

500MG

A074111 003 Feb 28, 1995

L PERRIGO CO 250MG

A077339 001 Apr 27, 2005

375MG

A077339 002 Apr 27, 2005

500MG

A077339 003 Apr 27, 2005

MYLAN 250MG

A074121 001 Dec 21, 1993

375MG

A074121 002 Dec 21, 1993

500MG

A074121 003 Dec 21, 1993

PLIVA 250MG

A074182 001 Jun 27, 1996

375MG

A074182 002 Jun 27, 1996

500MG

A074182 003 Jun 27, 1996

PUREPAC PHARM 250MG

A074263 001 Dec 21, 1993

375MG

A074263 002 Dec 21, 1993

500MG

A074263 003 Dec 21, 1993

ROXANE 250MG

A074211 001 Feb 28, 1994

375MG

A074211 002 Feb 28, 1994

500MG

A074211 003 Feb 28, 1994

TEVA 250MG

A074129 001 Dec 21, 1993

250MG

A074216 001 Apr 11, 1996

375MG

A074129 002 Dec 21, 1993

375MG

A074216 002 Apr 11, 1996

500MG

A074129 003 Dec 21, 1993

500MG

A074216 003 Apr 11, 1996

TEVA PHARMS 250MG

A074207 001 Dec 21, 1993

375MG

A074207 002 Dec 21, 1993

500MG

A074207 003 Dec 21, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXEN

TABLET; ORAL

NAPROXEN

WATSON LABS	250MG	A074457 001	May 31, 1995
	375MG	A074457 002	May 31, 1995
	500MG	A074457 003	May 31, 1995
WATSON LABS TEVA	250MG	A074163 001	Feb 10, 1995
	375MG	A074163 002	Feb 10, 1995
	500MG	A074163 003	Feb 10, 1995

TABLET, DELAYED RELEASE; ORAL

NAPROXEN

ACTAVIS ELIZABETH	375MG	A074936 001	Feb 24, 1998
	500MG	A074936 002	Feb 24, 1998
AUROBINDO PHARMA USA	375MG	A075390 001	Apr 19, 2001
	500MG	A075390 002	Apr 19, 2001
FOSUN PHARMA	375MG	A075061 001	Feb 18, 1998
	500MG	A075061 002	Feb 18, 1998
PLIVA	375MG	A075337 001	May 26, 1999
	500MG	A075337 002	May 26, 1999

NAPROXEN SODIUM

CAPSULE; ORAL

NAPROXEN SODIUM

STRIDES PHARMA	EQ 200MG BASE	A215472 001	Aug 17, 2022
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TABLET; ORAL

ANAPROX

+ ATNAHS PHARMA US	EQ 250MG BASE **	N018164 001	
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NAPROXEN SODIUM

ABLE	EQ 250MG BASE	A076544 001	Aug 22, 2003
	EQ 500MG BASE	A076544 002	Aug 22, 2003
CONTRACT PHARMACAL	220MG	A074789 001	Feb 27, 1997
HAMILTON PHARMS	EQ 250MG BASE	A074106 001	Aug 31, 1993
	EQ 500MG BASE	A074106 002	Aug 31, 1993
HIKMA	EQ 250MG BASE	A074480 002	Feb 18, 1998
	EQ 500MG BASE	A074480 001	May 14, 1996
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A074230 001	Mar 14, 1995
	EQ 500MG BASE	A074230 002	Mar 14, 1995
MYLAN	EQ 250MG BASE	A074367 001	Aug 31, 1994
	EQ 500MG BASE	A074367 002	Aug 31, 1994
PLD ACQUISITIONS LLC	220MG	A074646 001	Jan 13, 1997
PLIVA	EQ 250MG BASE	A074242 001	Jun 20, 1996
	EQ 500MG BASE	A074242 002	Jun 20, 1996
PUREPAC PHARM	EQ 250MG BASE	A074319 001	Mar 20, 1995
	EQ 500MG BASE	A074319 002	Mar 20, 1995
ROXANE	EQ 250MG BASE	A074257 001	Dec 21, 1993
	EQ 500MG BASE	A074257 002	Dec 21, 1993
SANDOZ	EQ 250MG BASE	A074162 001	Dec 21, 1993
	EQ 250MG BASE	A074495 001	Dec 05, 1994
	EQ 500MG BASE	A074162 002	Dec 21, 1993
	EQ 500MG BASE	A074495 002	Dec 05, 1994
TEVA	EQ 250MG BASE	A074142 001	Dec 21, 1993
	EQ 250MG BASE	A074198 001	Dec 21, 1993
	EQ 500MG BASE	A074142 002	Dec 21, 1993
	EQ 500MG BASE	A074198 002	Dec 21, 1993
TEVA PHARMS	EQ 250MG BASE	A074289 001	Jan 27, 1994
	EQ 500MG BASE	A074289 002	Jan 27, 1994
WATSON LABS	EQ 250MG BASE	A074195 001	Dec 21, 1993
	EQ 250MG BASE	A074455 001	May 31, 1995
	EQ 500MG BASE	A074195 002	Dec 21, 1993
	EQ 500MG BASE	A074455 002	May 31, 1995

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALEVE-D SINUS & COLD

+ BAYER	220MG;120MG **	N021076 001	Nov 29, 1999
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NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA	220MG;120MG	A211360 001	Jun 01, 2022
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET; ORAL

TREXIMET

+ CURRAX 60MG;EQ 10MG BASE N021926 002 May 14, 2015

NARATRIPTAN HYDROCHLORIDE

TABLET; ORAL

AMERGE

+ GLAXOSMITHKLINE LLC EQ 1MG BASE N020763 002 Feb 10, 1998

+ EQ 2.5MG BASE N020763 001 Feb 10, 1998

NARATRIPTAN

ANI PHARMS EQ 1MG BASE A078751 001 Jul 07, 2010

EQ 2.5MG BASE A078751 002 Jul 07, 2010

APOTEX CORP EQ 1MG BASE A091373 001 Apr 22, 2011

EQ 2.5MG BASE A091373 002 Apr 22, 2011

AUROBINDO PHARMA USA EQ 1MG BASE A202431 001 May 31, 2012

EQ 2.5MG BASE A202431 002 May 31, 2012

CHARTWELL RX EQ 1MG BASE A090288 001 Jul 07, 2010

EQ 2.5MG BASE A090288 002 Jul 07, 2010

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

ALVOGEN 60MG A205055 001 Dec 11, 2015

120MG A205055 002 Dec 11, 2015

TEVA PHARMS 60MG A077467 001 Sep 09, 2009

120MG A077467 002 Sep 09, 2009

STARLIX

+ NOVARTIS 60MG ** N021204 001 Dec 22, 2000

+ 120MG ** N021204 002 Dec 22, 2000

NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

NEBIVOLOL HYDROCHLORIDE

AJANTA PHARMA LTD EQ 2.5MG BASE A213349 001 Mar 31, 2022

EQ 5MG BASE A213349 002 Mar 31, 2022

EQ 10MG BASE A213349 003 Mar 31, 2022

EQ 20MG BASE A213349 004 Mar 31, 2022

ALKEM LABS LTD EQ 2.5MG BASE A203741 001 Jun 24, 2015

EQ 5MG BASE A203741 002 Jun 24, 2015

EQ 10MG BASE A203741 003 Jun 24, 2015

EQ 20MG BASE A203741 004 Jun 24, 2015

GLENMARK PHARMS LTD EQ 2.5MG BASE A203821 001 May 25, 2017

EQ 5MG BASE A203821 002 May 25, 2017

EQ 10MG BASE A203821 003 May 25, 2017

EQ 20MG BASE A203821 004 May 25, 2017

UNICHEM EQ 2.5MG BASE A213830 001 Mar 15, 2022

EQ 5MG BASE A213830 002 Mar 15, 2022

EQ 10MG BASE A213830 003 Mar 15, 2022

EQ 20MG BASE A213830 004 Mar 15, 2022

WATSON LABS INC EQ 2.5MG BASE A203683 001 Nov 27, 2015

EQ 5MG BASE A203683 002 Nov 27, 2015

EQ 10MG BASE A203683 003 Nov 27, 2015

EQ 20MG BASE A203683 004 Nov 27, 2015

NEBIVOLOL HYDROCHLORIDE; VALSARTAN

TABLET; ORAL

BYVALSON

+ ABBVIE EQ 5MG BASE;80MG N206302 001 Jun 03, 2016

NEDOCROMIL SODIUM

AEROSOL, METERED; INHALATION

TILADE

KING PHARMS LLC 1.75MG/INH N019660 001 Dec 30, 1992

SOLUTION; INHALATION

TILADE

SANOFI AVENTIS US 0.5% N020750 001 Oct 01, 1997

SOLUTION/DROPS; OPHTHALMIC

ALOCRIL

+ ALLERGAN 2% N021009 001 Dec 08, 1999

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEDOCROMIL SODIUM

SOLUTION/DROPS;OPHTHALMIC

NEDOCROMIL SODIUM

AKORN

2%

A090638 001 Aug 22, 2012

NEFAZODONE HYDROCHLORIDE

TABLET;ORAL

NEFAZODONE HYDROCHLORIDE

ANI PHARMS

50MG

A076072 001 Sep 16, 2003

100MG

A076072 002 Sep 16, 2003

150MG

A076072 003 Sep 16, 2003

200MG

A076072 004 Sep 16, 2003

250MG

A076072 005 Sep 16, 2003

AUROBINDO PHARMA USA 100MG

A076129 002 Sep 16, 2003

150MG

A076129 003 Sep 16, 2003

200MG

A076129 004 Sep 16, 2003

250MG

A076129 005 Sep 16, 2003

DR REDDYS LABS INC 50MG

A076309 001 Sep 16, 2003

100MG

A076309 002 Sep 16, 2003

150MG

A076309 003 Sep 16, 2003

200MG

A076309 004 Sep 16, 2003

250MG

A076309 005 Sep 16, 2003

FOSUN PHARMA 50MG

A076302 001 Sep 16, 2003

100MG

A076302 002 Sep 16, 2003

150MG

A076302 003 Sep 16, 2003

200MG

A076302 004 Sep 16, 2003

250MG

A076302 005 Sep 16, 2003

IVAX SUB TEVA PHARMS 50MG

A075763 001 Sep 16, 2003

100MG

A075763 002 Sep 16, 2003

150MG

A075763 003 Sep 16, 2003

200MG

A075763 004 Sep 16, 2003

250MG

A075763 005 Sep 16, 2003

ROXANE 50MG

A076196 001 Sep 16, 2003

100MG

A076196 002 Sep 16, 2003

150MG

A076196 003 Sep 16, 2003

200MG

A076196 004 Sep 16, 2003

250MG

A076196 005 Sep 16, 2003

SUN PHARM INDS LTD 50MG

A076409 001 Sep 16, 2003

100MG

A076409 002 Sep 16, 2003

150MG

A076409 003 Sep 16, 2003

200MG

A076409 004 Sep 16, 2003

250MG

A076409 005 Sep 16, 2003

WATSON LABS 100MG

A076073 002 Sep 16, 2003

150MG

A076073 003 Sep 16, 2003

200MG

A076073 004 Sep 16, 2003

250MG

A076073 005 Sep 16, 2003

SERZONE

+ BRISTOL MYERS SQUIBB 50MG **

N020152 001 Dec 22, 1994

+ 100MG **

N020152 002 Dec 22, 1994

+ 150MG **

N020152 003 Dec 22, 1994

+ 200MG **

N020152 004 Dec 22, 1994

+ 250MG **

N020152 005 Dec 22, 1994

+ 300MG **

N020152 006 Dec 22, 1994

NELFINAVIR MESYLATE

POWDER;ORAL

VIRACEPT

AGOURON PHARMS

EQ 50MG BASE/SCOOPFUL

N020778 001 Mar 14, 1997

NEOMYCIN SULFATE

SOLUTION;ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 87.5MG BASE/5ML

N050285 001

NEO-FRADIN

X GEN PHARMS EQ 87.5MG BASE/5ML

A065010 001 May 23, 2002

TABLET;ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 350MG BASE

A060520 001

NEOBIOTIC

PFIZER EQ 350MG BASE

A060475 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

BRISTOL MYERS SQUIBB	500MG	A060365	001	
CHARTWELL MOLECULAR	500MG	A204435	001	Jun 10, 2016
LANNETT	500MG	A060607	001	
LILLY	500MG	A060385	001	
NOSTRUM LABS INC	500MG	A065468	001	Mar 29, 2010
ROXANE	500MG	A062173	001	
SANDOZ	500MG	A061586	001	

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL

NEOSPORIN

GLAXOSMITHKLINE	EQ 3.5MG BASE/GM;10,000 UNITS/GM	N050176	002	Jan 14, 1985
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OINTMENT; OPHTHALMIC

STATROL

ALCON	EQ 3.5MG BASE/GM;10,000 UNITS/GM	N050344	002	
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SOLUTION/DROPS; OPHTHALMIC

STATROL

ALCON	EQ 3.5MG BASE/ML;16,250 UNITS/ML	A062339	001	Nov 30, 1984
	EQ 3.5MG BASE/ML;16,250 UNITS/ML	N050456	001	

NEOMYCIN SULFATE; POLYMYXIN B SULFATE; PREDNISOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

POLY-PRED

ALLERGAN	EQ 0.35% BASE;10,000 UNITS/ML;0.5%	N050081	002	
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NEOMYCIN SULFATE; PREDNISOLONE ACETATE

OINTMENT; OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN	EQ 3.5MG BASE/GM;0.25%	A061039	002	
	EQ 3.5MG BASE/GM;0.5%	A061039	001	

SUSPENSION/DROPS; OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN	EQ 3.5MG BASE/ML;0.25%	A061037	001	
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NEOMYCIN SULFATE; PREDNISOLONE SODIUM PHOSPHATE

OINTMENT; OPHTHALMIC

NEO-HYDELTRASOL

MERCK	EQ 3.5MG BASE/GM;EQ 0.25% PHOSPHATE	N050378	001	
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NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYTRES A

SAVAGE LABS	EQ 3.5MG BASE/GM;0.1%	A062598	001	Jul 21, 1986
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NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA	EQ 3.5MG BASE/GM;0.1%	A062600	001	Jul 21, 1986
PHARMADERM	EQ 3.5MG BASE/GM;0.1%	A062595	001	Jul 21, 1986

OINTMENT; TOPICAL

MYTRES A

SAVAGE LABS	EQ 3.5MG BASE/GM;0.1%	A062609	001	May 23, 1986
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NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA	EQ 3.5MG BASE/GM;0.1%	A062608	001	May 23, 1986
PHARMADERM	EQ 3.5MG BASE/GM;0.1%	A062607	001	May 23, 1986

NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

NEOSTIGMINE METHYLSULFATE

AM REGENT	5MG/10ML (0.5MG/ML)	A209182	001	May 04, 2018
	10MG/10ML (1MG/ML)	A209182	002	May 04, 2018

NESIRITIDE RECOMBINANT

FOR SOLUTION; INTRAVENOUS

NATRECOR

+ SCIOS LLC	1.5MG/VIAL **	N020920	001	Aug 10, 2001
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NETILMICIN SULFATE

INJECTABLE; INJECTION

NETROMYCIN

SCHERING	EQ 10MG BASE/ML	N050544 001	Feb 28, 1983
	EQ 25MG BASE/ML	N050544 002	Feb 28, 1983
	EQ 100MG BASE/ML	N050544 003	Feb 28, 1983

NEVIRAPINE

TABLET; ORAL

NEVIRAPINE

APOTEX INC	200MG	A203021 001	May 22, 2012
MYLAN LABS	200MG	A078864 001	May 22, 2012
PRINSTON INC	200MG	A078644 001	May 22, 2012
TECH ORGANIZED	200MG	A203176 001	May 22, 2012

VIRAMUNE

+ BOEHRINGER INGELHEIM	200MG	N020636 001	Jun 21, 1996
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TABLET, EXTENDED RELEASE; ORAL

NEVIRAPINE

ALVOGEN	100MG	A204621 002	Nov 09, 2015
	400MG	A204621 001	Jul 10, 2015
APOTEX	400MG	A205258 001	Apr 03, 2014
AUROBINDO PHARMA	100MG	A208616 001	Nov 23, 2016
	400MG	A207698 001	Feb 28, 2017
CIPLA	400MG	A206448 001	Oct 15, 2015
MYLAN	100MG	A206271 001	Nov 09, 2015
TECH ORGANIZED	100MG	A207467 001	Jul 31, 2017
	400MG	A207467 002	Jul 31, 2017

NIACIN

CAPSULE; ORAL

WAMPOCAP

MEDPOINTE PHARM HLC	500MG	N011073 003	
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TABLET; ORAL

NIACIN

EVERYLIFE	500MG	A083203 001	
HALSEY	500MG	A083453 001	
HIKMA	500MG	A083718 001	
IMPAX LABS	500MG	A083115 001	
IVAX SUB TEVA PHARMS	500MG	A083180 001	
MK LABS	500MG	A083525 001	
PUREPAC PHARM	500MG	A083271 001	
SANDOZ	500MG	A083306 001	
TABLICAPS	500MG	A084237 001	
WATSON LABS	500MG	A083136 001	
	500MG	A083305 001	
	500MG	A085172 001	
WOCKHARDT	500MG	A081134 001	Apr 28, 1992

NICOLAR

+ SANOFI AVENTIS US	500MG	A083823 001	
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TABLET, EXTENDED RELEASE; ORAL

NIACIN

BEIJING	500MG	A214428 001	Nov 22, 2021
	1GM	A214428 002	Nov 22, 2021
JUBILANT GENERICS	500MG	A209156 001	May 14, 2018
	750MG	A209156 002	May 14, 2018
	1GM	A209156 003	May 14, 2018
RISING	500MG	A203742 001	Feb 22, 2019
	750MG	A203742 002	Feb 22, 2019
	1GM	A203742 003	Feb 22, 2019
YICHANG HUMANWELL	500MG	A212017 001	Jun 10, 2019
	750MG	A212017 002	Jun 10, 2019
	1GM	A212017 003	Jun 10, 2019

NIASPAN

ABBVIE	375MG	N020381 001	Jul 28, 1997
+	500MG **	N020381 002	Jul 28, 1997
+	750MG **	N020381 003	Jul 28, 1997
+	1GM **	N020381 004	Jul 28, 1997

NIASPAN TITRATION STARTER PACK

ABBVIE	375MG; 500MG; 750MG	N020381 005	Jul 28, 1997
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; TYROSINE

SUSPENSION; ORAL

TPN

INTL MINERALS 15MG/5ML; 3.75MG/5ML; 600MG/5ML N008378 003

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

CARDENE

+ CHIESI 20MG ** N019488 001 Dec 21, 1988

+ 30MG ** N019488 002 Dec 21, 1988

NICARDIPINE HYDROCHLORIDE

ANI PHARMS 20MG A074439 001 Dec 10, 1996

20MG A074540 001 Oct 28, 1996

30MG A074439 002 Dec 10, 1996

30MG A074540 002 Oct 28, 1996

MYLAN 20MG A074642 001 Jul 18, 1996

30MG A074642 002 Jul 18, 1996

CAPSULE, EXTENDED RELEASE; ORAL

CARDENE SR

+ CHIESI 30MG ** N020005 001 Feb 21, 1992

+ 45MG ** N020005 002 Feb 21, 1992

+ 60MG ** N020005 003 Feb 21, 1992

INJECTABLE; INJECTION

CARDENE

+ CHIESI 25MG/10ML (2.5MG/ML) ** N019734 001 Jan 30, 1992

NICARDIPINE HYDROCHLORIDE

NAVINTA LLC 25MG/10ML (2.5MG/ML) A090125 001 Nov 17, 2009

RK PHARMA 25MG/10ML (2.5MG/ML) A090664 001 Nov 17, 2009

SUN PHARM 25MG/10ML (2.5MG/ML) N078405 001 Nov 17, 2009

WEST-WARD PHARMS INT 25MG/10ML (2.5MG/ML) A078714 001 Dec 28, 2009

WOCKHARDT 25MG/10ML (2.5MG/ML) A090671 001 Nov 17, 2009

INJECTABLE; INTRAVENOUS

CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER

+ CHIESI 40MG/200ML (0.2MG/ML) N019734 005 Nov 07, 2008

NICLOSAMIDE

TABLET, CHEWABLE; ORAL

NICLOCIDE

BAYER PHARMS 500MG N018669 001 May 14, 1982

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

NICOTROL

MCNEIL CONS 15MG/16HR N020536 001 Jul 03, 1996

PROSTEP

AVEVA 11MG/24HR N019983 003 Dec 23, 1998

22MG/24HR N019983 004 Dec 23, 1998

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICOTINE POLACRILEX

FERTIN PHARMA EQ 4MG BASE A214354 001 Dec 21, 2022

IVAX SUB TEVA PHARMS EQ 2MG BASE A076880 001 Feb 18, 2009

EQ 4MG BASE A077850 001 Feb 18, 2009

L PERRIGO CO EQ 2MG BASE A076776 001 Sep 16, 2004

EQ 4MG BASE A076778 001 Sep 16, 2004

PERRIGO R AND D EQ 2MG BASE A078967 001 Apr 23, 2008

EQ 4MG BASE A078968 001 Apr 23, 2008

THRIVE

GLAXOSMITHKLINE CONS EQ 2MG BASE A077658 001 Jun 19, 2007

EQ 4MG BASE A077656 001 Jun 19, 2007

NIFEDIPINE

CAPSULE; ORAL

ADALAT

BAYER PHARMS 10MG N019478 001 Nov 27, 1985

20MG N019478 002 Sep 17, 1986

NIFEDIPINE

CHASE LABS NJ 10MG A072409 001 Jul 04, 1990

20MG A073421 001 Jun 19, 1991

TEVA 10MG A072651 001 Feb 19, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NIFEDIPINE

CAPSULE;ORAL

PROCARDIA

+ PFIZER 20MG ** N018482 002 Jul 24, 1986

TABLET, EXTENDED RELEASE;ORAL

ADALAT CC

+ NORWICH 30MG ** N020198 001 Apr 21, 1993

+ 60MG ** N020198 002 Apr 21, 1993

+ 90MG ** N020198 003 Apr 21, 1993

AFEDITAB CR

WATSON LABS 60MG A075659 001 Oct 26, 2001

WATSON LABS TEVA 30MG A075128 001 Mar 10, 2000

NIFEDIPINE

AUROBINDO PHARMA USA 30MG A090649 001 Jun 21, 2010

60MG A090649 002 Jun 21, 2010

90MG A090649 003 Jun 21, 2010

MARTEC USA LLC 90MG A075414 003 Mar 23, 2004

MYLAN 30MG A075108 001 Dec 17, 1999

PAR PHARM 30MG A077899 001 Dec 13, 2006

60MG A077899 002 Dec 13, 2006

90MG A077899 003 May 25, 2012

RISING 30MG A090602 001 Sep 13, 2010

30MG A201071 001 Dec 03, 2010

60MG A090602 002 Sep 13, 2010

60MG A201071 002 Dec 03, 2010

90MG A090602 003 Sep 13, 2010

90MG A201071 003 Dec 03, 2010

VITRUVIAS 30MG A216019 001 Nov 18, 2022

60MG A216019 002 Nov 18, 2022

NILUTAMIDE

TABLET;ORAL

NILANDRON

CONCORDIA 50MG N020169 001 Sep 19, 1996

NIMODIPINE

CAPSULE;ORAL

NIMODIPINE

SOFGEN PHARMS 30MG A201832 001 Jul 24, 2015

SUN PHARM INDS INC 30MG A077067 001 Apr 17, 2007

NIMOTOP

+ BAYER PHARMS 30MG ** N018869 001 Dec 28, 1988

SOLUTION;ORAL

NYMALIZE

+ AZURITY 3MG/ML ** N203340 001 May 10, 2013

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

NISOLDIPINE

AMTA 8.5MG A216606 001 Apr 10, 2023

SULAR

+ COVIS 10MG ** N020356 001 Feb 02, 1995

+ 20MG ** N020356 002 Feb 02, 1995

+ 25.5MG ** N020356 006 Jan 02, 2008

+ 30MG ** N020356 003 Feb 02, 1995

+ 40MG ** N020356 004 Feb 02, 1995

NITRIC OXIDE

GAS; INHALATION

INOMAX

+ MALLINCKRODT HOSP 100PPM ** N020845 002 Dec 23, 1999

NITROFURANTOIN

CAPSULE;ORAL

NITROFURANTOIN

WATSON LABS 50MG A084326 001

100MG A084326 002

TABLET;ORAL

FURADANTIN

PROCTER AND GAMBLE 50MG N008693 001

100MG N008693 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROFURANTOIN

TABLET; ORAL

FURALAN

CHARTWELL MOLECULAR	50MG	A080017	001
	100MG	A080017	002

NITROFURANTOIN

ELKINS SINN	50MG	A080003	001
	100MG	A080003	002
IVAX SUB TEVA PHARMS	50MG	A080078	002
	100MG	A080078	001
SANDOZ	50MG	A080043	001
	100MG	A080043	002
WATSON LABS	50MG	A080447	001
	50MG	A085797	001
	100MG	A080447	002
	100MG	A085796	001
WHITEWORTH TOWN PLSN	100MG	A084085	002

NITROFURANTOIN SODIUM

INJECTABLE; INJECTION

IVADANTIN

PROCTER AND GAMBLE	EQ 180MG BASE/VIAL	N012402	001
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NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN

ATHEM	25MG	A074336	001	Jan 25, 1995
	50MG	A074336	002	Jan 25, 1995
	100MG	A074336	003	Jan 25, 1995
WATSON LABS	25MG	A073696	001	Dec 31, 1992
	50MG	A073696	002	Dec 31, 1992
	100MG	A073696	003	Dec 31, 1992

NITROFURANTOIN MACROCRYSTALLINE

WATSON LABS	50MG	A070248	001	Jun 24, 1988
	100MG	A070249	001	Jun 24, 1988

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)

ALVOGEN	75MG; 25MG	A215002	001	Jul 20, 2022
AUROBINDO PHARMA USA	75MG; 25MG	A076648	001	Mar 22, 2004
RANBAXY LABS LTD	75MG; 25MG	A076951	001	Mar 30, 2005

NITROFURAZONE

CREAM; TOPICAL

FURACIN

SHIRE	0.2%	A083789	001
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DRESSING; TOPICAL

ACTIN-N

SHERWOOD MEDCL	0.2%	N017343	001
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OINTMENT; TOPICAL

FURACIN

SHIRE	0.2%	N005795	001
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NITROFURAZONE

AMBIX	0.2%	A086077	001
LANNETT	0.2%	A084393	001
PERRIGO NEW YORK	0.2%	A084968	001
TARO	0.2%	A086156	001
WENDT	0.2%	A086766	001

POWDER; TOPICAL

FURACIN

SHIRE	0.2%	A083791	001
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SOLUTION; TOPICAL

NITROFURAZONE

PERRIGO NEW YORK	0.2%	A085130	001
WENDT	0.2%	A087081	001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROGLYCERIN

AEROSOL;SUBLINGUAL

NITROLINGUAL

POHL BOSKAMP 0.4MG/SPRAY

N018705 001 Oct 31, 1985

FILM, EXTENDED RELEASE;TRANSDERMAL

MINITRAN

BAUSCH 0.4MG/HR

A089773 001 Aug 30, 1996

VALEANT PHARMS 0.1MG/HR

A089771 001 Aug 30, 1996

0.6MG/HR

A089774 001 Aug 30, 1996

VALEANT PHARMS NORTH 0.2MG/HR

A089772 001 Aug 30, 1996

NITROGLYCERIN

LANNETT CO INC 0.2MG/HR

A075115 001 Aug 10, 2004

0.4MG/HR

A075115 002 Aug 10, 2004

MYLAN TECHNOLOGIES 0.1MG/HR

A074992 004 Nov 12, 1999

0.2MG/HR

A074992 003 Nov 12, 1999

0.4MG/HR

A074992 002 Nov 12, 1999

0.6MG/HR

A074992 001 Nov 12, 1999

TRANSDERM-NITRO

+ NOVARTIS 0.1MG/HR **

N020144 001 Feb 27, 1996

+ 0.2MG/HR **

N020144 002 Feb 27, 1996

+ 0.4MG/HR **

N020144 003 Feb 27, 1996

+ 0.6MG/HR **

N020144 004 Feb 27, 1996

+ 0.8MG/HR **

N020144 005 Feb 27, 1996

INJECTABLE;INJECTION

NITRO IV

POHL BOSKAMP 5MG/ML

N018672 002 Aug 30, 1983

NITRO-BID

SANOFI AVENTIS US 5MG/ML

N018621 001 Jan 05, 1982

10MG/ML

A071159 001 Feb 28, 1990

NITROGLYCERIN

ABRAXIS PHARM 5MG/ML

A070077 001 Dec 13, 1985

5MG/ML

A071203 001 May 08, 1987

+ HOSPIRA 5MG/ML **

N018531 001

INTL MEDICATION 5MG/ML

A070026 001 Sep 10, 1985

LUITPOLD 5MG/ML

A071492 001 May 24, 1988

SMITH AND NEPHEW 5MG/ML

A070633 001 Jun 19, 1986

5MG/ML

A070634 001 Jun 19, 1986

NITROGLYCERIN IN DEXTROSE 5%

HOSPIRA 0.1MG/ML

A074083 001 Oct 26, 1994

10MG/100ML

A071846 001 Aug 31, 1990

20MG/100ML

A071847 001 Aug 31, 1990

40MG/100ML

A071848 001 Aug 31, 1990

NITROL

RORER 0.8MG/ML

N018774 001 Jan 19, 1983

NITRONAL

POHL BOSKAMP 1MG/ML

N018672 001 Aug 30, 1983

NITROSTAT

PARKE DAVIS 0.8MG/ML

N018588 001

5MG/ML

A070863 001 Jan 08, 1987

5MG/ML

N018588 002 Dec 23, 1983

10MG/ML

A070871 001 Jan 08, 1987

10MG/ML

A070872 001 Jan 08, 1987

TRIDIL

HOSPIRA 0.5MG/ML

N018537 002 Jun 16, 1983

5MG/ML

N018537 001

POWDER;SUBLINGUAL

GONITRO

+ POHL BOSKAMP 0.4MG/PACKET

N208424 001 Jun 08, 2016

TABLET;SUBLINGUAL

NITROGLYCERIN

ACTAVIS LABS FL INC 0.3MG

A203693 001 Oct 16, 2017

0.4MG

A203693 002 Oct 16, 2017

0.6MG

A203693 003 Oct 16, 2017

SIGMAPHARM LABS LLC 0.3MG

A207745 001 May 07, 2018

0.4MG

A207745 002 May 07, 2018

0.6MG

A207745 003 May 07, 2018

ZYDUS PHARMS 0.3MG

A210153 001 Mar 08, 2022

0.4MG

A210153 002 Mar 08, 2022

0.6MG

A210153 003 Mar 08, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NIZATIDINECAPSULE; ORAL
AXID

SMITHKLINE BEECHAM	150MG	N019508 001	Apr 12, 1988
	300MG	N019508 002	Apr 12, 1988

NIZATIDINE

ANI PHARMS	150MG	A075461 001	Jul 08, 2002
	150MG	A075668 001	Sep 12, 2002
	300MG	A075461 002	Jul 08, 2002
	300MG	A075668 002	Sep 12, 2002
APOTEX INC	150MG	A076383 001	Jan 23, 2003
	300MG	A076383 002	Jan 23, 2003
MYLAN PHARMS INC	150MG	A075806 001	Jul 05, 2002
	150MG	A075934 001	Jul 09, 2002
	300MG	A075806 002	Jul 05, 2002
	300MG	A075934 002	Jul 09, 2002

SOLUTION; ORAL

AXID

+ BRAINTREE	15MG/ML **	N021494 001	May 25, 2004
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NIZATIDINE

AMNEAL PHARMS	15MG/ML	A090576 001	Nov 18, 2009
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NONOXYNOL-9

AEROSOL; VAGINAL

DELFIN

PERSONAL PRODS	12.5%	N014349 002	
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SPONGE; VAGINAL

TODAY

+ MAYER LABS INC	1GM	N018683 001	Apr 01, 1983
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NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

METRICS PHARM	EQ 1MG BASE/ML	A040522 001	Sep 30, 2004
ZYDUS PHARMS	EQ 1MG BASE/ML	A216341 001	Jul 19, 2022

SOLUTION; INTRAVENOUS

NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE

+ LONG GROVE PHARMS	EQ 4MG BASE/250 ML (EQ 16MCG BASE/ML)	N214628 001	Oct 06, 2022
	EQ 8MG BASE/250ML (EQ 32MCG BASE/ML)	N214628 002	Oct 06, 2022
	EQ 16MG BASE/250ML (EQ 64MCG BASE/ML)	N214628 003	Oct 06, 2022

NOREPINEPHRINE BITARTRATE; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAINE AND NOVOCAIN W/ LEVOPHED

EASTMAN KODAK	EQ 0.033MG BASE/ML; 2%; 0.4%	N008592 003	
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NORETHINDRONE

TABLET; ORAL

NORLUTIN

PARKE DAVIS	5MG	N010895 002	
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TABLET; ORAL-28

MICRONOR

+ JANSSEN PHARMS	0.35MG **	N016954 001	
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NORETHINDRONE

AMNEAL PHARMS	0.35MG	A202260 001	Aug 01, 2013
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NORETHINDRONE ACETATE

TABLET; ORAL

AYGESTIN

+ DURAMED RES	5MG **	N018405 001	Apr 21, 1982
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NORETHINDRONE ACETATE

AUROBINDO PHARMA LTD	5MG	A204236 001	Jan 08, 2016
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NORLUTATE

PARKE DAVIS	5MG	N012184 002	
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NORFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

CHIBROXIN

MERCK	0.3%	N019757 001	Jun 17, 1991
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TABLET; ORAL

NOROXIN

+ MERCK	400MG **	N019384 002	Oct 31, 1986
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NORGESTREL

TABLET; ORAL

OPILL

+ LABORATOIRE HRA 0.075MG N017031 001

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

AVENTYL HYDROCHLORIDE

LILLY EQ 10MG BASE N014684 001

EQ 25MG BASE N014684 002

NORTRIPTYLINE HYDROCHLORIDE

ANI PHARMS

EQ 10MG BASE

A074054 001

Dec 31, 1992

EQ 25MG BASE

A074054 002

Dec 31, 1992

EQ 50MG BASE

A074054 003

Dec 31, 1992

EQ 75MG BASE

A074054 004

Dec 31, 1992

AUROBINDO PHARMA LTD

EQ 10MG BASE

A074835 001

Jun 30, 1997

EQ 25MG BASE

A074835 002

Jun 30, 1997

EQ 50MG BASE

A074835 003

Jun 30, 1997

EQ 75MG BASE

A074835 004

Jun 30, 1997

MYLAN

EQ 10MG BASE

A074234 001

Jul 26, 1993

EQ 25MG BASE

A074234 002

Jul 26, 1993

EQ 50MG BASE

A074234 003

Jul 26, 1993

EQ 75MG BASE

A074234 004

Jul 26, 1993

TEVA

EQ 10MG BASE

A073667 001

Apr 11, 1996

EQ 25MG BASE

A073667 002

Apr 11, 1996

EQ 50MG BASE

A073667 003

Apr 11, 1996

EQ 75MG BASE

A073667 004

Apr 11, 1996

ZYDUS

EQ 10MG BASE

A213441 001

Feb 24, 2021

EQ 25MG BASE

A213441 002

Feb 24, 2021

EQ 50MG BASE

A213441 003

Feb 24, 2021

EQ 75MG BASE

A213441 004

Feb 24, 2021

SOLUTION; ORAL

AVENTYL

+ RANBAXY

EQ 10MG BASE/5ML **

N014685 001

NORTRIPTYLINE HYDROCHLORIDE

TARO

EQ 10MG BASE/5ML

A077965 001

Jun 20, 2006

PAMELOR

SPECGX LLC

EQ 10MG BASE/5ML

N018012 001

NYSTATIN

CREAM; TOPICAL

CANDEX

BAYER PHARMS

100,000 UNITS/GM

A061810 001

MYCOSTATIN

DELacor ASSET CORP

100,000 UNITS/GM **

A060575 001

MYKINAC

ALPHARMA US PHARMS

100,000 UNITS/GM

A062387 001

Jul 29, 1982

NILSTAT

LEDERLE

100,000 UNITS/GM

A061445 001

NYSTATIN

STRIDES PHARMA

100,000 UNITS/GM

A065315 001

May 31, 2006

TARO

100,000 UNITS/GM

A062457 001

Jul 28, 1983

LOTION; TOPICAL

CANDEX

BAYER PHARMS

100,000 UNITS/ML

N050233 001

OINTMENT; TOPICAL

MYCOSTATIN

DELacor ASSET CORP

100,000 UNITS/GM **

A060571 001

MYKINAC

ALPHARMA US PHARMS

100,000 UNITS/GM

A062731 001

Sep 22, 1986

NILSTAT

LEDERLE

100,000 UNITS/GM

A061444 001

NYSTATIN

TORRENT

100,000 UNITS/GM

A211838 001

Jan 28, 2019

PASTILLE; ORAL

MYCOSTATIN

DELacor ASSET CORP

200,000 UNITS

N050619 001

Apr 09, 1987

POWDER; ORAL

BARSTATIN 100

BARLAN

100%

A062489 001

Apr 27, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN

POWDER; ORAL

NILSTAT

+ STRIDES PHARMA 100% ** N050576 001 Dec 22, 1983

NYSTATIN

PADDOCK LLC 100% A062613 001 Nov 26, 1985

POWDER; TOPICAL

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/GM ** A060578 001

NYSTATIN

NESHER PHARMS 100,000 UNITS/GM A065321 001 Aug 18, 2006

SUPPOSITORY; VAGINAL

NYSERT

WARNER CHILCOTT 100,000 UNITS N050478 001

SUSPENSION; ORAL

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/ML A061533 001

NILSTAT

+ CHARTWELL MOLECULES 100,000 UNITS/ML ** N050299 001

NYSTATIN

AKORN 100,000 UNITS/ML A064042 001 Feb 28, 1994

ALPHARMA US PHARMS 100,000 UNITS/ML A062571 001 Oct 29, 1985

COSETTE 100,000 UNITS/ML A062776 001 Dec 17, 1987

G AND W LABS INC 100,000 UNITS/ML A062349 001 Jul 14, 1982

MORTON GROVE 100,000 UNITS/ML A062835 001 Nov 19, 1987

PHARMADERM 100,000 UNITS/ML A062518 001 Jul 06, 1984

PHARMAFAIR 100,000 UNITS/ML A062541 001 Jan 16, 1985

TEVA 100,000 UNITS/ML A062670 001 Jun 18, 1987

VISTAPHARM 100,000 UNITS/ML A064142 001 Jun 25, 1998

100,000 UNITS/ML A064142 002 Mar 07, 2011

NYSTEX

SAVAGE LABS 100,000 UNITS/ML A062519 001 Jul 06, 1984

TABLET; ORAL

MYCOSTATIN

DELCOR ASSET CORP 500,000 UNITS A060574 001

NILSTAT

LEDERLE 500,000 UNITS A061151 001

NYSTATIN

CHARTWELL RX 500,000 UNITS A062524 001 Nov 26, 1985

QUANTUM PHARMICS 500,000 UNITS A062525 001 Oct 29, 1984

SANDOZ 500,000 UNITS A062065 001

WATSON LABS 500,000 UNITS A062402 001 Dec 16, 1982

TABLET; VAGINAL

KOROSTATIN

HOLLAND RANTOS 100,000 UNITS A061718 001

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS A060577 001

NILSTAT

LEDERLE 100,000 UNITS A061325 001

NYSTATIN

FOUGERA 100,000 UNITS A062459 001 Nov 09, 1983

ODYSSEY PHARMS 100,000 UNITS A062615 001 Oct 17, 1985

PHARMADERM 100,000 UNITS A062460 001 Nov 09, 1983

QUANTUM PHARMICS 100,000 UNITS A062509 001 Apr 03, 1984

SANDOZ 100,000 UNITS A061965 001

TEVA 100,000 UNITS A062502 001 Dec 23, 1983

WATSON LABS 100,000 UNITS A062176 001

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYCO-TRIA CET II

TEVA 100,000 UNITS/GM; 0.1% A061954 002 Sep 20, 1985

MYCOLOG-II

DELCOR ASSET CORP 100,000 UNITS/GM; 0.1% ** A060576 002 May 01, 1985

MYLAN 100,000 UNITS/GM; 0.1% ** A062606 001 May 15, 1985

MYTRES F

SAVAGE LABS 100,000 UNITS/GM; 0.1% A062597 001 Oct 08, 1985

NYSTATIN AND TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS 100,000 UNITS/GM; 0.1% A063010 001 Dec 20, 1988

PERRIGO NEW YORK 100,000 UNITS/GM; 0.1% A062186 002 Jun 06, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

NYSTATIN AND TRIAMCINOLONE ACETONIDE

PHARMAFAIR	100,000 UNITS/GM;0.1%	A062657	001	Jul 30, 1986
TARO	100,000 UNITS/GM;0.1%	A062347	001	Mar 30, 1987
TORRENT	100,000 UNITS/GM;0.1%	A213142	001	Jul 14, 2020

NYSTATIN TRIAMCINOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM;0.1%	A062596	001	Oct 08, 1985
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OINTMENT; TOPICAL

MYCO-TRIA CET II

TEVA	100,000 UNITS/GM;0.1%	A062045	002	Nov 26, 1985
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MYCOLOG-II

MYLAN	100,000 UNITS/GM;0.1% **	A060572	001	Jun 28, 1985
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MYTRES F

SAVAGE LABS	100,000 UNITS/GM;0.1%	A062601	001	Oct 09, 1985
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NYSTATIN AND TRIAMCINOLONE ACETONIDE

CHARTWELL RX	100,000 UNITS/GM;0.1%	A207316	001	Nov 18, 2019
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CROWN LABS INC	100,000 UNITS/GM;0.1%	A207731	001	Dec 26, 2017
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PAI HOLDINGS PHARM	100,000 UNITS/GM;0.1%	A208287	001	Dec 30, 2016
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PERRIGO NEW YORK	100,000 UNITS/GM;0.1%	A062280	002	Oct 10, 1985
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PHARMAFAIR	100,000 UNITS/GM;0.1%	A062656	001	Jul 30, 1986
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NYSTATIN-TRIAMCINOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM;0.1%	A062603	001	Oct 09, 1985
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OBETICHOLIC ACID

TABLET; ORAL

OBETICHOLIC ACID

APOTEX	5MG	A214862	001	May 30, 2023
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	10MG	A214862	002	May 30, 2023
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LUPIN LTD	5MG	A214980	001	May 30, 2023
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	10MG	A214980	002	May 30, 2023
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MSN	5MG	A215017	001	May 30, 2023
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	10MG	A215017	002	May 30, 2023
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OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

SUN PHARM INDS	EQ 0.05MG BASE/ML	A077329	001	Mar 04, 2008
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	EQ 0.05MG BASE/ML	A077372	001	Aug 14, 2007
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	EQ 0.1MG BASE/ML	A077329	002	Mar 04, 2008
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	EQ 0.1MG BASE/ML	A077372	002	Aug 14, 2007
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	EQ 0.2MG BASE/ML	A077330	001	Mar 04, 2008
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	EQ 0.2MG BASE/ML	A077373	001	Aug 14, 2007
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	EQ 0.5MG BASE/ML	A077329	003	Mar 04, 2008
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	EQ 0.5MG BASE/ML	A077372	003	Aug 14, 2007
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	EQ 1MG BASE/ML	A077331	001	Mar 04, 2008
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	EQ 1MG BASE/ML	A077373	002	Aug 14, 2007
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WOCKHARDT USA	EQ 0.2MG BASE/ML	A090986	001	May 11, 2011
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	EQ 1MG BASE/ML	A090986	002	May 11, 2011
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OCTREOTIDE ACETATE PRESERVATIVE FREE

WOCKHARDT USA	EQ 0.05MG BASE/ML	A090985	001	May 11, 2011
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	EQ 0.1MG BASE/ML	A090985	002	May 11, 2011
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	EQ 0.5MG BASE/ML	A090985	003	May 11, 2011
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SANDOSTATIN

+ NOVARTIS	EQ 0.2MG BASE/ML **	N019667	004	Jun 12, 1991
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+	EQ 1MG BASE/ML **	N019667	005	Jun 12, 1991
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OFLOXACIN

INJECTABLE; INJECTION

FLOXIN

ORTHO MCNEIL PHARM	20MG/ML	N020087	002	Mar 31, 1992
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	40MG/ML	N020087	003	Mar 31, 1992
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FLOXIN IN DEXTROSE 5%

ORTHO MCNEIL PHARM	400MG/100ML	N020087	001	Mar 31, 1992
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FLOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

ORTHO MCNEIL PHARM	4MG/ML	N020087	004	Mar 31, 1992
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	400MG/100ML	N020087	005	Mar 31, 1992
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OFLOXACIN

BEDFORD	40MG/ML	A075762	001	Jan 16, 2002
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OFLOXACIN

SOLUTION/DROPS;OPHTHALMIC

OFLOXACIN

AKORN	0.3%	A076615	001	May 14, 2004
ALVOGEN	0.3%	A076830	001	Aug 31, 2004
SANDOZ	0.3%	A076231	001	May 14, 2004
	0.3%	A076848	001	Nov 25, 2008

SOLUTION/DROPS;OTIC

FLOXIN OTIC

+ DAIICHI	0.3% **	N020799	001	Dec 16, 1997
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OFLOXACIN

ALVOGEN	0.3%	A090395	001	Aug 11, 2009
FDC LTD	0.3%	A215038	001	Jan 19, 2022
SANDOZ	0.3%	A078222	001	Mar 17, 2008

TABLET;ORAL

FLOXIN

+ JANSSEN PHARMS	200MG **	N019735	001	Dec 28, 1990
	300MG **	N019735	002	Dec 28, 1990
	400MG **	N019735	003	Dec 28, 1990

OFLOXACIN

LARKEN LABS	200MG	A076093	001	Sep 02, 2003
	300MG	A076093	002	Sep 02, 2003
RANBAXY LABS LTD	200MG	A076220	001	Sep 02, 2003
	300MG	A076220	002	Sep 02, 2003
	400MG	A076220	003	Sep 02, 2003

OLANZAPINE

TABLET;ORAL

OLANZAPINE

AJANTA PHARMA LTD	2.5MG	A206711	001	Aug 30, 2016
	5MG	A206711	002	Aug 30, 2016
	7.5MG	A206711	003	Aug 30, 2016
	10MG	A206711	004	Aug 30, 2016
	15MG	A206711	005	Aug 30, 2016
	20MG	A206711	006	Aug 30, 2016
HIKMA	2.5MG	A204866	001	Jun 16, 2017
	5MG	A204866	002	Jun 16, 2017
	7.5MG	A204866	003	Jun 16, 2017
	10MG	A204866	004	Jun 16, 2017
	15MG	A204866	005	Jun 16, 2017
	20MG	A204866	006	Jun 16, 2017
HISUN PHARM HANGZHOU	2.5MG	A206924	001	Dec 31, 2020
	5MG	A206924	002	Dec 31, 2020
	7.5MG	A206924	003	Dec 31, 2020
	10MG	A206924	004	Dec 31, 2020
	15MG	A206924	005	Dec 31, 2020
	20MG	A206924	006	Dec 31, 2020
IVAX PHARMS INC	20MG	A077301	001	Apr 29, 2015
JIANGSU HANSOH PHARM	2.5MG	A209399	001	Sep 24, 2018
	5MG	A209399	002	Sep 24, 2018
	10MG	A209399	003	Sep 24, 2018
MYLAN	2.5MG	A076866	001	Apr 23, 2012
	5MG	A076866	002	Apr 23, 2012
	7.5MG	A076866	003	Apr 23, 2012
	10MG	A076866	004	Apr 23, 2012
	15MG	A076866	005	Apr 23, 2012
	20MG	A076866	006	Apr 23, 2012
SUNSHINE	2.5MG	A206238	001	Nov 19, 2018
	5MG	A206238	002	Nov 19, 2018
	7.5MG	A206238	003	Nov 19, 2018
	10MG	A206238	004	Nov 19, 2018
	15MG	A206238	005	Nov 19, 2018
	20MG	A206238	006	Nov 19, 2018
TEVA PHARMS	2.5MG	A076000	001	Oct 24, 2011
	5MG	A076000	002	Oct 24, 2011
	7.5MG	A076000	003	Oct 24, 2011
	10MG	A076000	004	Oct 24, 2011
	15MG	A076000	005	Oct 24, 2011
TORRENT PHARMS LTD	2.5MG	A091434	001	Apr 23, 2012
	5MG	A091434	002	Apr 23, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OLANZAPINETABLET; ORAL
OLANZAPINE

7.5MG	A091434	003	Apr 23, 2012
10MG	A091434	004	Apr 23, 2012
15MG	A091434	005	Apr 23, 2012
20MG	A091434	006	Apr 23, 2012

TABLET, ORALLY DISINTEGRATING; ORAL
OLANZAPINE

AJANTA PHARMA LTD	5MG	A204320	001	May 30, 2017
	10MG	A204320	002	May 30, 2017
	15MG	A204320	003	May 30, 2017
	20MG	A204320	004	May 30, 2017
ZYDUS PHARMS	5MG	A202889	001	Mar 09, 2023
	10MG	A202889	002	Mar 09, 2023
	15MG	A202889	003	Mar 09, 2023
	20MG	A202889	004	Mar 09, 2023

OLAPARIB

CAPSULE; ORAL

LYNPARZA

+ ASTRAZENECA 50MG

N206162 001 Dec 19, 2014

OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL

AMNEAL	5MG	A207480	001	Jul 11, 2023
	20MG	A207480	002	Jul 11, 2023
	40MG	A207480	003	Jul 11, 2023
INVENTIA	5MG	A208659	001	May 18, 2020
	20MG	A208659	002	May 18, 2020
	40MG	A208659	003	May 18, 2020
JUBILANT GENERICS	5MG	A205482	001	Apr 24, 2017
	20MG	A205482	002	Apr 24, 2017
	40MG	A205482	003	Apr 24, 2017
LUPIN LTD	5MG	A206631	001	Apr 27, 2017
	20MG	A206631	002	Apr 27, 2017
	40MG	A206631	003	Apr 27, 2017
RISING	5MG	A078276	001	Oct 26, 2016
	20MG	A078276	002	Oct 26, 2016
	40MG	A078276	003	Oct 26, 2016
SANDOZ	5MG	A090237	001	Apr 13, 2020
	20MG	A090237	002	Apr 13, 2020
	40MG	A090237	003	Apr 13, 2020
TEVA PHARMS USA	5MG	A091079	001	Apr 24, 2017
	20MG	A091079	002	Apr 24, 2017
	40MG	A091079	003	Apr 24, 2017
TORRENT	5MG	A202375	001	Apr 24, 2017
	20MG	A202375	002	Apr 24, 2017
	40MG	A202375	003	Apr 24, 2017

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

AKORN	EQ 0.1% BASE	A204532	001	Jan 10, 2017
	EQ 0.2% BASE	A204723	001	Dec 05, 2017
BAUSCH AND LOMB INC	EQ 0.1% BASE	A206046	001	Jul 26, 2017
	EQ 0.2% BASE	A206087	001	Dec 05, 2017
FDC LTD	EQ 0.1% BASE	A209282	001	Sep 26, 2019
FLORIDA	EQ 0.7% BASE	A208637	001	Feb 19, 2020
GLENMARK PHARMS INC	EQ 0.1% BASE	A200810	001	Jun 28, 2017
RISING	EQ 0.1% BASE	A204392	001	Mar 21, 2018
	EQ 0.2% BASE	A204620	001	Jun 16, 2020
ZAMBON SPA	EQ 0.1% BASE	A204706	001	Dec 07, 2015

SPRAY, METERED; NASAL

PATANASE

+ NOVARTIS 0.665MG/SPRAY

N021861 001 Apr 15, 2008

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET;ORAL

TECHNIVIE

+ ABBVIE

12.5MG;75MG;50MG **

N207931 001 Jul 24, 2015

OMEGA-3-ACID ETHYL ESTERS

CAPSULE;ORAL

OMEGA-3-ACID ETHYL ESTERS

STRIDES PHARMA

1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

A091018 001 Jun 24, 2014

ZYDUS

1GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

A210107 001 Jun 14, 2019

OMEGA-3-ACID ETHYL ESTERS TYPE A

CAPSULE;ORAL

OMTRYG

+ OSMOTICA PHARM US

1.2GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

N204977 001 Apr 23, 2014

OMEGA-3-CARBOXYLIC ACIDS

CAPSULE;ORAL

EPANOVA

+ ASTRAZENECA

1GM CONTAINS AT LEAST 850MG OF
POLYUNSATURATED FATTY ACIDS

N205060 001 May 05, 2014

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

LUPIN LTD

40MG

A202384 001 Aug 25, 2015

MYLAN

10MG

A205070 001 Jun 29, 2018

20MG

A205070 002 Jun 29, 2018

40MG

A205070 003 Jun 29, 2018

STRIDES PHARMA

10MG

A075876 001 May 29, 2003

20MG

A075876 002 May 29, 2003

40MG

A075876 003 Jan 21, 2009

TEVA PHARMS USA

40MG

A204661 002 Jun 13, 2017

PRILOSEC

+ ASTRAZENECA

10MG **

N019810 003 Oct 05, 1995

+

20MG **

N019810 001 Sep 14, 1989

+

40MG **

N019810 002 Jan 15, 1998

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA LTD EQ 20MG BASE

A213201 001 Apr 28, 2023

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

PERRIGO R AND D EQ 20MG BASE

A204152 001 Jul 30, 2015

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

CHARTWELL RX 20MG;1.1GM

A204137 001 Jul 15, 2016

STRIDES PHARMA 20MG;1.1GM

A078966 001 May 25, 2010

20MG;1.1GM

A201946 001 Jul 15, 2016

40MG;1.1GM

A078966 002 May 25, 2010

OMIDENEPAG ISOPROPYL

SOLUTION;OPHTHALMIC

OMLONTI

+ VISIOX PHARMA

0.002%

N215092 001 Sep 22, 2022

ONDANSETRON

FILM;ORAL

ZUPLENZ

+ AQUESTIVE

4MG

N022524 001 Jul 02, 2010

+

8MG

N022524 002 Jul 02, 2010

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

BARR

4MG

A076693 001 Jun 25, 2007

8MG

A076693 002 Jun 25, 2007

NESHER PHARMS

4MG

A077717 001 Jun 25, 2007

8MG

A077717 002 Jun 25, 2007

TEVA

4MG

A076810 001 Jun 25, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

	8MG	A076810	002	Jun 25, 2007
ZOFRAN ODT				
+ SANDOZ	4MG **	N020781	001	Jan 27, 1999
+	8MG **	N020781	002	Jan 27, 1999

ONDANSETRON HYDROCHLORIDE

INJECTABLE;INJECTION

ONDANSETRON HYDROCHLORIDE

APOTEX INC	EQ 2MG BASE/ML	A077368	001	Dec 26, 2006
AVET LIFESCIENCES	EQ 2MG BASE/ML	A090424	001	Apr 16, 2010
BAXTER HLTHCARE CORP	EQ 2MG BASE/ML	A078288	001	Feb 22, 2013
CHARTWELL MOLECULAR	EQ 2MG BASE/ML	A090116	001	Apr 14, 2010
EUGIA PHARMA	EQ 2MG BASE/ML	A202599	001	Dec 21, 2012
HOSPIRA	EQ 2MG BASE/ML	A076695	001	Dec 26, 2006
	EQ 2MG BASE/ML	A077840	001	Jan 19, 2007
LANNETT CO INC	EQ 2MG BASE/ML	A090883	001	Aug 05, 2010
LUITPOLD	EQ 2MG BASE/ML	A077582	001	Dec 26, 2006
	EQ 2MG BASE/ML	A079039	001	Nov 18, 2008
PLIVA HRVATSKA DOO	EQ 2MG BASE/ML	A077544	001	Dec 26, 2006
RISING	EQ 2MG BASE/ML	A204906	001	Jul 31, 2017
SAGENT PHARMS	EQ 2MG BASE/ML	A078180	001	Mar 26, 2007
STERISCIENCE	EQ 2MG BASE/ML	A078257	001	Apr 23, 2008
SUN PHARM INDS (IN)	EQ 2MG BASE/ML	A077172	001	Dec 26, 2006
TEVA	EQ 2MG BASE/ML	A076876	001	Nov 22, 2006

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

HOSPIRA	EQ 0.64MG BASE/ML	A076978	001	Feb 26, 2007
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ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

AM REGENT	EQ 2MG BASE/ML	A079032	001	Nov 18, 2008
APOTEX INC	EQ 2MG BASE/ML	A077343	001	Dec 26, 2006
AVET LIFESCIENCES	EQ 2MG BASE/ML	A078945	001	Jan 03, 2013
EUGIA PHARMA	EQ 2MG BASE/ML	A202600	001	Dec 21, 2012
HIKMA FARMACEUTICA	EQ 2MG BASE/ML	A076780	001	Dec 26, 2006
HOSPIRA	EQ 2MG BASE/ML	A076696	001	Dec 26, 2006
LUITPOLD	EQ 2MG BASE/ML	A077387	001	Dec 26, 2006
STERISCIENCE	EQ 2MG BASE/ML	A078244	001	Apr 23, 2008
SUN PHARM INDS LTD	EQ 2MG BASE/ML	A077173	001	Dec 26, 2006
TARO PHARMS IRELAND	EQ 2MG BASE/ML	A078014	001	Mar 21, 2008
TEVA	EQ 2MG BASE/ML	A076759	001	Nov 22, 2006

ZOFRAN

+ SANDOZ	EQ 2MG BASE/ML **	N020007	001	Jan 04, 1991
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ZOFRAN AND DEXTROSE IN PLASTIC CONTAINER

+ GLAXOSMITHKLINE	EQ 0.64MG BASE/ML **	N020403	001	Jan 31, 1995
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ZOFRAN PRESERVATIVE FREE

+ SANDOZ	EQ 2MG BASE/ML **	N020007	003	Dec 10, 1993
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SOLUTION;ORAL

ZOFRAN

+ SANDOZ	EQ 4MG BASE/5ML	N020605	001	Jan 24, 1997
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TABLET;ORAL

ONDANSETRON HYDROCHLORIDE

AUROBINDO PHARMA	EQ 24MG BASE	A078539	003	Jul 31, 2007
CASI PHARMS INC	EQ 4MG BASE	A077517	001	Jun 25, 2007
	EQ 8MG BASE	A077517	002	Jun 25, 2007
	EQ 24MG BASE	A077517	003	Jun 25, 2007
DR REDDYS LABS LTD	EQ 16MG BASE	A076183	004	Dec 26, 2006
	EQ 24MG BASE	A076183	001	Dec 26, 2006
GLENMARK GENERICS	EQ 24MG BASE	A077535	003	Jun 25, 2007
HIKMA INTL PHARMS	EQ 4MG BASE	A077545	001	Sep 06, 2007
	EQ 8MG BASE	A077545	002	Sep 06, 2007
	EQ 24MG BASE	A077545	003	Sep 06, 2007
PLIVA HRVATSKA DOO	EQ 4MG BASE	A077112	001	Jun 25, 2007
	EQ 8MG BASE	A077112	002	Jun 25, 2007
	EQ 24MG BASE	A077112	003	Jun 25, 2007
RISING	EQ 4MG BASE	A076930	001	Jun 25, 2007
	EQ 8MG BASE	A076930	002	Jun 25, 2007
	EQ 24MG BASE	A076930	004	Jun 25, 2007
TARO	EQ 4MG BASE	A077729	001	Mar 28, 2011
	EQ 8MG BASE	A077729	002	Mar 28, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ONDANSETRON HYDROCHLORIDE

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

	EQ 24MG BASE	A077729 003	Mar 28, 2011
TEVA	EQ 4MG BASE	A076252 001	Jun 25, 2007
	EQ 8MG BASE	A076252 002	Jun 25, 2007
	EQ 24MG BASE	A076252 003	Jun 25, 2007
ZOFRAN			
+ SANDOZ	EQ 4MG BASE **	N020103 001	Dec 31, 1992
+	EQ 8MG BASE **	N020103 002	Dec 31, 1992
+	EQ 24MG BASE **	N020103 003	Aug 27, 1999

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

NORFLEX

+ PAI HOLDINGS PHARM	30MG/ML **	N013055 001	
ORPHENADRINE CITRATE			
WATSON LABS	30MG/ML	A087062 001	
TABLET, EXTENDED RELEASE; ORAL			
NORFLEX			
+ BAUSCH	100MG **	N012157 001	
ORPHENADRINE CITRATE			
ASCOT	100MG	A088067 001	Apr 06, 1983
IMPAX PHARMS	100MG	A040368 001	Jun 23, 2000
SANDOZ	100MG	A085046 001	
WATSON LABS	100MG	A084303 001	

ORPHENADRINE HYDROCHLORIDE

TABLET; ORAL

DISIPAL

3M 50MG N010653 001

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATE

ACCORD HLTHCARE	EQ 30MG BASE	A214726 001	Jul 25, 2022
	EQ 45MG BASE	A214726 002	Jul 25, 2022
	EQ 75MG BASE	A214726 003	Jul 25, 2022
RISING	EQ 30MG BASE	A210157 001	Jan 21, 2021
	EQ 45MG BASE	A210157 002	Jan 21, 2021
	EQ 75MG BASE	A210157 003	Jan 21, 2021
FOR SUSPENSION; ORAL			
OSELTAMIVIR PHOSPHATE			
CHEMISTRY HLTH	EQ 6MG BASE/ML	A214949 001	Feb 28, 2022
TAMIFLU			
ROCHE	EQ 12MG BASE/ML	N021246 001	Dec 14, 2000

OSILODROSTAT PHOSPHATE

TABLET; ORAL

ISTURISA

+ RECORDATI RARE EQ 10MG BASE N212801 003 Mar 06, 2020

OXACILLIN SODIUM

CAPSULE; ORAL

BACTOCILL

GLAXOSMITHKLINE	EQ 250MG BASE	A061336 001	
	EQ 250MG BASE	A062241 001	
	EQ 500MG BASE	A061336 002	
	EQ 500MG BASE	A062241 002	

OXACILLIN SODIUM

ANI PHARMS	EQ 250MG BASE	A062222 001	
	EQ 500MG BASE	A062222 002	
APOTHECON	EQ 250MG BASE	A061450 002	
	EQ 500MG BASE	A061450 001	

PROSTAPHLIN

APOTHECON EQ 500MG BASE N050118 002

FOR SOLUTION; ORAL

BACTOCILL

GLAXOSMITHKLINE EQ 250MG BASE/5ML A062321 001

OXACILLIN SODIUM

APOTHECON	EQ 250MG BASE/5ML	A061457 001	
TEVA	EQ 250MG BASE/5ML	A062252 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXACILLIN SODIUM

FOR SOLUTION; ORAL

PROSTAPHLIN

APOTHECON

EQ 250MG BASE/5ML

N050194 001

INJECTABLE; INJECTION

BACTOCILL

GLAXOSMITHKLINE

EQ 500MG BASE/VIAL **

A061334 009 Mar 26, 1982

EQ 1GM BASE/VIAL **

A061334 006 Mar 26, 1982

EQ 1GM BASE/VIAL **

A062736 001 Dec 19, 1986

EQ 2GM BASE/VIAL **

A061334 007 Mar 26, 1982

EQ 2GM BASE/VIAL **

A062736 002 Dec 19, 1986

EQ 4GM BASE/VIAL **

A061334 008 Mar 26, 1982

EQ 10GM BASE/VIAL **

A061334 010

OXACILLIN SODIUM

+ APOTHECON

EQ 250MG BASE/VIAL **

N050195 001

+ APOTHECON

EQ 500MG BASE/VIAL **

N050195 002

+ APOTHECON

EQ 1GM BASE/VIAL **

N050195 003

+ APOTHECON

EQ 2GM BASE/VIAL **

N050195 004

+ APOTHECON

EQ 4GM BASE/VIAL **

N050195 005

ELKINS SINN

EQ 250MG BASE/VIAL

A062711 001 Feb 03, 1989

EQ 500MG BASE/VIAL

A062711 002 Feb 03, 1989

EQ 1GM BASE/VIAL

A062711 003 Feb 03, 1989

EQ 2GM BASE/VIAL

A062711 004 Feb 03, 1989

EQ 4GM BASE/VIAL

A062711 005 Feb 03, 1989

EQ 10GM BASE/VIAL

A062711 006 Feb 03, 1989

HOSPIRA

EQ 1GM BASE/VIAL

A203950 001 Dec 11, 2015

EQ 2GM BASE/VIAL

A203950 002 Dec 11, 2015

ISTITUTO BIO ITA SPA

EQ 125MG BASE/VIAL

A062798 003 Dec 11, 1995

EQ 250MG BASE/VIAL

A062798 004 Dec 11, 1995

EQ 500MG BASE/VIAL

A062798 005 Dec 11, 1995

EQ 1GM BASE/VIAL

A062798 001 Dec 11, 1995

EQ 2GM BASE/VIAL

A062798 002 Dec 11, 1995

PIRAMAL CRITICAL

EQ 1GM BASE/VIAL

A206681 001 Sep 11, 2017

EQ 2GM BASE/VIAL

A206681 002 Sep 11, 2017

EQ 10GM BASE/VIAL

A206760 001 Oct 26, 2017

SANDOZ

EQ 250MG BASE/VIAL

A061490 001

EQ 500MG BASE/VIAL

A061490 002

EQ 1GM BASE/VIAL

A061490 003

EQ 2GM BASE/VIAL

A061490 004

EQ 10GM BASE/VIAL

A061490 006 May 09, 1991

STERISCIENCE

EQ 1GM BASE/VIAL

A091486 001 Aug 25, 2014

WATSON LABS INC

EQ 250MG BASE/VIAL

A062856 001 Oct 26, 1988

EQ 500MG BASE/VIAL

A062856 002 Oct 26, 1988

EQ 1GM BASE/VIAL

A062856 003 Oct 26, 1988

EQ 2GM BASE/VIAL

A062856 004 Oct 26, 1988

EQ 4GM BASE/VIAL

A062856 005 Oct 26, 1988

EQ 10GM BASE/VIAL

A062984 001 Sep 29, 1988

POWDER; INTRAVENOUS

OXACILLIN SODIUM

SANDOZ

EQ 1GM BASE/VIAL

A062737 001 Dec 23, 1986

EQ 2GM BASE/VIAL

A062737 002 Dec 23, 1986

OXALIPLATIN

INJECTABLE; INTRAVENOUS

ELOXATIN

+ SANOFI AVENTIS US

50MG/VIAL **

N021492 001 Aug 09, 2002

+ SANOFI AVENTIS US

100MG/VIAL **

N021492 002 Aug 09, 2002

+ SANOFI AVENTIS US

200MG/40ML (5MG/ML) **

N021759 003 Nov 17, 2006

OXALIPLATIN

ACCORD HLTHCARE

200MG/40ML (5MG/ML)

A207474 003 Mar 21, 2017

AM REGENT

50MG/10ML (5MG/ML)

A204378 001 May 12, 2017

100MG/20ML (5MG/ML)

A204378 002 May 12, 2017

CHARTWELL MOLECULAR

50MG/10ML (5MG/ML)

A208523 001 Feb 10, 2017

100MG/20ML (5MG/ML)

A208523 002 Feb 10, 2017

FRESENIUS KABI ONCOL

50MG/VIAL

A078810 001 Aug 07, 2009

100MG/VIAL

A078810 002 Aug 07, 2009

FRESENIUS KABI USA

200MG/40ML (5MG/ML)

A090030 003 Jan 31, 2017

GLAND

200MG/40ML (5MG/ML)

A207325 003 Oct 18, 2017

HOSPIRA INC

50MG/VIAL

A078815 001 Sep 30, 2009

100MG/VIAL

A078815 002 Sep 30, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXALIPLATIN

INJECTABLE; INTRAVENOUS

OXALIPLATIN

MYLAN LABS LTD	50MG/VIAL	A200979 001	Aug 08, 2012
	100MG/VIAL	A200979 002	Aug 08, 2012
	200MG/40ML (5MG/ML)	A091358 003	Nov 14, 2017
SANDOZ	50MG/10ML (5MG/ML)	A078812 001	Aug 07, 2009
	50MG/VIAL	A090849 001	Apr 28, 2011
	100MG/20ML (5MG/ML)	A078812 002	Aug 07, 2009
	100MG/VIAL	A090849 002	Apr 28, 2011
SUN PHARM	50MG/VIAL	A078818 001	Aug 07, 2009
	50MG/10ML (5MG/ML)	A202922 001	Apr 08, 2014
	100MG/VIAL	A078818 002	Aug 07, 2009
	100MG/20ML (5MG/ML)	A202922 002	Apr 08, 2014
	200MG/40ML (5MG/ML)	A202922 003	Feb 15, 2019

OXAMNIOUINE

CAPSULE; ORAL

VANSIL

PFIZER	250MG	N018069 001	
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OXANDROLONE

TABLET; ORAL

OXANDROLONE

PAR PHARM	2.5MG	A077827 001	Jun 22, 2007
	10MG	A077827 002	Jun 22, 2007
ROXANE	2.5MG	A077249 001	Jul 10, 2007
	10MG	A077249 002	Jul 10, 2007
SANDOZ	2.5MG	A076897 001	Dec 01, 2006
	10MG	A076897 002	Dec 01, 2006
UPSHER SMITH LABS	2.5MG	A076761 001	Dec 01, 2006
	10MG	A078033 001	Mar 22, 2007

OXAPROZIN

TABLET; ORAL

OXAPROZIN

ACTAVIS ELIZABETH	600MG	A075843 001	Oct 03, 2001
BEXIMCO PHARMS USA	600MG	A075842 001	Apr 12, 2001
IVAX SUB TEVA PHARMS	600MG	A075846 001	May 13, 2002
MYLAN	600MG	A075851 001	Aug 17, 2001
MYLAN PHARMS INC	600MG	A075847 001	Feb 28, 2001
SANDOZ	600MG	A075850 001	Apr 27, 2001
SUN PHARM INDS INC	600MG	A075844 001	Jan 03, 2002
WATSON LABS	600MG	A075848 001	Feb 09, 2001

OXAPROZIN POTASSIUM

TABLET; ORAL

DAYPRO ALTA

PFIZER	600MG	N020776 001	Oct 17, 2002
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OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AM THERAP	10MG	A071955 001	Mar 03, 1988
	15MG	A071956 001	Mar 03, 1988
	30MG	A071957 001	Mar 03, 1988
IVAX SUB TEVA PHARMS	10MG	A070943 001	Aug 03, 1987
	15MG	A070944 001	Aug 03, 1987
	30MG	A070945 001	Aug 03, 1987
MYLAN	10MG	A071713 001	Oct 20, 1987
	15MG	A071714 001	Oct 20, 1987
	30MG	A071715 001	Oct 20, 1987
WATSON LABS	15MG	A072953 001	Sep 28, 1990
	30MG	A072954 001	Sep 28, 1990
WATSON LABS TEVA	10MG	A072952 001	Sep 28, 1990

SERAX

+ ALPHARMA US PHARMS	10MG **	N015539 002	
+	15MG **	N015539 004	
+	30MG **	N015539 006	

ZAXOPAM

QUANTUM PHARMICS	10MG	A070650 001	Mar 01, 1988
	15MG	A070640 001	Mar 01, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXAZEPAMCAPSULE; ORAL
ZAXOPAM

30MG

A070641 001 Mar 01, 1988

TABLET; ORAL

OXAZEPAM

PARKE DAVIS

15MG

A071508 001 Feb 02, 1987

SUN PHARM INDUSTRIES

15MG

A070683 001 Jan 16, 1987

WATSON LABS

15MG

A071494 001 Apr 21, 1987

SERAX

ALPHARMA US PHARMS

15MG **

N015539 008

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

HIKMA

300MG/5ML

A201193 001 Oct 03, 2012

RENEW PHARMS

300MG/5ML

A211420 001 Jul 09, 2021

TABLET; ORAL

OXCARBAZEPINE

ANI PHARMS

150MG

A078005 001 Dec 11, 2007

300MG

A078005 002 Dec 11, 2007

600MG

A078005 003 Dec 11, 2007

HIKMA

150MG

A077795 001 Oct 09, 2007

300MG

A077795 002 Oct 09, 2007

600MG

A077795 003 Oct 09, 2007

JUBILANT CADISTA

150MG

A090239 001 Jan 25, 2010

300MG

A090239 002 Jan 25, 2010

600MG

A090239 003 Jan 25, 2010

TABLET, EXTENDED RELEASE; ORAL

OXCARBAZEPINE

APOTEX

150MG

A213369 001 Jul 13, 2023

300MG

A213369 002 Jul 13, 2023

600MG

A213369 003 Jul 13, 2023

OPRENOLOL HYDROCHLORIDE

CAPSULE; ORAL

TRASICOR

NOVARTIS

20MG

N018166 001 Dec 28, 1983

40MG

N018166 002 Dec 28, 1983

80MG

N018166 003 Dec 28, 1983

160MG

N018166 004 Dec 28, 1983

OXTRIPHYLLINE

SOLUTION; ORAL

CHOLEDYL

PARKE DAVIS

100MG/5ML

N009268 012 Nov 27, 1984

OXTRIPHYLLINE

MORTON GROVE

100MG/5ML

A088243 001 Dec 05, 1983

SYRUP; ORAL

CHOLEDYL

PARKE DAVIS

50MG/5ML

N009268 011

OXTRIPHYLLINE PEDIATRIC

MORTON GROVE

50MG/5ML

A088242 001 Dec 05, 1983

TABLET, DELAYED RELEASE; ORAL

CHOLEDYL

PARKE DAVIS

100MG

N009268 003

200MG

N009268 007

OXTRIPHYLLINE

WATSON LABS

100MG

A087866 001 Aug 25, 1983

200MG

A087835 001 Aug 25, 1983

TABLET, EXTENDED RELEASE; ORAL

CHOLEDYL SA

WARNER CHILCOTT LLC

400MG

A087863 001 May 24, 1983

600MG

A086742 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYBUTYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL

OXYBUTYNIN

BARR LABS DIV TEVA 3.9MG/24HR

A090526 001 Mar 04, 2014

GEL, METERED;TRANSDERMAL

GELNIQUE 3%

+ ALLERGAN 3%

N202513 001 Dec 07, 2011

OXYBUTYNIN CHLORIDE

GEL;TRANSDERMAL

GELNIQUE

+ ABBVIE 10% (100MG/PACKET)

N022204 001 Jan 27, 2009

OXYBUTYNIN CHLORIDE

CHARTWELL RX 10% (100MG/PACKET)

A207329 001 May 31, 2018

SYRUP;ORAL

DITROPAN

+ ORTHO MCNEIL JANSSEN 5MG/5ML **

N018211 001

OXYBUTYNIN CHLORIDE

LANNETT CO INC 5MG/5ML

A076682 001 Dec 28, 2004

PHARM ASSOC 5MG/5ML

A074997 001 Oct 15, 1997

PHARMOBEDIANT CNSLTG 5MG/5ML

A074868 001 Feb 12, 1997

TABLET;ORAL

DITROPAN

+ JANSSEN PHARMS 5MG **

N017577 001

OXYBUTYNIN CHLORIDE

AVET LIFESCIENCES 5MG

A211682 001 May 10, 2019

QUANTUM PHARMICS 5MG

A072296 001 Dec 08, 1988

USL PHARMA 5MG

A070746 001 Mar 10, 1988

WATSON LABS 5MG

A072485 001 Apr 19, 1989

TABLET, EXTENDED RELEASE;ORAL

DITROPAN XL

+ JANSSEN PHARMS 5MG

N020897 001 Dec 16, 1998

+ 10MG

N020897 002 Dec 16, 1998

+ 15MG **

N020897 003 Jun 22, 1999

OXYBUTYNIN CHLORIDE

IMPAX PHARMS 5MG

A076745 002 May 09, 2007

10MG

A076745 003 May 09, 2007

15MG

A076745 001 Nov 09, 2006

MYLAN 5MG

A076702 001 Nov 09, 2006

MYLAN PHARMS INC 10MG

A076644 001 Nov 09, 2006

15MG

A076644 002 May 10, 2007

OXYCODONE HYDROCHLORIDE

CAPSULE;ORAL

OXYCODONE HYDROCHLORIDE

DR REDDYS LABS SA 5MG

A203107 001 Jul 26, 2012

LANNETT CO INC 5MG

A203823 001 Aug 01, 2014

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

ANI PHARMS 100MG/5ML

A203447 001 Aug 30, 2017

AUROLIFE PHARMA LLC 5MG/5ML

A212429 001 Jan 27, 2020

100MG/5ML

A212429 002 Jan 27, 2020

CHARTWELL MOLECULAR 100MG/5ML

A204085 001 Sep 09, 2014

DR REDDYS LABS SA 100MG/5ML

A204092 001 Jun 05, 2014

HIKMA 100MG/5ML

A203208 001 Jul 12, 2013

NOVEL LABS INC 100MG/5ML

A204603 001 Apr 29, 2015

PHARMOBEDIANT CNSLTG 5MG/5ML

A206456 001 Jun 16, 2015

RHODES PHARMS 100MG/5ML

A205853 001 Apr 29, 2020

+ VISTAPHARM 5MG/5ML

N201194 001 Jan 12, 2012

100MG/5ML

A202537 001 Jul 30, 2012

TABLET;ORAL

OXAYDO

+ ZYLA 5MG

N202080 001 Jun 17, 2011

+ 7.5MG

N202080 002 Jun 17, 2011

OXYCODONE HYDROCHLORIDE

ACTAVIS ELIZABETH 5MG

A076636 003 Apr 07, 2015

15MG

A076636 001 Feb 06, 2004

30MG

A076636 002 Feb 06, 2004

DR REDDYS LABS SA 5MG

A091313 001 Feb 18, 2011

10MG

A091313 004 Apr 29, 2016

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

	15MG	A091313 002	Feb 18, 2011
	20MG	A091313 005	Apr 29, 2016
	30MG	A091313 003	Feb 18, 2011
NESHER PHARMS	5MG	A077290 001	Dec 08, 2005
	10MG	A077290 002	Dec 08, 2005
	15MG	A077290 003	Dec 08, 2005
	20MG	A077290 004	Dec 08, 2005
	30MG	A077290 005	Dec 08, 2005
STRIDES PHARMA	5MG	A077712 003	Mar 02, 2009
	10MG	A077712 004	Apr 13, 2015
	15MG	A077712 001	Jan 31, 2007
	20MG	A077712 005	Apr 13, 2015
	30MG	A077712 002	Jan 31, 2007

TABLET, EXTENDED RELEASE; ORAL

ROXICODONE

ROXANE	10MG	N020932 001	Oct 26, 1998
	30MG	N020932 002	Oct 26, 1998

OXYMETAZOLINE HYDROCHLORIDE

CREAM; TOPICAL

OXYMETAZOLINE HYDROCHLORIDE

TARO	1%	A213584 001	Oct 04, 2021
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SOLUTION/DROPS; OPHTHALMIC

OCUCLEAR

BAYER HEALTHCARE LLC	0.025%	N018471 001	May 30, 1986
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OXYMETHOLONE

TABLET; ORAL

ANADROL-50

+ MYLAN SPECIALITY LP	50MG	N016848 001	
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OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

OPANA

+ ENDO PHARMS	1MG/ML	N011707 002	
	1.5MG/ML	N011707 001	

SUPPOSITORY; RECTAL

NUMORPHAN

ENDO PHARMS	5MG	N011738 004	
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TABLET; ORAL

OPANA

+ ENDO PHARMS	5MG **	N021611 001	Jun 22, 2006
	10MG **	N021611 002	Jun 22, 2006

OXYMORPHONE HYDROCHLORIDE

SPECGX LLC	5MG	A202321 001	Apr 25, 2013
	10MG	A202321 002	Apr 25, 2013

TABLET, EXTENDED RELEASE; ORAL

OPANA ER

+ ENDO PHARMS	5MG **	N021610 001	Jun 22, 2006
	5MG	N201655 001	Dec 09, 2011
	7.5MG **	N021610 005	Feb 29, 2008
	7.5MG	N201655 002	Dec 09, 2011
	10MG **	N021610 002	Jun 22, 2006
	10MG	N201655 003	Dec 09, 2011
	15MG **	N021610 006	Feb 29, 2008
	15MG	N201655 004	Dec 09, 2011
	20MG **	N021610 003	Jun 22, 2006
	20MG	N201655 005	Dec 09, 2011
	30MG **	N021610 007	Feb 29, 2008
	30MG	N201655 006	Dec 09, 2011
	40MG **	N021610 004	Jun 22, 2006
	40MG	N201655 007	Dec 09, 2011

OXYMORPHONE HYDROCHLORIDE

ACTAVIS ELIZABETH	5MG	A079046 003	Jul 11, 2013
	7.5MG	A079046 001	Dec 13, 2010
	10MG	A079046 004	Jul 11, 2013
	15MG	A079046 002	Dec 13, 2010
	20MG	A079046 005	Jul 11, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYMORPHONE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
OXYMORPHONE HYDROCHLORIDE

	30MG	A079046 006	Jul 11, 2013
	40MG	A079046 007	Jul 11, 2013
HIKMA	5MG	A200822 002	Jul 15, 2013
	7.5MG	A200822 003	Jul 15, 2013
	10MG	A200822 004	Jul 15, 2013
	15MG	A200822 005	Jul 15, 2013
	20MG	A200822 006	Jul 15, 2013
	30MG	A200822 007	Jul 15, 2013
	40MG	A200822 001	Jul 15, 2013
PAR PHARM	5MG	A200792 001	Oct 24, 2014
	7.5MG	A200792 002	Oct 24, 2014
	10MG	A200792 003	Oct 24, 2014
	15MG	A200792 004	Oct 24, 2014
	20MG	A200792 005	Oct 24, 2014
	30MG	A200792 006	Oct 24, 2014
	40MG	A200792 007	Oct 24, 2014
SPECGX LLC	5MG	A202946 001	Jun 27, 2014
	7.5MG	A202946 002	Jun 27, 2014
	10MG	A202946 003	Jun 27, 2014
	15MG	A202946 004	Jun 27, 2014
	20MG	A202946 005	Jun 27, 2014
	30MG	A202946 006	Jun 27, 2014
	40MG	A202946 007	Jun 27, 2014
SUN PHARM INDS LTD	5MG	A203506 001	Apr 24, 2015
	7.5MG	A203506 002	Apr 24, 2015
	10MG	A203506 003	Apr 24, 2015
	15MG	A203506 004	Apr 24, 2015
	20MG	A203506 005	Apr 24, 2015
	30MG	A203506 006	Apr 24, 2015
	40MG	A203506 007	Apr 24, 2015

OXYPHENBUTAZONE

TABLET;ORAL

OXYPHENBUTAZONE

WATSON LABS	100MG	A088399 001	Sep 17, 1984
TANDEARIL			
NOVARTIS	100MG	N012542 004	Sep 03, 1982

OXYPHENCYCLIMINE HYDROCHLORIDE

TABLET;ORAL

DARICON

PFIZER	10MG	N011612 001	
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OXYPHENONIUM BROMIDE

TABLET;ORAL

ANTRENYL

NOVARTIS	5MG	N008492 002	
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OXYTETRACYCLINE

TABLET;ORAL

TERRAMYCIN

PFIZER	250MG	N050287 001	
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OXYTETRACYCLINE CALCIUM

SYRUP;ORAL

TERRAMYCIN

PFIZER	EQ 125MG BASE/5ML	A060595 001	
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OXYTETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

OXY-KESSO-TETRA

FERRANTE	EQ 250MG BASE	A060179 001	
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OXYTETRACYCLINE HYDROCHLORIDE

HIKMA	EQ 250MG BASE	A060770 001	
IMPAX LABS	EQ 250MG BASE	A060760 001	
PROTER	EQ 250MG BASE	A060869 001	
PUREPAC PHARM	EQ 250MG BASE	A060634 001	
TERRAMYCIN			
PFIZER	EQ 125MG BASE	N050286 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYTETRACYCLINE HYDROCHLORIDECAPSULE; ORAL
TERRAMYCIN

EQ 250MG BASE

N050286 002

INJECTABLE; INJECTION
TERRAMYCIN

PFIZER

EQ 250MG BASE/VIAL

A060586 001

EQ 500MG BASE/VIAL

A060586 002

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

TERRAMYCIN W/ POLYMYXIN B SULFATE

CASPER PHARMA LLC

EQ 5MG BASE/GM;10,000 UNITS/GM

N061015 001

OINTMENT; OTIC

TERRAMYCIN W/ POLYMYXIN

PFIZER

EQ 5MG BASE/GM;10,000 UNITS/GM

A061841 001

TABLET; VAGINAL

TERRAMYCIN-POLYMYXIN

PFIZER

EQ 100MG BASE;100,000 UNITS

A061009 001

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

DR REDDYS

10USP UNITS/ML (10USP UNITS/ML)

A077453 001 Jan 24, 2008

100USP UNITS/10ML (10USP UNITS/ML)

A077453 002 Jan 24, 2008

OXYTOCIN 10 USP UNITS IN DEXTROSE 5%

+ ABBOTT

1USP UNITS/100ML **

N019185 004 Mar 29, 1985

+

2USP UNITS/100ML **

N019185 003 Mar 29, 1985

OXYTOCIN 20 USP UNITS IN DEXTROSE 5%

+ ABBOTT

2USP UNITS/100ML **

N019185 002 Mar 29, 1985

OXYTOCIN 5 USP UNITS IN DEXTROSE 5%

+ ABBOTT

1USP UNITS/100ML **

N019185 001 Mar 29, 1985

SYNTOCINON

NOVARTIS

10USP UNITS/ML

N018245 001

SOLUTION; NASAL

SYNTOCINON

RTRX

40USP UNITS/ML

N012285 001

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL

ACCORD HLTHCARE

6MG/ML

A075436 001 Nov 12, 2004

DASH PHARMS

6MG/ML

A091540 001 Sep 29, 2011

HOSPIRA

6MG/ML

A076233 001 Aug 01, 2002

MYLAN

6MG/ML

A075278 001 Jan 25, 2002

PLIVA LACHEMA

6MG/ML

A077413 001 Mar 12, 2008

SANDOZ

6MG/ML

A078167 001 Dec 26, 2007

TEVA PHARMS USA

6MG/ML

A075297 001 Jan 25, 2002

TAXOL

+ HQ SPCLT PHARMA

6MG/ML **

N020262 001 Dec 29, 1992

POWDER; INTRAVENOUS

PACLITAXEL

+ TEVA PHARMS INC

100MG/VIAL

N216338 001 May 11, 2023

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA

+ JANSSEN PHARMS

12MG **

N021999 004 Dec 19, 2006

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

PALIPERIDONE PALMITATE

TEVA PHARMS USA

39MG/0.25ML (39MG/0.25ML)

A211149 001 Jul 06, 2021

78MG/0.5ML (78MG/0.5ML)

A211149 002 Jul 06, 2021

117MG/0.75ML (117MG/0.75ML)

A211149 003 Jul 06, 2021

156MG/ML (156MG/ML)

A211149 004 Jul 06, 2021

234MG/1.5ML (156MG/ML)

A211149 005 Jul 06, 2021

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

ALOXI

+ HELSINN HLTHCARE EQ 0.5MG BASE ** N022233 001 Aug 22, 2008

INJECTABLE; INTRAVENOUS

ALOXI

+ HELSINN HLTHCARE EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML) ** N021372 002 Feb 29, 2008+ EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)
** N021372 001 Jul 25, 2003

PALONOSETRON HYDROCHLORIDE

ACCORD HLTHCARE EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A204615 001 Mar 15, 2021

CIPLA EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A206396 001 Sep 19, 2018

DR REDDYS EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML) A201533 001 Apr 21, 2016HOSPIRA EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML) A207005 002 Sep 19, 2018

NOVAST LABS EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) A208789 001 May 22, 2020

QILU PHARM HAINAN EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML) A205648 002 Sep 19, 2018TEVA PHARMS USA EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML) A090713 002 Mar 23, 2018

SOLUTION; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

AVYXA PHARMA EQ 0.075MG BASE/1.5ML (EQ 0.05MG
BASE/ML) N203050 001 Mar 01, 2016

EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) N203050 002 Mar 01, 2016

+ FRESENIUS KABI USA EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) N208109 001 Nov 21, 2017

HIKMA EQ 0.25MG BASE/2ML (EQ 0.125MG BASE/ML) N207963 001 Aug 22, 2016

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

AREZIA

+ NOVARTIS 30MG/VIAL ** N020036 001 Oct 31, 1991

60MG/VIAL N020036 003 May 06, 1993

90MG/VIAL N020036 004 May 06, 1993

PAMIDRONATE DISODIUM

AESGEN 30MG/VIAL A075594 001 May 06, 2002

90MG/VIAL A075594 002 May 06, 2002

AM REGENT 30MG/10ML (3MG/ML) A078942 001 Jul 25, 2008

90MG/10ML (9MG/ML) A078942 002 Jul 25, 2008

FRESENIUS KABI USA 30MG/VIAL A075773 001 May 06, 2002

30MG/10ML (3MG/ML) A076207 001 May 17, 2002

90MG/VIAL A075773 002 May 06, 2002

90MG/10ML (9MG/ML) A076207 002 May 17, 2002

MN PHARMS 30MG/VIAL A078300 001 Mar 10, 2009

90MG/VIAL A078300 002 Mar 10, 2009

SUN PHARMA GLOBAL 30MG/VIAL A077703 001 Dec 24, 2008

90MG/VIAL A077703 002 Dec 24, 2008

TEVA PHARMS USA 30MG/10ML (3MG/ML) A076153 001 Mar 27, 2002

90MG/10ML (9MG/ML) A076153 002 Mar 27, 2002

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

ELKINS SINN 1MG/ML A072058 001 Mar 23, 1988

2MG/ML A072059 001 Mar 23, 1988

2MG/ML A072060 001 Mar 23, 1988

HOSPIRA 2MG/ML A072321 001 Jan 19, 1989

IGI LABS INC 1MG/ML A072210 001 Mar 31, 1988

2MG/ML A072211 001 Mar 31, 1988

2MG/ML A072212 001 Mar 31, 1988

2MG/ML A072213 001 Mar 31, 1988

PAVULON

+ ORGANON USA INC 1MG/ML ** N017015 002

+ 2MG/ML ** N017015 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PANOBINOSTAT LACTATE

CAPSULE; ORAL

FARYDAK

+	SECURA	EQ 10MG BASE	N205353 001	Feb 23, 2015
+		EQ 15MG BASE	N205353 002	Feb 23, 2015
+		EQ 20MG BASE	N205353 003	Feb 23, 2015

PANTOPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

PANTOPRAZOLE SODIUM

MYLAN LABS LTD

EQ 40MG BASE/VIAL A208580 001 May 04, 2018

TABLET, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

JUBILANT GENERICS

EQ 20MG BASE A090901 001 Aug 30, 2011

EQ 40MG BASE A090901 002 Aug 30, 2011

L PERRIGO CO

EQ 20MG BASE A203024 001 May 07, 2014

MACLEODS PHARMS LTD

EQ 20MG BASE A200821 001 Feb 16, 2012

EQ 40MG BASE A200821 002 Feb 16, 2012

SUN PHARM

EQ 20MG BASE A077058 001 Sep 10, 2007

EQ 40MG BASE A077058 002 Sep 10, 2007

SUN PHARM INDS LTD

EQ 20MG BASE A200794 001 May 02, 2012

EQ 40MG BASE A200794 002 May 02, 2012

TEVA

EQ 20MG BASE A077056 001 Aug 02, 2007

EQ 40MG BASE A077056 002 Aug 02, 2007

PARAMETHADIONE

CAPSULE; ORAL

PARADIONE

ABBVIE

150MG N006800 003

300MG N006800 001

SOLUTION; ORAL

PARADIONE

ABBVIE

300MG/ML N006800 002

PARAMETHASONE ACETATE

TABLET; ORAL

HALDRONE

LILLY

1MG N012772 005

2MG N012772 006

PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTONYL

ABBOTT

10MG N013448 002

25MG N013448 003

50MG N013448 004

PARICALCITOL

CAPSULE; ORAL

PARICALCITOL

LOTUS PHARM CO LTD

1MCG A206710 001 Feb 24, 2016

2MCG A206710 002 Feb 24, 2016

4MCG A206710 003 Feb 24, 2016

ZEMPLAR

+ ABBVIE

4MCG ** N021606 003 May 26, 2005

SOLUTION; INTRAVENOUS

PARICALCITOL

AKORN

0.01MG/2ML (0.005MG/ML) A207692 001 Oct 16, 2017

RISING

0.002MG/ML (0.002MG/ML) A203897 001 Nov 02, 2017

PAROMOMYCIN SULFATE

CAPSULE; ORAL

HUMATIN

KING PFIZER

EQ 250MG BASE A062310 001

PARKEDALE

EQ 250MG BASE A060521 001

PAROMOMYCIN SULFATE

SUN PHARM INDS INC

EQ 250MG BASE A064171 001 Jun 30, 1997

SYRUP; ORAL

HUMATIN

PARKE DAVIS

EQ 125MG BASE/5ML A060522 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PAROXETINE HYDROCHLORIDE

CAPSULE; ORAL

PAXIL

+	APOTEX	EQ 10MG BASE **	N020885 001	Oct 09, 1998
+		EQ 20MG BASE **	N020885 002	Oct 09, 1998
+		EQ 30MG BASE **	N020885 003	Oct 09, 1998
+		EQ 40MG BASE **	N020885 004	Oct 09, 1998

SUSPENSION; ORAL

PAROXETINE HYDROCHLORIDE

APOTEX INC

EQ 10MG BASE/5ML A077395 001 Dec 05, 2006

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

JUBILANT GENERICS

		EQ 10MG BASE	A205528 001	Nov 27, 2015
		EQ 20MG BASE	A205528 002	Nov 27, 2015
		EQ 30MG BASE	A205528 003	Nov 27, 2015
		EQ 40MG BASE	A205528 004	Nov 27, 2015

MYLAN PHARMS INC

		EQ 10MG BASE	A075716 001	Mar 08, 2004
		EQ 20MG BASE	A075716 002	Mar 08, 2004
		EQ 30MG BASE	A075716 003	Mar 08, 2004
		EQ 40MG BASE	A075716 004	Mar 08, 2004

ROXANE

		EQ 10MG BASE	A078026 001	Jun 29, 2007
		EQ 20MG BASE	A078026 002	Jun 29, 2007
		EQ 30MG BASE	A078026 003	Jun 29, 2007
		EQ 40MG BASE	A078026 004	Jun 29, 2007

SUN PHARM INDS INC

		EQ 10MG BASE	A078194 001	Jun 29, 2007
		EQ 20MG BASE	A078194 002	Jun 29, 2007
		EQ 30MG BASE	A078194 003	Jun 29, 2007
		EQ 40MG BASE	A078194 004	Jun 29, 2007

TEVA PHARMS

		EQ 10MG BASE	A077082 001	Jun 29, 2007
		EQ 20MG BASE	A077082 002	Jun 29, 2007
		EQ 30MG BASE	A077082 003	Jun 29, 2007
		EQ 40MG BASE	A077082 004	Jun 29, 2007

UPSHER SMITH LABS

		EQ 10MG BASE	A075566 001	Mar 08, 2004
		EQ 20MG BASE	A075566 002	Mar 08, 2004
		EQ 30MG BASE	A075566 003	Mar 08, 2004
		EQ 40MG BASE	A075566 004	Mar 08, 2004

PAXIL

APOTEX

EQ 50MG BASE N020031 004 Dec 29, 1992

TABLET, EXTENDED RELEASE; ORAL

PAROXETINE HYDROCHLORIDE

EPIC PHARMA LLC

		EQ 12.5MG BASE	A213612 001	Aug 11, 2021
		EQ 25MG BASE	A213612 002	Aug 11, 2021
		EQ 37.5MG BASE	A213612 003	May 26, 2022

PAROXETINE MESYLATE

TABLET; ORAL

PEXEVA

+	SEBELA IRELAND LTD	EQ 10MG BASE	N021299 001	Jul 03, 2003
+		EQ 20MG BASE	N021299 002	Jul 03, 2003
+		EQ 30MG BASE	N021299 003	Jul 03, 2003
+		EQ 40MG BASE	N021299 004	Jul 03, 2003

PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT

+ NOVARTIS

EQ 400MG BASE ** N022465 002 Oct 19, 2009

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

+ BAUSCH AND LOMB INC

EQ 0.3MG ACID/0.09ML N021756 001 Dec 17, 2004

PEGINESATIDE ACETATE

SOLUTION; INTRAVENOUS, SUBCUTANEOUS

OMONTYS

TAKEDA PHARMS USA

		EQ 10MG BASE/ML (EQ 10MG BASE/ML)	N202799 007	Mar 27, 2012
		EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N202799 008	Mar 27, 2012

OMONTYS PRESERVATIVE FREE

TAKEDA PHARMS USA

		EQ 1MG BASE/0.5ML (EQ 1MG BASE/0.5ML)	N202799 001	Mar 27, 2012
		EQ 2MG BASE/0.5ML (EQ 2MG BASE/0.5ML)	N202799 002	Mar 27, 2012
		EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)	N202799 003	Mar 27, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PEGINESATIDE ACETATESOLUTION; INTRAVENOUS, SUBCUTANEOUS
OMONTYS PRESERVATIVE FREE

EQ 4MG BASE/0.5ML (EQ 4MG BASE/0.5ML)	N202799 004	Mar 27, 2012
EQ 5MG BASE/0.5ML (EQ 5MG BASE/0.5ML)	N202799 005	Mar 27, 2012
EQ 6MG BASE/0.5ML (EQ 6MG BASE/0.5ML)	N202799 006	Mar 27, 2012

PEMETREXED DISODIUMPOWDER; INTRAVENOUS
PEMETREXED DISODIUM
AMNEAL

EQ 100MG BASE/VIAL	A210047 002	May 08, 2023
EQ 500MG BASE/VIAL	A210047 001	Aug 04, 2022
BIOCON PHARMA	EQ 1GM BASE/VIAL	A211090 001
HOSPIRA	EQ 100MG BASE/VIAL	A202111 001
	EQ 500MG BASE/VIAL	A202111 002
	EQ 1GM BASE/VIAL	A202111 003
MYLAN LABS LTD	EQ 500MG BASE/VIAL	A203628 001

PEMETREXED DITROMETHAMINEPOWDER; INTRAVENOUS
PEMETREXED DITROMETHAMINE
+ HOSPIRA

EQ 1GM BASE/VIAL	N208746 003	Jun 10, 2022
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PEMIROLAST POTASSIUMSOLUTION/DROPS; OPHTHALMIC
ALAMAST
SANTEN

0.1%	N021079 001	Sep 24, 1999
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PEMOLINETABLET; ORAL
CYLERT

ABBOTT	18.75MG	N016832 001
	37.5MG	N016832 002
	75MG	N016832 003

PEMOLINE

ACTAVIS ELIZABETH	18.75MG	A075595 001	Feb 28, 2000
	37.5MG	A075595 002	Feb 28, 2000
	75MG	A075595 003	Feb 28, 2000
FOSUN PHARMA	18.75MG	A075286 001	Dec 27, 1999
	37.5MG	A075286 002	Jun 30, 1999
	75MG	A075286 003	Jun 30, 1999
MALLINCKRODT	18.75MG	A075726 003	Mar 30, 2001
	37.5MG	A075726 002	Mar 30, 2001
	75MG	A075726 001	Mar 30, 2001
TEVA PHARMS	18.75MG	A075030 003	Feb 22, 2000
	37.5MG	A075030 001	Jan 29, 1999
	75MG	A075030 002	Jan 29, 1999
VINTAGE PHARMS	18.75MG	A075328 001	Apr 19, 2000
	37.5MG	A075328 002	Apr 19, 2000
	75MG	A075328 003	Apr 19, 2000
WATSON LABS	18.75MG	A075287 001	Jun 13, 2001
	37.5MG	A075287 002	Sep 18, 2000
	75MG	A075287 003	Sep 18, 2000

TABLET, CHEWABLE; ORAL

CYLERT

ABBOTT	37.5MG	N017703 001
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PEMOLINE

ACTAVIS ELIZABETH	37.5MG	A075678 001	Jul 26, 2000
TEVA PHARMS	37.5MG	A075555 001	Feb 18, 2000

PENBUTOLOL SULFATE

TABLET; ORAL

LEVATOL

+ AUXILIUM PHARMS LLC	10MG **	N018976 001	Dec 30, 1987
+	20MG **	N018976 004	Jan 05, 1989

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLAMINECAPSULE; ORAL
CUPRIMINE

VALEANT PHARMS INTL 125MG N019853 002

TABLET; ORAL

PENICILLAMINE

TEVA PHARMS USA 250MG A211497 001 Feb 13, 2020

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

+ KING PHARMS LLC 300,000 UNITS/ML N050141 003

WYETH AYERST 300,000 UNITS/ML N050131 001

PERMAPEN

CASPER PHARMA LLC 600,000 UNITS/ML N060014 001

SUSPENSION; ORAL

BICILLIN

WYETH AYERST 300,000 UNITS/5ML N050126 002

TABLET; ORAL

BICILLIN

WYETH AYERST 200,000 UNITS N050128 001

PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

+ KING PHARMS LLC 150,000 UNITS/ML; 150,000 UNITS/ML N050138 002

PENICILLIN G POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN

TEVA 200,000 UNITS/5ML A060307 002

400,000 UNITS/5ML A060307 004

PENICILLIN G POTASSIUM

MYLAN 200,000 UNITS/5ML A060752 003

250,000 UNITS/5ML A060752 002

400,000 UNITS/5ML A060752 001

PUREPAC PHARM 250,000 UNITS/5ML A061740 001

400,000 UNITS/5ML A061740 002

PENICILLIN-2

TEVA 250,000 UNITS/5ML A060307 003

PENTIDS '200'

APOTHECON 200,000 UNITS/5ML A062149 001

PENTIDS '400'

APOTHECON 400,000 UNITS/5ML A062149 002

PFIZERPEN G

PFIZER 400,000 UNITS/5ML A060587 001

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

APOTHECON 1,000,000 UNITS/VIAL A060362 001

5,000,000 UNITS/VIAL A060362 003

10,000,000 UNITS/VIAL A060362 004

20,000,000 UNITS/VIAL A060362 002

CONSOLIDATED PHARM 500,000 UNITS/VIAL A060806 001

1,000,000 UNITS/VIAL A060806 002

5,000,000 UNITS/VIAL A060806 003

10,000,000 UNITS/VIAL A060806 004

LILLY 200,000 UNITS/VIAL A060384 004

500,000 UNITS/VIAL A060384 003

1,000,000 UNITS/VIAL A060384 002

5,000,000 UNITS/VIAL A060384 001

20,000,000 UNITS/VIAL A060384 005

20,000,000 UNITS/VIAL A060601 001

PARKE DAVIS 1,000,000 UNITS/VIAL A062003 001

5,000,000 UNITS/VIAL A062003 002

PFIZER 20,000,000 UNITS/VIAL A060074 003

SANDOZ 1,000,000 UNITS/VIAL ** A065079 001 Aug 30, 2002

WATSON LABS INC 1,000,000 UNITS/VIAL A062991 001 Sep 13, 1988

5,000,000 UNITS/VIAL A062991 002 Sep 13, 1988

10,000,000 UNITS/VIAL A062991 003 Sep 13, 1988

20,000,000 UNITS/VIAL A062991 004 Sep 13, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PFIZERPEN

PFIZER 1,000,000 UNITS/VIAL ** A060657 001

TABLET; ORAL

PENICILLIN G POTASSIUM

APOTHECON 250,000 UNITS A060392 003

IVAX SUB TEVA PHARMS 400,000 UNITS A060073 004

LILLY 250,000 UNITS A060403 001

MYLAN 200,000 UNITS A060781 001

250,000 UNITS A060781 002

400,000 UNITS A060781 003

500,000 UNITS A060781 005

800,000 UNITS A060781 004

PUREPAC PHARM 200,000 UNITS A061588 001

250,000 UNITS A061588 002

400,000 UNITS A061588 003

TEVA 200,000 UNITS A060306 001

250,000 UNITS A060306 002

400,000 UNITS A060306 003

500,000 UNITS A060306 004

WYETH AYERST 200,000 UNITS A060413 001

250,000 UNITS A060413 002

400,000 UNITS A060413 003

PENTIDS '200'

APOTHECON 200,000 UNITS A062155 001

PENTIDS '250'

APOTHECON 250,000 UNITS A062155 002

PENTIDS '400'

APOTHECON 400,000 UNITS A060392 004

400,000 UNITS A062155 003

PENTIDS '800'

APOTHECON 800,000 UNITS A060392 005

800,000 UNITS A062155 004

PFIZERPEN G

PFIZER 50,000 UNITS A060075 001

100,000 UNITS A060075 002

200,000 UNITS A060075 003

250,000 UNITS A060075 004

400,000 UNITS A060075 005

800,000 UNITS A060075 006

PENICILLIN G PROCAINE

INJECTABLE; INJECTION

DURACILLIN A.S.

LILLY 300,000 UNITS/ML A060093 001

PENICILLIN G PROCAINE

CONSOLIDATED PHARM 300,000 UNITS/ML A060800 001

600,000 UNITS/1.2ML A060800 002

PARKE DAVIS 300,000 UNITS/ML A062029 001

PFIZER 300,000 UNITS/VIAL A060099 001

1,500,000 UNITS/VIAL A060099 002

PFIZERPEN-AS

PFIZER 300,000 UNITS/ML A060286 001

600,000 UNITS/ML A060286 002

PENICILLIN G SODIUM

INJECTABLE; INJECTION

PENICILLIN G SODIUM

BRISTOL MYERS SQUIBB 5,000,000 UNITS/VIAL A061935 001

COPANOS 5,000,000 UNITS/VIAL A061051 001

PHARMACIA AND UPJOHN 1,000,000 UNITS/VIAL A061046 001

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

PENICILLIN G SODIUM

WATSON LABS INC 5,000,000 UNITS/VIAL A063014 001 Sep 13, 1988

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN VFOR SUSPENSION;ORAL
V-CILLIN

LILLY 125MG/0.6ML A060002 001

PENICILLIN V POTASSIUM

FOR SOLUTION;ORAL

BEEPEN-VK

GLAXOSMITHKLINE EQ 125MG BASE/5ML A062270 001
EQ 250MG BASE/5ML A062270 002

BETAPEN-VK

APOTHECON EQ 125MG BASE/5ML A061149 001
EQ 250MG BASE/5ML A061149 002

LEDERCILLIN VK

LEDERLE EQ 125MG BASE/5ML A060136 001
EQ 250MG BASE/5ML A060136 002

PEN-VEE K

WYETH AYERST EQ 125MG BASE/5ML A060007 001
EQ 250MG BASE/5ML A060007 002

PENAPAR-VK

PARKE DAVIS EQ 125MG BASE/5ML A062002 001
EQ 250MG BASE/5ML A062002 002

PENICILLIN V POTASSIUM

BELCHER PHARMS EQ 125MG BASE/5ML A061529 001
EQ 250MG BASE/5ML A061529 002CHARTWELL RX EQ 125MG BASE/5ML A062981 001 Feb 10, 1989
EQ 250MG BASE/5ML A062981 002 Feb 10, 1989MYLAN EQ 125MG BASE/5ML A061624 002
EQ 250MG BASE/5ML A061624 001PUREPAC PHARM EQ 125MG BASE/5ML A061758 001
EQ 250MG BASE/5ML A061758 002

PFIZERPEN VK

PFIZER EQ 125MG BASE/5ML A061815 001
EQ 250MG BASE/5ML A061815 002

V-CILLIN K

LILLY EQ 125MG BASE/5ML A060004 001
EQ 250MG BASE/5ML A060004 002

VEETIDS

APOTHECON EQ 125MG BASE/5ML A061410 001
EQ 250MG BASE/5ML A061410 002

VEETIDS '125'

APOTHECON EQ 125MG BASE/5ML A061206 001
EQ 125MG BASE/5ML A062153 001

VEETIDS '250'

APOTHECON EQ 250MG BASE/5ML A061206 002
EQ 250MG BASE/5ML A062153 002

TABLET;ORAL

BEEPEN-VK

GLAXOSMITHKLINE EQ 250MG BASE A062273 001
EQ 500MG BASE A062273 002

BETAPEN-VK

BRISTOL EQ 250MG BASE A061150 001
EQ 500MG BASE A061150 002

LEDERCILLIN VK

LEDERLE EQ 250MG BASE A060134 001
EQ 500MG BASE A060134 002

PEN-VEE K

WYETH AYERST EQ 125MG BASE A060006 001
EQ 250MG BASE A060006 002
EQ 500MG BASE A060006 003

PENAPAR-VK

PARKE DAVIS EQ 250MG BASE A062001 001
EQ 500MG BASE A062001 002

PENICILLIN V POTASSIUM

BELCHER PHARMS EQ 250MG BASE A061528 001
EQ 500MG BASE A061528 002IVAX SUB TEVA PHARMS EQ 125MG BASE A060518 001
EQ 250MG BASE A060518 002
EQ 500MG BASE A060518 003

MYLAN EQ 250MG BASE A061530 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN V POTASSIUM

TABLET; ORAL

PENICILLIN V POTASSIUM

	EQ 500MG BASE	A061530 002
PUREPAC PHARM	EQ 125MG BASE	A061571 001
	EQ 250MG BASE	A061571 002
	EQ 500MG BASE	A061571 003
PFIZERPEN VK		
PFIZER	EQ 250MG BASE	A061836 001
	EQ 500MG BASE	A061836 002
UTICILLIN VK		
PHARMACIA AND UPJOHN	EQ 250MG BASE	A061651 001
	EQ 500MG BASE	A061651 002
V-CILLIN K		
LILLY	EQ 125MG BASE **	A060003 001
	EQ 250MG BASE **	A060003 002
	EQ 500MG BASE **	A060003 003
VEETIDS		
APOTHECON	EQ 250MG BASE	A061411 001
	EQ 500MG BASE	A061411 002
VEETIDS '250'		
APOTHECON	EQ 250MG BASE	A061164 001
	EQ 250MG BASE	A062156 002
VEETIDS '500'		
APOTHECON	EQ 500MG BASE	A061164 002
	EQ 500MG BASE	A062156 001

PENTAGASTRIN

INJECTABLE; INJECTION

PEPTAVLON

+ WYETH AYERST 0.25MG/ML ** N017048 001

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

FRESENIUS KABI USA	600MG/VIAL	N019887 002	Mar 22, 1996
INJECTABLE; INJECTION			
PENTACARINAT			
ARMOUR PHARM	300MG/VIAL	A073447 001	Apr 28, 1994
PENTAMIDINE ISETHIONATE			
BAXTER HLTHCARE	300MG/VIAL	A073617 001	Dec 18, 1995
HOSPIRA	300MG/VIAL	A073479 001	Jun 30, 1992
WATSON LABS	300MG/VIAL	A074303 001	Aug 17, 1995

PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN 50

SANOFI AVENTIS US EQ 50MG BASE N016732 001

PENTAZOCINE LACTATE

INJECTABLE; INJECTION

TALWIN

+ HOSPIRA EQ 30MG BASE/ML N016194 001

PENTETATE CALCIUM TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE CALCIUM TRISODIUM

+ HAMELN EQ 1GM BASE/5ML (EQ 200MG BASE/ML) ** N021749 001 Aug 11, 2004

PENTETATE CALCIUM TRISODIUM YB-169

INJECTABLE; INJECTION

YTTERBIUM YB 169 DTPA

3M 2mCi/ML N017518 001

PENTETATE ZINC TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE ZINC TRISODIUM

+ HAMELN PHARMA PLUS EQ 1GM BASE/5ML (EQ 200MG BASE/ML) ** N021751 001 Aug 11, 2004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENTOBARBITAL

ELIXIR; ORAL

NEMBUTAL

AKORN 18.2MG/5ML A083244 001

PENTOBARBITAL SODIUM

CAPSULE; ORAL

NEMBUTAL SODIUM

AKORN 30MG A084095 001

50MG A084093 001

100MG A083245 001

PENTOBARBITAL SODIUM

LANNETT 50MG A085937 001

100MG A085915 001

VITARINE 100MG A083284 001

WHITEWORTH TOWN PLSN 100MG A083338 001

SODIUM PENTOBARBITAL

ANABOLIC 100MG A084590 001

ELKINS SINN 100MG A083368 001

EVERYLIFE 100MG A083259 001

HALSEY 100MG A084677 001

IVAX SUB TEVA PHARMS 50MG A083461 001

100MG A083461 002

PARKE DAVIS 100MG A084156 001

PERRIGO 100MG A084560 001

PUREPAC PHARM 100MG A083301 001

VALEANT PHARM INTL 100MG A083264 001

WATSON LABS 100MG A085791 001

WYETH AYERST 100MG A083239 001

INJECTABLE; INJECTION

PENTOBARBITAL SODIUM

ELKINS SINN 50MG/ML A083270 001

SODIUM PENTOBARBITAL

WYETH AYERST 50MG/ML A083261 001

SUPPOSITORY; RECTAL

NEMBUTAL

AKORN 30MG A083247 001 Jan 25, 1982

60MG A083247 002 Jan 25, 1982

120MG A083247 003 Jan 25, 1982

200MG A083247 004 Jan 25, 1982

TABLET; ORAL

PENTOBARBITAL SODIUM

VITARINE 100MG A083285 001

SODIUM PENTOBARBITAL

NEXGEN PHARMA INC 100MG A084238 001

PENTOLINIUM TARTRATE

INJECTABLE; INJECTION

ANSOLYSEN

WYETH AYERST 10MG/ML N009372 001

PENTOSTATIN

INJECTABLE; INJECTION

PENTOSTATIN

RISING 10MG/VIAL A203554 001 Sep 19, 2014

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

ANI PHARMS 400MG A074878 001 Jul 09, 1997

400MG A075107 001 Sep 04, 1998

400MG A075199 001 Sep 03, 1999

HERITAGE PHARMS 400MG A074877 001 Jul 08, 1997

IMPAX LABS 400MG A075093 001 Aug 10, 1999

PLIVA 400MG A074874 001 May 25, 1999

RISING 400MG A074425 001 Jul 08, 1997

PENTOXIL

UPSHER SMITH LABS 400MG A074962 001 Mar 31, 1999

TRENAL

+ VALIDUS PHARMS 400MG ** N018631 001 Aug 30, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PERFLUBRONLIQUID; ORAL
IMAGENT

ALLIANCE PHARM 100% N020091 001 Aug 13, 1993

PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE

PASTE; TOPICAL

SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS

US ARMY MED RES 50%;50% N021084 001 Feb 17, 2000

PERGOLIDE MESYLATE

TABLET; ORAL

PERGOLIDE MESYLATE

IVAX SUB TEVA PHARMS EQ 0.05MG BASE A076094 001 Sep 04, 2003

EQ 0.25MG BASE A076094 002 Sep 04, 2003

EQ 1MG BASE A076094 003 Sep 04, 2003

STRIDES PHARMA EQ 0.05MG BASE A076061 001 Nov 27, 2002

EQ 0.25MG BASE A076061 002 Nov 27, 2002

EQ 1MG BASE A076061 003 Nov 27, 2002

PERMAX

VALEANT PHARM INTL EQ 0.05MG BASE N019385 001 Dec 30, 1988

EQ 0.25MG BASE N019385 002 Dec 30, 1988

EQ 1MG BASE N019385 003 Dec 30, 1988

PERINDOPRIL ERBUMINE

TABLET; ORAL

ACEON

+ SYMPLMED PHARMS LLC 2MG N020184 001 Dec 30, 1993

+ 4MG N020184 002 Dec 30, 1993

+ 8MG N020184 003 Dec 30, 1993

PERINDOPRIL ERBUMINE

ANI PHARMS 2MG A078138 001 Nov 10, 2009

4MG A078138 002 Nov 10, 2009

8MG A078138 003 Nov 10, 2009

APOTEX 2MG A090463 001 Aug 30, 2010

4MG A090463 002 Aug 30, 2010

8MG A090463 003 Aug 30, 2010

HIKMA 2MG A090072 001 Nov 10, 2009

4MG A090072 002 Nov 10, 2009

8MG A090072 003 Nov 10, 2009

LUPIN LTD 2MG A078263 001 Jan 27, 2010

4MG A078263 002 Jan 27, 2010

8MG A078263 003 Jan 27, 2010

PERMETHRIN

LOTION; TOPICAL

NIX

GLAXOSMITHKLINE 1% ** N019435 001 Mar 31, 1986

PERPHENAZINE

CONCENTRATE; ORAL

PERPHENAZINE

PHARM ASSOC 16MG/5ML A040360 001 May 25, 2001

TRILAFON

SCHERING 16MG/5ML N011557 001

INJECTABLE; INJECTION

TRILAFON

SCHERING 5MG/ML N011213 002

SYRUP; ORAL

TRILAFON

SCHERING 2MG/5ML N011294 002

TABLET; ORAL

PERPHENAZINE

ANI PHARMS 2MG ** A089707 001 Sep 10, 1987

4MG ** A089708 001 Sep 10, 1987

8MG A089456 001 Sep 10, 1987

16MG A089457 001 Sep 10, 1987

MYLAN 2MG A206691 001 Apr 14, 2017

4MG A206691 002 Apr 14, 2017

8MG A206691 003 Apr 14, 2017

16MG A206691 004 Apr 14, 2017

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PERPHENAZINE

TABLET; ORAL

TRILAFON

+	SCHERING	2MG **	N010775 001
+		4MG **	N010775 002
+		8MG **	N010775 003
+		16MG **	N010775 004

TABLET, EXTENDED RELEASE; ORAL

TRILAFON

	SCHERING	8MG	N011361 002
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PEXIDARTINIB HYDROCHLORIDE

CAPSULE; ORAL

TURALIO

+	DAIICHI SANKYO INC	EQ 200MG BASE	N211810 001 Aug 02, 2019
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PHENACEMIDE

TABLET; ORAL

PHENURONE

+	ABBVIE	500MG **	N007707 001
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PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE

TABLET; ORAL

AZO GANTANOL

+	ROCHE	100MG; 500MG **	N013294 001 Sep 10, 1987
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PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM AND PHENAZOPYRIDINE HYDROCHLORIDE

	ABLE	200MG, N/A, N/A; N/A, 800MG, 160MG	N021105 001 Jun 26, 2001
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PHENAZOPYRIDINE HYDROCHLORIDE; SULFISOXAZOLE

TABLET; ORAL

AZO GANTRISIN

+	ROCHE	50MG; 500MG **	N019358 001 Aug 31, 1990
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PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENAZINE

	MAST MM	35MG	A086523 001
		35MG	A086524 001
		35MG	A086525 001

PHENDIMETRAZINE TARTRATE

	SANDOZ	35MG	A085633 001
		35MG	A085694 001
		35MG	A085702 001
	VIRTUS	35MG	A085695 001
	VITARINE	35MG	A085634 001
		35MG	A085645 001
		35MG	A085670 001
		35MG	A086403 001
		35MG	A086408 001
		35MG	A086410 001
		35MG	A087424 001

SPRX-3

	SOLVAY	35MG	A085897 001
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STATOBEX

	TEVA	35MG	A085507 001
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X-TROZINE

	SHIRE RICHWOOD	35MG	A087394 001 Sep 22, 1982
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CAPSULE, EXTENDED RELEASE; ORAL

BONTRIL

	VALEANT	105MG	A088021 001 Sep 21, 1982
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MELFIAT-105

	NUMARK	105MG	A087487 001 Oct 13, 1982
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PHENDIMETRAZINE TARTRATE

	GRAHAM DM	105MG	A087214 001 May 26, 1982
		105MG	A088020 001 Aug 16, 1982
		105MG	A088028 001 Aug 16, 1982
		105MG	A088062 001 Sep 13, 1982
		105MG	A088063 001 Sep 10, 1982
		105MG	A088111 001 Oct 18, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

PHENDIMETRAZINE TARTRATE

VIRTUS 105MG

A087378 001

SPRX-105

NUMARK 105MG

A088024 001 Dec 22, 1982

X-TROZINE L.A.

SHIRE RICHWOOD 105MG

A087371 001 Aug 24, 1982

TABLET;ORAL

ADPHEN

FERNDALE LABS 35MG

A083655 001

ALPHAZINE

SANDOZ 35MG

A085034 001

CAM-METRAZINE

ABC HOLDING 35MG

A085511 001

CAMALL 35MG

A085756 001

CHARTWELL RX 35MG

A083922 001

35MG

A085318 001

35MG

A085320 001

35MG

A085321 001

DI-METREX

PVT FORM 35MG

A085698 001

MELFIAT

NUMARK 35MG

A083790 002

METRA

FOREST PHARMS 35MG

A083754 001

PHENAZINE

MAST MM 35MG

A087305 001

PHENAZINE-35

ABC HOLDING 35MG

A085512 001

PHENDIMETRAZINE TARTRATE

BARR 35MG

A083644 001

35MG

A083684 001

35MG

A083686 001

35MG

A083687 001

35MG

A084831 001

35MG

A084834 001

35MG

A084835 001

CHARTWELL RX 35MG

A085761 001

35MG

A085941 001 Jun 27, 1983

FERNDALE LABS 35MG

A086834 001 Sep 15, 1983

INWOOD LABS 35MG

A084740 001

35MG

A084741 001

35MG

A084742 001

35MG

A084743 001

IVAX PHARMS 35MG

A085611 001

35MG

A085612 001

IVAX SUB TEVA PHARMS 35MG

A083682 001

KV PHARM 35MG

A084138 001

35MG

A084141 001

35MG

A085525 001

MFG CHEMISTS 35MG

A085914 001

NEXGEN PHARMA INC 35MG

A086020 001

NOSTRUM LABS INC 35MG

A203600 001 Dec 27, 2017

NUMARK 35MG

A083790 001

PVT FORM 35MG

A085199 001

35MG

A085697 001

SANDOZ 35MG

A085402 001

35MG

A085830 001

35MG

A086370 001

SOLVAY 35MG

A083993 001

UPSHER SMITH LABS 35MG

A084399 001

USL PHARMA 35MG

A083805 001

35MG

A084398 001

VIRTUS 35MG

A085497 001

35MG

A086365 001

VITARINE 35MG

A085519 001

35MG

A086005 001

35MG

A086106 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

TABLET; ORAL

PHENDIMETRAZINE TARTRATE

WATSON LABS

35MG

A085767 001

35MG

A085768 001

35MG

A085770 001

35MG

A085773 001

PLEGINE

WYETH AYERST

35MG **

N012248 001

STATOBEX

TEVA

35MG

A086013 001

STATOBEX-G

TEVA

35MG

A085095 001

X-TROZINE

SHIRE RICHWOOD

35MG

A086550 001

35MG

A086551 001

35MG

A086552 001

35MG

A086553 001

35MG

A086554 001

PHENINDIONE

TABLET; ORAL

HEDULIN

SANOFI AVENTIS US

50MG

N008767 002

PHENMETRAZINE HYDROCHLORIDE

TABLET; ORAL

PRELUDIN

BOEHRINGER INGELHEIM

25MG

N010460 005

TABLET, EXTENDED RELEASE; ORAL

PRELUDIN

BOEHRINGER INGELHEIM

50MG

N011752 004

75MG

N011752 003

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENOXYBENZAMINE HYDROCHLORIDE

HIKMA

10MG

A201050 001 Jul 16, 2012

PHENPROCOUMON

TABLET; ORAL

LIQUAMAR

ORGANON USA INC

3MG

N011228 001

PHENSUXIMIDE

CAPSULE; ORAL

MILONTIN

PARKE DAVIS

500MG

N008855 004

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

FASTIN

GLAXOSMITHKLINE

30MG **

N017352 001

OBESTIN-30

FERNDALE LABS

30MG

A087144 001

OBY-TRIM

SHIRE RICHWOOD

30MG

A087764 001 Mar 18, 1982

ONA-MAST

MAST MM

30MG

A086511 001

30MG

A086516 001

PHENTERMINE HYDROCHLORIDE

ABC HOLDING

30MG

A085411 001

ABLE

15MG

A040497 001 Mar 13, 2003

30MG

A040403 001 Aug 30, 2001

30MG

A040427 001 Aug 30, 2001

BARR

15MG

A090591 001 Mar 18, 2010

30MG

A090591 002 Mar 18, 2010

CAMALL

15MG

A086735 001

30MG

A087226 001

CHARTWELL RX

18.75MG

A088576 001 May 23, 1984

30MG

A085417 001

30MG

A086732 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

	30MG	A087215	001	
	37.5MG	A087915	001	Dec 22, 1983
	37.5MG	A087918	001	Dec 22, 1983
	37.5MG	A087930	001	Oct 14, 1983
	37.5MG	A088610	001	Jun 04, 1984
	37.5MG	A088611	001	Jun 04, 1984
	37.5MG	A088625	001	Aug 23, 1984
DURAMED PHARMS BARR	30MG	A088948	001	Apr 25, 1986
ELITE LABS INC	15MG	A040460	001	Jan 14, 2003
	30MG	A040227	001	Jun 18, 1997
	30MG	A040448	001	Jan 22, 2003
IVAX PHARMS	30MG	A086329	001	
LANNETT	15MG	A087022	002	Jan 20, 2012
	30MG	A087022	001	Feb 03, 1983
LANNETT CO INC	30MG	A091359	001	Jul 16, 2010
	37.5MG	A201961	001	Jul 20, 2011
SANDOZ	30MG	A087208	001	
	30MG	A087223	001	
	37.5MG	A088414	001	Oct 19, 1983
SUN PHARM INDUSTRIES	30MG	A040525	001	Oct 23, 2003
	37.5MG	A040527	001	Oct 23, 2003
TEVA	30MG	A086911	001	
	30MG	A087126	001	
	30MG	A087777	001	Nov 01, 1985
	30MG	A088612	001	Apr 04, 1984
	30MG	A088613	001	Apr 09, 1984
	30MG	A088614	001	Apr 09, 1984
TG UNITED INC	30MG	A040083	001	Mar 07, 1997
UPSHER SMITH LABS	30MG	A084487	001	Apr 09, 1982
	30MG	A088430	001	Mar 27, 1984
USL PHARMA	30MG	A088797	001	Dec 10, 1984
VITARINE	30MG	A087202	001	
	30MG	A087235	001	
WATSON LABS	30MG	A086740	001	Mar 21, 1985
TABLET; ORAL				
ONA-MAST				
MAST MM	8MG	A086260	001	
PHENTERMINE HYDROCHLORIDE				
ABLE	37.5MG	A040402	001	Aug 30, 2001
ACTAVIS ELIZABETH	37.5MG	A040276	001	Nov 25, 1998
BARR	37.5MG	A090470	001	Aug 31, 2009
CHARTWELL RX	8MG	A083923	001	
	8MG	A085319	001	
	37.5MG	A087805	001	Dec 06, 1982
	37.5MG	A088596	001	Apr 04, 1984
IVAX PHARMS	8MG	A085553	001	
LANNETT	37.5MG	A040555	001	Apr 15, 2005
NOVAST LABS	37.5MG	A091451	001	Sep 21, 2012
SANDOZ	8MG	A085671	001	
	8MG	A085689	001	
SANDOZ INC	30MG	A088605	001	Sep 28, 1987
SUN PHARM INDS INC	37.5MG	A040790	001	Aug 21, 2007
+ USL PHARMA	8MG	A083804	001	
	37.5MG	A088910	001	Jul 17, 1985
	37.5MG	A088917	001	Jul 17, 1985
VITARINE	8MG	A086453	001	
	8MG	A086456	001	
WATSON LABS	8MG	A085739	001	
TORA				
SOLVAY	8MG	A084035	001	
WILPO				
+ SANDOZ	8MG **	N012737	001	
TABLET, ORALLY DISINTEGRATING; ORAL				
SUPRENZA				
CITIUS PHARMS	15MG **	N202088	001	Jun 13, 2011
	30MG **	N202088	002	Jun 13, 2011

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENTERMINE HYDROCHLORIDETABLET, ORALLY DISINTEGRATING;ORAL
SUPRENZA

37.5MG **

N202088 003 Mar 27, 2012

PHENTERMINE RESIN COMPLEXCAPSULE, EXTENDED RELEASE;ORAL
IONAMIN

UCB INC

EQ 15MG BASE **

N011613 004

EQ 30MG BASE **

N011613 002

PHENTERMINE RESIN 30

QUANTUM PHARMICS

EQ 30MG BASE

A089120 001 Feb 04, 1988

PHENTERMINE RESIN COMPLEX

LANNETT CO INC

EQ 15MG BASE

A040872 001 Jul 28, 2011

EQ 30MG BASE

A040872 002 Jul 28, 2011

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

REGITINE

+ NOVARTIS

5MG/VIAL **

N008278 003

PHENYL AMINOSALICYLATE

POWDER;ORAL

PHENY-PAS-TEBAMIN

PHARM RES ASSOC

50%

N011695 002

TABLET;ORAL

PHENY-PAS-TEBAMIN

PHARM RES ASSOC

500MG

N011695 003

PHENYLBUTAZONE

CAPSULE;ORAL

AZOLID

SANOFI AVENTIS US

100MG

A087260 001

BUTAZOLIDIN

NOVARTIS

100MG

N008319 009

PHENYLBUTAZONE

FOSUN PHARMA

100MG

A087774 001 Jun 16, 1982

IVAX PHARMS

100MG

A088218 001 Jun 24, 1983

SUN PHARM INDUSTRIES

100MG

A088994 001 Dec 04, 1985

WATSON LABS

100MG

A087756 001 Dec 17, 1982

TABLET;ORAL

AZOLID

SANOFI AVENTIS US

100MG

A087091 001

BUTAZOLIDIN

NOVARTIS

100MG

N008319 008

PHENYLBUTAZONE

FOSUN PHARMA

100MG

A084339 001

SUN PHARM INDUSTRIES

100MG

A088863 001 Dec 04, 1985

WATSON LABS

100MG

A086151 001

100MG

A087674 001 Apr 21, 1982

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

BIORPHEN

+ DR REDDYS LABS SA

10MG/ML (10MG/ML)

N212909 002 Mar 11, 2021

PHENYLEPHRINE HYDROCHLORIDE

PAR STERILE PRODUCTS

10MG/ML (10MG/ML)

A210025 001 Dec 21, 2018

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN VC

+ ANI PHARMS

5MG/5ML;6.25MG/5ML **

N008604 003 Apr 02, 1984

PHERAZINE VC

HALSEY

5MG/5ML;6.25MG/5ML

A088868 001 Mar 02, 1987

PROMETH VC PLAIN

+ G AND W LABS INC

5MG/5ML;6.25MG/5ML

A088761 001 Nov 08, 1984

PROMETHAZINE VC PLAIN

CENCI

5MG/5ML;6.25MG/5ML

A088815 001 Nov 22, 1985

XTTRIUM LABS INC

5MG/5ML;6.25MG/5ML

A088897 001 Jan 04, 1985

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENYLEPHRINE HYDROCHLORIDE; PYRILAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC

PREFRIN-A

ALLERGAN 0.12%;0.1% N007953 001

PHENYTOIN

SUSPENSION;ORAL

DILANTIN-30

VIATRIS 30MG/5ML N008762 002

PHENYTOIN

ACTAVIS MID ATLANTIC 125MG/5ML A089892 001 Sep 25, 1992

PHENYTOIN SODIUM

CAPSULE;ORAL

DIPHENYLAN SODIUM

CHARTWELL MOLECULAR 30MG PROMPT A080857 001

100MG PROMPT A080857 002

EXTENDED PHENYTOIN SODIUM

ANI PHARMS 100MG EXTENDED A040435 001 Jun 20, 2003

100MG EXTENDED A089441 001 Dec 18, 1986

LUPIN LTD 100MG EXTENDED A211633 001 Sep 30, 2019

MYLAN 100MG EXTENDED A040298 001 Dec 28, 1998

SUN PHARM INDS (IN) 100MG EXTENDED A040621 001 Dec 11, 2006

UNICHEM 100MG EXTENDED A213834 001 Oct 13, 2022

WOCKHARDT 30MG EXTENDED A040759 001 Dec 18, 2007

WOCKHARDT USA 100MG EXTENDED A040732 001 Jan 30, 2008

PHENYTEX

WATSON LABS 100MG EXTENDED A088711 001 Dec 21, 1984

PHENYTOIN SODIUM

PHARMERAL 100MG PROMPT A085435 001

WATSON LABS 100MG PROMPT A085894 001

PROMPT PHENYTOIN SODIUM

ANI PHARMS 100MG PROMPT A080259 001

WATSON LABS 100MG PROMPT A080905 001

INJECTABLE;INJECTION

DILANTIN

PARKE DAVIS 50MG/ML N010151 001

PHENYTOIN SODIUM

AM REGENT 50MG/ML A040781 001 Dec 04, 2007

FRESENIUS KABI USA 50MG/ML A089003 001 May 31, 1985

HOSPIRA 50MG/ML A089521 001 Mar 17, 1987

50MG/ML A089744 001 Dec 18, 1987

MARSAM PHARMS LLC 50MG/ML A089501 001 Oct 13, 1987

50MG/ML A089779 001 Nov 27, 1992

SMITH AND NEPHEW 50MG/ML A088519 001 Dec 19, 1984

50MG/ML A088521 001 Dec 18, 1984

SOLOPAK 50MG/ML A088520 001 Dec 17, 1984

WARNER CHILCOTT 50MG/ML A089900 001 Mar 30, 1990

WATSON LABS 50MG/ML A085434 001

PHYTONADIONE

INJECTABLE;INJECTION

AQUAMEPHYTON

+ PAI HOLDINGS PHARM 1MG/0.5ML ** N012223 002

+ 10MG/ML ** N012223 001

KONAKION

ROCHE 1MG/0.5ML N011745 001

10MG/ML N011745 003

PHYTONADIONE

CIPLA 1MG/0.5ML A212424 001 Apr 22, 2022

10MG/ML A212424 002 Apr 22, 2022

GLAXOSMITHKLINE 1MG/0.5ML A084060 001

10MG/ML A084060 002

VITAMIN K1

HOSPIRA 10MG/ML A087956 001 Jul 25, 1983

TABLET;ORAL

MEPHYTON

+ BAUSCH 5MG ** N010104 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PILOCARPINE

INSERT, EXTENDED RELEASE;OPHTHALMIC

OCUSERT PILO-20

EPIC PHARMA LLC

5MG

N017431 001

OCUSERT PILO-40

EPIC PHARMA LLC

11MG

N017548 001

PILOCARPINE HYDROCHLORIDE

GEL;OPHTHALMIC

PILOPINE HS

ALCON

4%

N018796 001 Oct 01, 1984

PIMAVANSERIN TARTRATE

TABLET;ORAL

NUPLAZID

+ ACADIA PHARMS INC

EQ 17MG BASE

N207318 001 Apr 29, 2016

PIMOZIDE

TABLET;ORAL

ORAP

+ TEVA

1MG **

N017473 003 Aug 27, 1997

+

2MG **

N017473 001 Jul 31, 1984

PINACIDIL

CAPSULE, EXTENDED RELEASE;ORAL

PINDAC

LEO PHARM

12.5MG

N019456 001 Dec 28, 1989

25MG

N019456 002 Dec 28, 1989

PINDOLOL

TABLET;ORAL

PINDOLOL

COSETTE

5MG

A073661 001 Oct 31, 1993

5MG

A073687 001 Feb 26, 1993

5MG

A074123 001 Apr 17, 1997

10MG

A073661 002 Oct 31, 1993

10MG

A073687 002 Feb 26, 1993

10MG

A074123 002 Apr 17, 1997

MYLAN PHARMS INC

5MG

A074013 001 Sep 24, 1992

10MG

A074018 001 Sep 24, 1992

NOSTRUM LABS

5MG

A074474 001 Oct 28, 1996

10MG

A074474 002 Oct 28, 1996

PUREPAC PHARM

5MG

A074125 001 Apr 28, 1993

10MG

A074125 002 Apr 28, 1993

WATSON LABS

5MG

A074437 001 Feb 27, 1995

10MG

A074437 002 Feb 27, 1995

ZYDUS PHARMS

5MG

A209866 001 Aug 18, 2017

10MG

A209866 002 Aug 18, 2017

VISKEN

+ NOVARTIS

5MG **

N018285 001 Sep 03, 1982

+

10MG **

N018285 002 Sep 03, 1982

PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

PIOGLITAZONE HYDROCHLORIDE

CHARTWELL RX

EQ 15MG BASE

A078383 001 Mar 12, 2013

EQ 30MG BASE

A078383 002 Mar 12, 2013

EQ 45MG BASE

A078383 003 Mar 12, 2013

MYLAN PHARMS INC

EQ 15MG BASE

A076801 001 Aug 17, 2012

EQ 30MG BASE

A076801 002 Aug 17, 2012

EQ 45MG BASE

A076801 003 Aug 17, 2012

NOSTRUM LABS INC

EQ 15MG BASE

A078472 001 Feb 13, 2013

EQ 30MG BASE

A078472 002 Feb 13, 2013

EQ 45MG BASE

A078472 003 Feb 13, 2013

TORRENT PHARMS LTD

EQ 15MG BASE

A091298 001 Feb 13, 2013

EQ 30MG BASE

A091298 002 Feb 13, 2013

EQ 45MG BASE

A091298 003 Feb 13, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PIPECURONIUM BROMIDE

INJECTABLE; INJECTION

ARDUAN

ORGANON USA INC 10MG/VIAL

N019638 001 Jun 26, 1990

PIPERACETAZINE

TABLET; ORAL

QUIDE

DOW PHARM 10MG
25MGN013615 001
N013615 002PIPERACILLIN SODIUM

INJECTABLE; INJECTION

PIPRACIL

WYETH PHARMS INC

EQ 2GM BASE/VIAL

A062750 001 Oct 13, 1987

+ EQ 2GM BASE/VIAL **

N050545 002

EQ 3GM BASE/VIAL

A062750 002 Oct 13, 1987

+ EQ 3GM BASE/VIAL **

N050545 003

EQ 4GM BASE/VIAL

A062750 003 Oct 13, 1987

+ EQ 4GM BASE/VIAL **

N050545 004

+ EQ 40GM BASE/VIAL **

N050545 006 Sep 30, 1985

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

HOSPIRA INC

EQ 2GM BASE/VIAL; EQ 250MG BASE/VIAL

A065386 001 Sep 15, 2009

EQ 3GM BASE/VIAL; EQ 375MG BASE/VIAL

A065386 002 Sep 15, 2009

EQ 4GM BASE/VIAL; EQ 500MG BASE/VIAL

A065386 003 Sep 15, 2009

EQ 36GM BASE/VIAL; EQ 4.5GM BASE/VIAL

A065446 001 Sep 15, 2009

ZOSYN

+ WYETH PHARMS

EQ 2GM BASE/VIAL; EQ 250MG BASE/VIAL **

N050684 001 Oct 22, 1993

+ EQ 3GM BASE/VIAL; EQ 375MG BASE/VIAL **

N050684 002 Oct 22, 1993

+ EQ 4GM BASE/VIAL; EQ 500MG BASE/VIAL **

N050684 003 Oct 22, 1993

+ EQ 36GM BASE/VIAL; EQ 4.5GM BASE/VIAL **

N050684 004 Oct 22, 1993

PIPERAZINE CITRATE

SYRUP; ORAL

ANTEPAR

GLAXOSMITHKLINE

EQ 500MG BASE/5ML

N009102 001

BRYREL

SANOFI AVENTIS US

EQ 500MG BASE/5ML

N017796 001

MULTIFUGE

BLULINE

EQ 500MG BASE/5ML

N009452 001

PIPERAZINE CITRATE

ALPHARMA US PHARMS

EQ 500MG BASE/5ML

A080774 001

LANNETT

EQ 500MG BASE/5ML

A080963 001

LUITPOLD

EQ 500MG BASE/5ML

A080671 001

VERMIDOL

SOLVAY

EQ 500MG BASE/5ML

A080992 001

TABLET; ORAL

ANTEPAR

GLAXOSMITHKLINE

EQ 500MG BASE

N009102 003

PIPERAZINE CITRATE

IMPAX LABS

EQ 250MG BASE

A080874 001

PIPERONYL BUTOXIDE; PYRETHRINS

AEROSOL; TOPICAL

RID MOUSSE

BAYER HEALTHCARE LLC 4%; EQ 0.33% BASE

N021043 001 Mar 07, 2000

PIPOBROMAN

TABLET; ORAL

VERCYTE

ABBOTT

10MG

N016245 001

25MG

N016245 002

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

BAUSCH

EQ 0.2MG BASE/INH

N019009 001 Dec 30, 1986

EQ 0.2MG BASE/INH

N020014 001 Nov 30, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PIRFENIDONE

CAPSULE; ORAL

PIRFENIDONE

APOTEX

267MG

A212687 001 Jun 09, 2023

LUPIN LTD

267MG

A212404 001 Aug 28, 2023

TABLET; ORAL

ESBRIET

+ GENENTECH INC

534MG **

N208780 002 Jan 11, 2017

PIRFENIDONE

LUPIN LTD

267MG

A212403 001 Aug 23, 2023

801MG

A212403 002 Aug 23, 2023

PIROXICAM

CAPSULE; ORAL

PIROXICAM

BRECKENRIDGE

10MG

A208991 001 Feb 21, 2018

20MG

A208991 002 Feb 21, 2018

CYCLE

10MG

A073651 001 Feb 26, 1993

20MG

A073651 002 Feb 26, 1993

EGIS

10MG

A074808 001 Jul 08, 1997

20MG

A074808 002 Jul 08, 1997

HIKMA

10MG

A209256 001 Aug 11, 2017

20MG

A209256 002 Aug 11, 2017

IVAX SUB TEVA PHARMS

10MG

A074148 001 Jun 03, 1996

20MG

A074148 002 Jun 03, 1996

MYLAN

10MG

A074043 001 Sep 22, 1992

10MG

A074102 001 Jul 31, 1992

20MG

A074043 002 Sep 22, 1992

20MG

A074102 002 Jul 31, 1992

SCS

10MG

A074036 001 May 29, 1992

20MG

A074036 002 May 29, 1992

SUN PHARM INDUSTRIES

10MG

A073536 002 Jan 23, 2008

20MG

A073536 001 Mar 12, 1993

TEVA

10MG

A073637 001 Jan 28, 1994

20MG

A073638 001 Jan 28, 1994

TEVA PHARMS

10MG

A074103 001 Aug 28, 1992

20MG

A074103 002 Aug 28, 1992

WATSON LABS

10MG

A074287 001 May 16, 1996

10MG

A074460 001 Sep 29, 1995

20MG

A074287 002 May 16, 1996

20MG

A074460 002 Sep 29, 1995

PITAVASTATIN MAGNESIUM

TABLET; ORAL

ZYPITAMAG

+ MEDICURE

EQ 1MG BASE

N208379 001 Jul 14, 2017

PITAVASTATIN SODIUM

TABLET; ORAL

NIKITA

+ LUPIN LTD

EQ 1MG BASE

N209875 001 Aug 04, 2017

+

EQ 2MG BASE

N209875 002 Aug 04, 2017

+

EQ 4MG BASE

N209875 003 Aug 04, 2017

PLICAMYCIN

INJECTABLE; INJECTION

MITHRACIN

PFIZER

2.5MG/VIAL

N050109 001

PODOFILOX

SOLUTION; TOPICAL

PODOFILOX

BAUSCH AND LOMB INC

0.5%

A090184 001 Jul 21, 2010

POLIDOCANOL

SOLUTION; INTRAVENOUS

VARITHENA

+ PROVENSIS

77.5MG/7.75ML (10MG/ML)

N205098 002 Dec 21, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POLYESTRADIOL PHOSPHATEINJECTABLE; INJECTION
ESTRADURIN

WYETH AYERST 40MG/AMP N010753 001

POLYETHYLENE GLYCOL 3350FOR SOLUTION; ORAL
GLYCOLAXLANNETT CO INC 17GM/SCOOPFUL A076652 001 Jul 02, 2004
17GM/PACKET A090600 001 Oct 06, 2009
17GM/SCOOPFUL A090600 002 Oct 06, 2009

POLYETHYLENE GLYCOL 3350

BRECKENRIDGE PHARM 17GM/SCOOPFUL A077736 001 May 26, 2006
NEXGEN PHARMA INC 17GM/SCOOPFUL A077706 001 Sep 27, 2006
PADDOCK LLC 17GM/SCOOPFUL A077893 001 May 26, 2006
17GM/SCOOPFUL A090567 001 Oct 15, 2009
TEVA PHARMS 17GM/SCOOPFUL A077445 001 May 04, 2006POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

LAX-LYTE WITH FLAVOR PACKS

L PERRIGO CO 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT A079232 001 Feb 25, 2010

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

MYLAN 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT A090409 001 Apr 02, 2010
NOSTRUM LABS INC 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT A202060 001 Mar 08, 2017
NOVEL LABS INC 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT A090019 001 May 27, 2009

TRILYTE

AUROBINDO PHARMA USA 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT A076491 001 Feb 05, 2004

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

CLENZ-LYTE

PADDOCK LLC 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A090769 001 Jun 07, 2010

SOLUTION; ORAL

OCL

HOSPIRA 6GM/100ML; 75MG/100ML; 168MG/100ML; 146MG/100ML; 1.29GM/100ML N019284 001 Apr 30, 1986

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION; ORAL

COLYTE

MYLAN SPECIALITY LP 120GM/PACKET; 1.49GM/PACKET; 3.36GM/PACKET; 2.92GM/PACKET; 11.36GM/PACKET ** N018983 005 Oct 26, 1984
227.1GM/PACKET; 2.82GM/PACKET; 6.36GM/PACKET; 5.53GM/PACKET; 21.5GM/PACKET ** N018983 004 Oct 26, 1984
227.1GM/BOT; 2.82GM/BOT; 6.36GM/BOT; 5.53GM/BOT; 21.5GM/BOT ** N018983 010 Jan 31, 1989
240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT ** N018983 007 Jun 12, 1987
360GM/PACKET; 4.47GM/PACKET; 10.08GM/PACKET; 8.76GM/PACKET; 34.08GM/PACKET ** N018983 006 Oct 26, 1984

COLYTE-FLAVORED

MYLAN SPECIALITY LP 227.1GM/BOT; 2.82GM/BOT; 6.36GM/BOT; 5.53GM/BOT; 21.5GM/BOT ** N018983 008 Nov 14, 1991
240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT ** N018983 009 Nov 14, 1991

GOLYTELY

+ BRAINTREE 227.1GM/PACKET; 2.82GM/PACKET; 6.36GM/PACKET; 5.53GM/PACKET; 21.5GM/PACKET N019011 002 Jun 02, 1992

PEG 3350 AND ELECTROLYTES

MYLAN 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A090928 001 Jan 28, 2010

POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

PADDOCK LLC 240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT A090712 001 Feb 25, 2010

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SUSPENSION; ORAL

CO-LAV

VINTAGE PHARMS	240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT	A073428 001	Jan 28, 1992
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COLOVAGE

DYNAPHARM	227.1GM/PACKET; 2.82GM/PACKET; 6.36GM/PACKET; 5.53GM/PACKET; 21.5GM/PACKET	A071320 001	Apr 20, 1988
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E-Z-EM PREP LYTE

E Z EM	236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	A071278 001	Nov 21, 1988
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GLYCOPREP

GOLDLINE	236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	A072319 001	Dec 23, 1988
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GO-EVAC

VINTAGE PHARMS	236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	A073433 001	Apr 28, 1992
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PEG-LYTE

SANDOZ	236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	A073098 001	Aug 31, 1993
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POLYMYXIN B SULFATE

INJECTABLE; INJECTION

AEROSPORIN

GLAXOSMITHKLINE	EQ 500,000 UNITS BASE/VIAL	A062036 001	
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POLYMYXIN B SULFATE

HIKMA	EQ 500,000 UNITS BASE/VIAL	A060716 001	
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RISING	EQ 500,000 UNITS BASE/VIAL	A090110 001	Jun 29, 2011
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POWDER; FOR RX COMPOUNDING

POLY-RX

X GEN PHARMS	100,000,000 UNITS/BOT	A061578 001	
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POLYMYXIN B SULFATE

PADDOCK LLC	100,000,000 UNITS/BOT	A062455 001	Jul 27, 1983
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POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS; OPHTHALMIC

POLYTRIM

+ ALLERGAN	10,000 UNITS/ML; EQ 1MG BASE/ML	N050567 001	Oct 20, 1988
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POLYTHIAZIDE

TABLET; ORAL

RENESE

PFIZER	1MG	N012845 001	
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	2MG	N012845 002	
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	4MG	N012845 003	
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POLYTHIAZIDE; PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIZIDE

PFIZER	0.5MG; EQ 1MG BASE	N017986 001	
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	0.5MG; EQ 2MG BASE	N017986 002	
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	0.5MG; EQ 5MG BASE	N017986 003	
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POLYTHIAZIDE; RESERPINE

TABLET; ORAL

RENESE-R

PFIZER	2MG; 0.25MG	N013636 001	
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POMALIDOMIDE

CAPSULE; ORAL

POMALIDOMIDE

BRECKENRIDGE	1MG	A210111 001	Oct 30, 2020
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	2MG	A210111 002	Oct 30, 2020
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	3MG	A210111 003	Oct 30, 2020
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	4MG	A210111 004	Oct 30, 2020
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EUGIA PHARMA	1MG	A210249 001	Oct 30, 2020
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	2MG	A210249 002	Oct 30, 2020
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	3MG	A210249 003	Oct 30, 2020
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	4MG	A210249 004	Oct 30, 2020
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MYLAN	1MG	A210275 001	Jan 26, 2022
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	2MG	A210275 002	Jan 26, 2022
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	3MG	A210275 003	Jan 26, 2022
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	4MG	A210275 004	Jan 26, 2022
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POMALIDOMIDE

CAPSULE; ORAL

POMALIDOMIDE

TEVA PHARMS USA	1MG	A209956 001	May 04, 2022
	2MG	A209956 002	May 04, 2022
	3MG	A209956 003	May 04, 2022
	4MG	A209956 004	May 04, 2022

PONATINIB HYDROCHLORIDE

TABLET; ORAL

PONATINIB HYDROCHLORIDE

APOTEX	EQ 15MG BASE	A215893 001	Jul 14, 2023
	EQ 45MG BASE	A215893 002	Jul 14, 2023

POSACONAZOLE

TABLET, DELAYED RELEASE; ORAL

POSACONAZOLE

ACTAVIS LABS FL INC	100MG	A207355 001	Nov 30, 2022
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POTASSIUM AMINOSALICYLATE

CAPSULE; ORAL

PASKALIUM

GLENWOOD	500MG	N009395 004	
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POWDER; ORAL

POTASSIUM AMINOSALICYLATE

HEXCEL	100%	A080098 001	
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TABLET; ORAL

PASKALIUM

GLENWOOD	1GM	N009395 003	
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POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

K-LEASE

SAVAGE LABS	8MEQ	A073398 001	Jan 28, 1992
	10MEQ	A072427 001	Mar 28, 1990

MICRO-K

+ NESHER PHARMS	8MEQ **	N018238 001	
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MICRO-K 10

+ NESHER PHARMS	10MEQ **	N018238 002	May 14, 1984
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POTASSIUM CHLORIDE

CHARTWELL MOLECULAR	8MEQ	A204210 001	Mar 28, 2016
	10MEQ	A204210 002	Mar 28, 2016
NESHER PHARMS	10MEQ	A070980 001	Feb 17, 1987
TEVA	8MEQ	A073531 001	Apr 26, 1996
	10MEQ	A073532 001	Apr 26, 1996
TRIS PHARMA INC	8MEQ	A201944 001	Mar 04, 2016
	10MEQ	A201944 002	Mar 04, 2016

FOR SUSPENSION, EXTENDED RELEASE; ORAL

MICRO-K LS

+ KV PHARM	20MEQ/PACKET **	N019561 003	Aug 26, 1988
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INJECTABLE; INJECTION

POTASSIUM CHLORIDE

+ ABRAXIS PHARM	2MEQ/ML	A080204 001	
	2MEQ/ML	A084290 001	
	2MEQ/ML	A086713 001	
	2MEQ/ML	A086714 001	
	2MEQ/ML	A087787 001	Apr 20, 1982
	2MEQ/ML	A087885 001	Feb 03, 1983
AKORN	2MEQ/ML	A088286 001	Sep 05, 1985
BAXTER HLTHCARE	2MEQ/ML	A080203 001	
	2MEQ/ML	A085499 001	
+ FRESENIUS KABI USA	2MEQ/ML	A080225 001	
	2MEQ/ML	A087817 001	Oct 20, 1982
	3MEQ/ML	A080225 003	
GD SEARLE LLC	1MEQ/ML	A086219 001	
	2MEQ/ML	A086219 002	
	2MEQ/ML	A086220 002	
	3MEQ/ML	A086219 003	
	3MEQ/ML	A086220 001	
	4MEQ/ML	A086219 004	
HOSPIRA	1MEQ/ML	A080205 003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDEINJECTABLE; INJECTION
POTASSIUM CHLORIDE

	1MEQ/ML	A083345	003	
	1.5MEQ/ML	A083345	001	
	2MEQ/ML	A083345	002	
	2.4MEQ/ML	A080205	004	
	3.2MEQ/ML	A080205	005	
INTL MEDICATION	2MEQ/ML	A083163	001	
LILLY	2MEQ/ML	N007865	002	
LUITPOLD	2MEQ/ML	A080221	001	
	2MEQ/ML	A080736	001	
	2MEQ/ML	A087584	001	
	2MEQ/ML	A087585	001	
MILES	1MEQ/ML	A080195	002	
	2MEQ/ML	A080195	001	
	3MEQ/ML	A080195	003	
	4MEQ/ML	A080195	004	
PHARMA SERVE NY	2MEQ/ML	A086297	001	
	2MEQ/ML	A087362	001	Mar 08, 1983
WATSON LABS	2MEQ/ML	A086208	001	
	2MEQ/ML	A089163	001	Mar 10, 1988
	2MEQ/ML	A089421	001	Jan 02, 1987
	3MEQ/ML	A086210	001	
POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER				
+ ICU MEDICAL INC	2.24GM/100ML	N020161	003	Aug 11, 1998
SOLUTION; ORAL				
POTASSIUM CHLORIDE				
STRIDES PHARMA	40MEQ/15ML	A211665	001	Nov 17, 2022
TRIS PHARMA INC	20MEQ/15ML	A214076	001	Jan 26, 2022
	40MEQ/15ML	A214076	002	Jan 26, 2022
TABLET, EXTENDED RELEASE; ORAL				
K+10				
FUTURE PAK	10MEQ	A070999	001	Oct 22, 1987
K+8				
FUTURE PAK	8MEQ	A070998	001	Jan 25, 1993
K-TAB				
+ ABBVIE	8MEQ **	N018279	002	Aug 01, 1988
+	10MEQ **	N018279	001	
+	20MEQ **	N018279	003	Nov 25, 2013
KAON CL				
SAVAGE LABS	6.7MEQ	N017046	001	
KAON CL-10				
SAVAGE LABS	10MEQ	N017046	002	
KLOTRIX				
APOTHECON	10MEQ	N017850	001	
POTASSIUM CHLORIDE				
AMNEAL	15MEQ	A212861	002	May 08, 2020
AUROBINDO PHARMA LTD	10MEQ	A214728	001	Mar 31, 2021
	15MEQ	A214728	002	Mar 31, 2021
	20MEQ	A214728	003	Mar 31, 2021
BRECKENRIDGE	10MEQ	A213588	001	Aug 21, 2020
	20MEQ	A213588	002	Aug 21, 2020
CHARTWELL RX	20MEQ	A210098	001	Apr 26, 2019
COPLEY PHARM	8MEQ	A070618	001	Sep 09, 1987
NESHER PHARMS	20MEQ	A076044	001	Apr 05, 2002
+ SCHERING	10MEQ **	N019439	002	Jun 13, 1986
+	20MEQ **	N019439	001	Jun 13, 1986
SIGMAPHARM LABS LLC	8MEQ	A207528	001	Aug 19, 2016
	10MEQ	A207528	002	Aug 19, 2016
STRIDES PHARMA	8MEQ	A206881	001	Jan 22, 2019
	10MEQ	A206630	001	Mar 29, 2019
	10MEQ	A206881	002	Jan 22, 2019
	10MEQ	A210097	001	Jun 17, 2019
	15MEQ	A206630	002	Mar 29, 2019
	20MEQ	A206630	003	Mar 29, 2019
SLOW-K				
NOVARTIS	8MEQ	N017476	002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDETABLET, EXTENDED RELEASE;ORAL
TEN-K

NOVARTIS 10MEQ N019381 001 Apr 16, 1986

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE;INJECTION

POTASSIUM CHLORIDE 0.037% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 37MG/100ML;900MG/100ML N019708 001 Sep 29, 1989

POTASSIUM CHLORIDE 0.075% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 75MG/100ML;900MG/100ML N019708 002 Sep 29, 1989

POTASSIUM CHLORIDE 0.11% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 110MG/100ML;900MG/100ML N019708 003 Sep 29, 1989

POTASSIUM CHLORIDE 0.22% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 220MG/100ML;900MG/100ML N019708 005 Sep 29, 1989

POTASSIUM CHLORIDE 0.224% IN SODIUM CHLORIDE 0.9%

+ BAXTER HLTHCARE 224MG/100ML;900MG/100ML N017648 003

POTASSIUM CHLORIDE 0.3% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 300MG/100ML;900MG/100ML N019708 006 Sep 29, 1989

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075%

BAXTER HLTHCARE 75MG/100ML;900MG/100ML N017648 004

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER

B BRAUN 75MG/100ML;900MG/100ML N018722 001 Nov 09, 1982

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

B BRAUN 150MG/100ML;900MG/100ML N018722 002 Nov 09, 1982

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER

B BRAUN 220MG/100ML;900MG/100ML N018722 003 Nov 09, 1982

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN 300MG/100ML;900MG/100ML N018722 004 Nov 09, 1982

POTASSIUM CHLORIDE; SODIUM CHLORIDE; TROMETHAMINE

INJECTABLE;INJECTION

THAM-E

HOSPIRA 370MG/VIAL;1.75GM/VIAL;36GM/VIAL N013025 001

POTASSIUM CITRATE

FOR SOLUTION;ORAL

POTASSIUM CITRATE

+ UT SW MEDCTR 10MEQ/PACKET ** N019647 002 Oct 13, 1988

+ 20MEQ/PACKET ** N019647 001 Oct 13, 1988

POTASSIUM IODIDE

SOLUTION;ORAL

POTASSIUM IODIDE

+ ROXANE 1GM/ML ** N018551 001 Feb 19, 1982

THYROSHIELD

ARCO PHARMS LLC 65MG/ML A077218 001 Jan 12, 2005

TABLET;ORAL

THYRO-BLOCK

MEDA PHARMS 130MG N018307 001

POTASSIUM PERCHLORATE

CAPSULE;ORAL

PERCHLORACAP

MALLINCKRODT 200MG N017551 001

POVIDONE-IODINE

SOLUTION;TOPICAL

E-Z PREP

CLINIPAD 10% N019382 001 Jul 25, 1989

SPONGE;TOPICAL

E-Z PREP

CLINIPAD 5% N019382 002 Jul 25, 1989

E-Z PREP 220

CLINIPAD 5% N019382 003 Jul 25, 1989

PRALIDOXIME CHLORIDE

INJECTABLE;INJECTION

PRALIDOXIME CHLORIDE

BAXTER HLTHCARE CORP 300MG/ML N018799 001 Dec 13, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRALIDOXIME CHLORIDE

SOLUTION; INTRAMUSCULAR

PRALIDOXIME CHLORIDE (AUTOINJECTOR)

+ MERIDIAN MEDCL TECHN 600MG/2ML (300MG/ML)

N018986 001 Apr 26, 1983

TABLET; ORAL

PROTOPAM CHLORIDE

WYETH AYERST

500MG

N014122 002

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

MIRAPEX

+ BOEHRINGER INGELHEIM 0.125MG **

N020667 001 Jul 01, 1997

+ 0.25MG **

N020667 002 Jul 01, 1997

+ 0.5MG **

N020667 006 Feb 12, 1998

+ 0.75MG **

N020667 007 Jul 30, 2007

+ 1MG **

N020667 003 Jul 01, 1997

+ 1.25MG

N020667 004 Jul 01, 1997

+ 1.5MG **

N020667 005 Jul 01, 1997

PRAMIPEXOLE DIHYDROCHLORIDE

ALEMBIC

0.125MG

A078894 001 Oct 08, 2010

0.25MG

A078894 002 Oct 08, 2010

0.5MG

A078894 003 Oct 08, 2010

1MG

A078894 004 Oct 08, 2010

1.5MG

A078894 005 Oct 08, 2010

CHARTWELL RX

0.125MG

A090241 001 Oct 08, 2010

0.25MG

A090241 002 Oct 08, 2010

0.5MG

A090241 003 Oct 08, 2010

0.75MG

A090241 004 Oct 08, 2010

1MG

A090241 005 Oct 08, 2010

1.5MG

A090241 006 Oct 08, 2010

HERITAGE PHARMA AVET

0.125MG

A077724 001 Feb 19, 2008

0.125MG

A078551 001 Oct 08, 2010

0.125MG

A091254 001 Nov 30, 2010

0.25MG

A077724 002 Feb 19, 2008

0.25MG

A078551 002 Oct 08, 2010

0.25MG

A091254 002 Nov 30, 2010

0.5MG

A077724 003 Feb 19, 2008

0.5MG

A078551 003 Oct 08, 2010

0.5MG

A091254 003 Nov 30, 2010

0.75MG

A091254 004 Nov 30, 2010

1MG

A077724 004 Feb 19, 2008

1MG

A078551 004 Oct 08, 2010

1MG

A091254 005 Nov 30, 2010

1.5MG

A077724 005 Feb 19, 2008

1.5MG

A078551 005 Oct 08, 2010

1.5MG

A091254 006 Nov 30, 2010

MACLEODS PHARMS LTD

0.125MG

A202164 001 Sep 20, 2012

0.25MG

A202164 002 Sep 20, 2012

0.5MG

A202164 003 Sep 20, 2012

1MG

A202164 004 Sep 20, 2012

1.5MG

A202164 005 Sep 20, 2012

MYLAN

0.125MG

A077854 001 Oct 08, 2010

0.25MG

A077854 002 Oct 08, 2010

0.5MG

A077854 003 Oct 08, 2010

0.75MG

A090764 001 Apr 09, 2010

1MG

A077854 004 Oct 08, 2010

1.5MG

A077854 005 Oct 08, 2010

NOSTRUM LABS INC

0.125MG

A091450 001 Oct 08, 2010

0.25MG

A091450 002 Oct 08, 2010

0.5MG

A091450 003 Oct 08, 2010

1MG

A091450 004 Oct 08, 2010

1.5MG

A091450 005 Oct 08, 2010

SANDOZ

0.125MG

A090190 001 Jul 06, 2010

0.25MG

A090190 002 Jul 06, 2010

0.5MG

A090190 003 Jul 06, 2010

0.75MG

A090190 006 Oct 08, 2010

1MG

A090190 004 Jul 06, 2010

1.5MG

A090190 005 Jul 06, 2010

SUN PHARM INDS INC

0.125MG

A091683 001 Mar 27, 2013

0.25MG

A091683 002 Mar 27, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

0.5MG	A091683 003	Mar 27, 2013
0.75MG	A091683 004	Mar 27, 2013
1MG	A091683 005	Mar 27, 2013
1.5MG	A091683 006	Mar 27, 2013
UNICHEM 0.125MG	A207011 001	Dec 19, 2018
0.25MG	A207011 002	Dec 19, 2018
0.5MG	A207011 003	Dec 19, 2018
0.75MG	A207011 004	Dec 19, 2018
1MG	A207011 005	Dec 19, 2018
1.5MG	A207011 006	Dec 19, 2018

TABLET, EXTENDED RELEASE;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

ANCHEN PHARMS

0.375MG	A202206 001	Feb 06, 2014
0.75MG	A202206 002	Feb 06, 2014
1.5MG	A202206 003	Feb 06, 2014
2.25MG	A202206 004	Feb 06, 2014
3MG	A202206 005	Feb 06, 2014
3.75MG	A202206 006	Feb 06, 2014
4.5MG	A202206 007	Feb 06, 2014

PRAMLINTIDE ACETATE

INJECTABLE;SUBCUTANEOUS

SYMLIN

ASTRAZENECA AB

EQ 3MG BASE/5ML (EQ 600MCG BASE/ML)

N021332 001 Mar 16, 2005

PRASUGREL HYDROCHLORIDE

TABLET;ORAL

PRASUGREL

CHARTWELL RX

EQ 5MG BASE

A205790 001 Oct 16, 2017

EQ 10MG BASE

A205790 002 Oct 16, 2017

DR REDDYS

EQ 5MG BASE

A205926 001 Jul 07, 2020

EQ 10MG BASE

A205926 002 Jul 07, 2020

LUPIN LTD

EQ 5MG BASE

A205930 001 Jan 09, 2023

EQ 10MG BASE

A205930 002 Jan 09, 2023

PRAVASTATIN SODIUM

TABLET;ORAL

PRAVACHOL

+ BRISTOL MYERS SQUIBB 10MG **

N019898 002 Oct 31, 1991

+ 20MG **

N019898 003 Oct 31, 1991

+ 40MG **

N019898 004 Mar 22, 1993

+ 80MG **

N019898 008 Dec 18, 2001

PRAVASTATIN SODIUM

HISUN PHARM HANGZHOU

20MG

A206061 001 Nov 23, 2018

40MG

A206061 002 Nov 23, 2018

80MG

A206061 003 Nov 23, 2018

MYLAN

10MG

A077013 001 Oct 23, 2006

20MG

A077013 002 Oct 23, 2006

40MG

A077013 003 Oct 23, 2006

80MG

A077013 004 Dec 28, 2007

MYLAN PHARMS INC

10MG

A079187 001 May 27, 2010

20MG

A079187 002 May 27, 2010

40MG

A079187 003 May 27, 2010

80MG

A079187 004 May 27, 2010

PLIVA HRVATSKA DOO

10MG

A077730 001 Nov 21, 2006

20MG

A077730 002 Nov 21, 2006

30MG

A077730 003 Nov 21, 2006

40MG

A077730 005 Nov 21, 2006

RANBAXY LABS LTD

10MG

A076445 001 Apr 23, 2007

20MG

A076445 002 Apr 23, 2007

40MG

A076445 003 Apr 23, 2007

80MG

A076445 004 Apr 23, 2007

ZYDUS PHARMS USA

10MG

A077751 001 Apr 30, 2008

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAZEPAM

CAPSULE; ORAL

CENTRAX

PARKE DAVIS	5MG	N018144	001	
	10MG	N018144	002	
	20MG	N018144	003	May 10, 1982

PRAZEPAM

USL PHARMA	5MG	A070427	001	Nov 06, 1987
	10MG	A070428	001	Nov 06, 1987

TABLET; ORAL

CENTRAX

PARKE DAVIS	10MG	N017415	001	
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PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

AM THERAP	EQ 1MG BASE	A072782	001	May 16, 1989
	EQ 2MG BASE	A072783	001	May 16, 1989
	EQ 5MG BASE	A072784	001	May 16, 1989
ANI PHARMS	EQ 1MG BASE	A072577	002	May 16, 1989
	EQ 2MG BASE	A072577	001	May 16, 1989
	EQ 5MG BASE	A072577	003	May 16, 1989
DAVA PHARMS INC	EQ 1MG BASE	A072705	001	May 16, 1989
	EQ 2MG BASE	A072706	001	May 16, 1989
	EQ 5MG BASE	A072707	001	May 16, 1989
PUREPAC PHARM	EQ 1MG BASE	A072991	001	May 16, 1989
	EQ 2MG BASE	A072921	001	May 16, 1989
	EQ 5MG BASE	A072992	001	May 16, 1989
WATSON LABS	EQ 1MG BASE	A072352	001	May 16, 1989
	EQ 2MG BASE	A072333	001	May 16, 1989
	EQ 5MG BASE	A072609	001	May 16, 1989

TABLET, EXTENDED RELEASE; ORAL

MINIPRESS XL

PFIZER	2.5MG	N019775	001	Jan 29, 1992
	5MG	N019775	002	Jan 29, 1992

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

+ VALEANT BERMUDA	0.1% **	N020279	001	Oct 29, 1993
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PREDNICARBATE

FOUGERA PHARMS	0.1%	A077287	001	Sep 19, 2006
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OINTMENT; TOPICAL

DERMATOP

+ VALEANT PHARMS NORTH	0.1% **	N019568	001	Sep 23, 1991
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PREDNISOLONE

CREAM; TOPICAL

METI-DERM

SCHERING	0.5%	N010209	002	
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SYRUP; ORAL

PREDNISOLONE

IVAX SUB TEVA PHARMS	15MG/5ML	A040287	001	May 28, 1999
NESHER PHARMS	5MG/5ML	A040423	001	Oct 22, 2001
	15MG/5ML	A040364	001	Apr 10, 2002
PHARM ASSOC	5MG/5ML	A040570	001	Aug 25, 2005
PHARMOBEDIANT CNSLTG	15MG/5ML	A040313	001	Sep 10, 2003
TEVA PHARMS	15MG/5ML	A040322	001	Jan 19, 2000
WE PHARMS	15MG/5ML	A040192	001	May 28, 1998

PRELONE

MURO	5MG/5ML	A089654	001	Jan 17, 1989
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TABLET; ORAL

CORTALONE

HALSEY	1MG	A080304	003	
	2.5MG	A080304	002	
	5MG	A080304	001	

DELTA-CORTEF

PHARMACIA AND UPJOHN	5MG	N009987	004	
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FERNISOLONE-P

FERNDALE LABS	5MG	A083941	001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE

TABLET; ORAL

PREDNISOLONE

BARR	5MG	A084426	002	
BUNDY	5MG	A083675	001	
CHARTWELL RX	5MG	A084542	001	
ELKINS SINN	5MG	A080625	001	
EVERYLIFE	1MG	A084439	001	
	2.5MG	A084439	002	
	5MG	A084439	003	
FERRANTE	2.5MG	A080562	001	
	5MG	A080562	002	
FOSUN PHARMA	5MG	A080339	001	
HEATHER	5MG	A080326	001	
IMPAX LABS	5MG	A080780	001	
INWOOD LABS	5MG	A080748	001	
IVAX SUB TEVA PHARMS	5MG	A080378	001	
MARSHALL PHARMA	5MG	A080307	001	
PANRAY	1MG	A080351	001	
	5MG	A080351	002	
PHOENIX LABS NY	5MG	A080322	001	
PUREPAC PHARM	5MG	A080325	001	
PVT FORM	5MG	A080211	001	
RISING	5MG	A084773	001	
ROXANE	5MG	A080327	002	
SPERTI	1MG	A080358	001	
	2.5MG	A080358	002	
	5MG	A080358	003	
SUPERPHARM	5MG	A088892	001	Feb 26, 1985
TABLICAPS	5MG	A085170	001	
TEVA	5MG	A080398	001	
UDL	5MG	A087987	001	Jan 18, 1983
VALEANT PHARM INTL	5MG	A080236	001	
VITARINE	5MG	A080534	001	
WATSON LABS	5MG	A085085	002	
	5MG	A085415	001	
	5MG	A085416	001	
WEST WARD	5MG	A080324	001	
WHITEWORTH TOWN PLSN	5MG	A080342	001	
STERANE				
PFIZER	5MG	N009996	001	

PREDNISOLONE ACETATE

INJECTABLE; INJECTION

METICORTELONE

SCHERING	25MG/ML	N010255	002	
PREDNISOLONE ACETATE				
AKORN	25MG/ML	A083032	001	
	50MG/ML	A084492	001	
BEL MAR	25MG/ML	A083738	001	
	50MG/ML	A083738	002	
CENT PHARMS	25MG/ML	A084717	001	
	50MG/ML	A084717	002	
WATSON LABS	25MG/ML	A083398	001	
	25MG/ML	A083654	001	
	40MG/ML	A083767	001	
	50MG/ML	A083764	001	
	50MG/ML	A085781	001	

STERANE

PFIZER 25MG/ML N011446 001

SUSPENSION; ORAL

FLO-PRED

TARO	EQ 5MG BASE/5ML	N022067	001	Jan 17, 2008
	EQ 15MG BASE/5ML	N022067	002	Jan 17, 2008

SUSPENSION/DROPS; OPHTHALMIC

ECONOPRED

HARROW EYE 0.125% N017468 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT;OPHTHALMIC

BLEPHAMIDE S.O.P.

ALLERGAN 0.2%;10% A087748 001 Dec 03, 1986

CETAPRED

ALCON 0.25%;10% A087771 001 Aug 06, 1993

METIMYD

SCHERING 0.5%;10% N010210 002 Sep 09, 1984

PREDSULFAIR

PHARMAFAIR 0.5%;10% A088032 001 Apr 15, 1983

VASOCIDIN

NOVARTIS 0.5%;10% A088791 001 Oct 05, 1984

SUSPENSION;OPHTHALMIC

BLEPHAMIDE

+ ALLERGAN 0.2%;10% N012813 002

ISOPTO CETAPRED

ALCON 0.25%;10% A087547 001

SUSPENSION/DROPS;OPHTHALMIC

METIMYD

SCHERING 0.5%;10% N010210 001

PREDAMIDE

AKORN 0.5%;10% A088059 001 Jul 29, 1983

PREDSULFAIR

PHARMAFAIR 0.5%;10% A088007 001 Apr 19, 1983

PREDSULFAIR II

PHARMAFAIR 0.2%;10% A088837 001 Dec 24, 1985

SULPHRIN

BAUSCH AND LOMB 0.5%;10% A088089 001 Dec 28, 1982

PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

HYDELTRASOL

MERCCK EQ 20MG PHOSPHATE/ML N011583 002

PREDNISOLONE SODIUM PHOSPHATE

WATSON LABS EQ 20MG PHOSPHATE/ML A080517 001

OINTMENT;OPHTHALMIC, OTIC

HYDELTRASOL

MERCCK EQ 0.25% PHOSPHATE N011028 001

SOLUTION;ORAL

ORAPRED

CONCORDIA PHARMS INC EQ 15MG BASE/5ML ** A075117 001 Dec 14, 2000

PREDNISOLONE SODIUM PHOSPHATE

AMNEAL PHARMS EQ 15MG BASE/5ML A078345 001 Mar 10, 2009

BAUSCH EQ 15MG BASE/5ML A075250 001 Jul 12, 2002

NESHER PHARMS EQ 5MG BASE/5ML A076982 001 May 24, 2005

EQ 15MG BASE/5ML A076988 001 May 24, 2005

PHARM ASSOC EQ 5MG BASE/5ML A076123 001 Dec 23, 2002

PHARMOBEDIANT CNSLTG EQ 5MG BASE/5ML A075099 001 Jun 28, 2002

VINTAGE EQ 15MG BASE/5ML A079010 001 May 26, 2009

VINTAGE PHARMS EQ 5MG BASE/5ML A078416 001 Oct 31, 2007

WE PHARMS EQ 5MG BASE/5ML A075181 001 Dec 23, 2002

XTTRIUM LABS INC EQ 15MG BASE/5ML A076895 001 Oct 04, 2004

SOLUTION/DROPS;OPHTHALMIC

INFLAMASE FORTE

+ NOVARTIS EQ 0.9% PHOSPHATE A080751 002

INFLAMASE MILD

+ NOVARTIS EQ 0.11% PHOSPHATE A080751 001

METRETON

SCHERING EQ 0.5% PHOSPHATE A083834 001

PREDAIR

PHARMAFAIR EQ 0.11% PHOSPHATE A088415 001 Feb 29, 1984

PREDAIR FORTE

PHARMAFAIR EQ 0.9% PHOSPHATE A088165 001 Mar 28, 1983

PREDNISOLONE SODIUM PHOSPHATE

AKORN EQ 0.11% PHOSPHATE A083358 001

EQ 0.9% PHOSPHATE A083358 002

ALCON PHARMS LTD EQ 0.11% PHOSPHATE A081043 001 Oct 24, 1991

EQ 0.9% PHOSPHATE A081044 001 Oct 24, 1991

BAUSCH AND LOMB EQ 0.11% PHOSPHATE A040065 001 Jul 29, 1994

SOLA BARNES HIND EQ 0.11% PHOSPHATE A084171 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION/DROPS;OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

EQ 0.9% PHOSPHATE	A084168	001
EQ 0.9% PHOSPHATE	A084169	001
EQ 0.9% PHOSPHATE	A084172	001

TABLET, ORALLY DISINTEGRATING;ORAL

PREDNISOLONE SODIUM PHOSPHATE

RISING	EQ 10MG BASE	A202179	001	Apr 10, 2013
	EQ 15MG BASE	A202179	002	Apr 10, 2013
	EQ 30MG BASE	A202179	003	Apr 10, 2013

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

SANDOZ	EQ 0.23% PHOSPHATE;10%	A073630	001	May 27, 1993
SULSTER				
AKORN	EQ 0.23% PHOSPHATE;10%	A074511	001	Jul 30, 1996
VASOCIDIN				
+ NOVARTIS	EQ 0.23% PHOSPHATE;10% **	N018988	001	Aug 26, 1988

PREDNISOLONE TEBUTATE

INJECTABLE; INJECTION

HYDELTRA-TBA

MERCK	20MG/ML	N010562	001
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PREDNISOLONE TEBUTATE

WATSON LABS	20MG/ML	A083362	001	Feb 17, 1984
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PREDNISON

SOLUTION;ORAL

PREDNISON

XTTRIUM LABS INC	5MG/5ML	A089726	001	Aug 02, 1988
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SYRUP;ORAL

LIQUID PRED

MURO	5MG/5ML	A087611	002	Sep 07, 1982
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TABLET;ORAL

CORTAN

HALSEY	20MG	A087480	001
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DELTA-DOME

BAYER PHARMS	5MG	A080293	001
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DELTASONE

+ PHARMACIA AND UPJOHN	2.5MG **	N009986	005
+	5MG **	N009986	002
+	10MG **	N009986	006
+	20MG **	N009986	007
+	50MG **	N009986	008

FERNISON

FERNDALE LABS	5MG	A083364	001
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METICORTEN

+ SCHERING	1MG **	N009766	002
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+	5MG **	N009766	001
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ORASONE

SOLVAY	1MG	A083009	001
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	5MG	A083009	002
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	10MG	A083009	003
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	20MG	A083009	004
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	50MG	A085999	001
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PARACORT

PARKE DAVIS	5MG	N010962	002
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PREDNICEN-M

SCHWARZ PHARMA	5MG	A084655	001
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PREDNISON

AM THERAP	5MG	A089387	001	Nov 06, 1986
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	10MG	A089388	001	Nov 06, 1986
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	20MG	A089389	001	Nov 06, 1986
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AMNEAL PHARMS NY	5MG	A089597	001	Oct 05, 1987
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	10MG	A089598	001	Oct 05, 1987
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	20MG	A089599	001	Oct 05, 1987
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BUNDY	5MG	A083676	001
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CHARTWELL MOLECULAR	5MG	A080514	001
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	20MG	A084275	001
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISON

TABLET; ORAL

PREDNISON

CHARTWELL RX	5MG	A083059	001	
CONTRACT PHARMACAL	5MG	A080209	001	
DURAMED PHARMS BARR	5MG	A088394	001	Oct 04, 1983
	10MG	A088395	001	Oct 04, 1983
	20MG	A088396	001	Oct 04, 1983
ELKINS SINN	5MG	A080491	001	
	20MG	A085811	001	
EVERYLIFE	1MG	A084440	001	
	2.5MG	A084440	002	
	5MG	A084440	003	
FERRANTE	2.5MG	A080563	001	
	5MG	A080563	002	
HALSEY	5MG	A080300	001	
HEATHER	5MG	A080320	001	
	10MG	A084341	001	
	20MG	A084417	001	
	20MG	A085543	001	
	50MG	A086946	001	
HIKMA PHARMS	1MG	A040890	001	Nov 01, 2010
	2.5MG	A040538	001	Jan 08, 2004
	50MG	A088465	001	Jun 01, 1984
IMPAX LABS	5MG	A080782	001	
INWOOD LABS	1MG	A080328	001	
	2.5MG	A080306	001	
	5MG	A080279	001	
IVAX SUB TEVA PHARMS	5MG	A080283	001	
	10MG	A084133	001	
	20MG	A084134	001	
KV PHARM	5MG	A084236	001	
LEDERLE	5MG	A086968	001	
MARSHALL PHARMA	5MG	A080301	001	
MUTUAL PHARM	5MG	A080701	001	
	10MG	A086595	001	
	20MG	A084634	001	
NYLOS	5MG	A085115	001	
PANRAY	1MG	A080350	001	
	2.5MG	A080350	002	
	5MG	A080350	003	
PHARMAVITE	5MG	A084662	002	
PHOENIX LABS NY	5MG	A080321	001	
	20MG	A083807	001	
PUREPAC PHARM	5MG	A080353	001	
	10MG	A086062	001	
	20MG	A086061	001	
PVT FORM	20MG	A085151	001	
REXALL	5MG	A080232	001	
RISING	5MG	A084774	001	
	10MG	A089983	001	Jan 12, 1989
	20MG	A085813	001	
	50MG	A089984	001	Jan 12, 1989
ROXANE	20MG	N017109	001	
+	25MG	A087833	001	May 04, 1982
SANDOZ	5MG	A080336	002	
SCHERER LABS	5MG	A080371	001	
SPERTI	1MG	A080359	001	
	2.5MG	A080359	002	
	5MG	A080359	003	
SUN PHARM INDUSTRIES	50MG	A086596	001	
SUPERPHARM	5MG	A088865	001	Oct 25, 1984
	10MG	A088866	001	Oct 25, 1984
	20MG	A088867	001	Oct 25, 1984
TEVA	5MG	A080397	001	
UDL	5MG	A087984	001	Jan 18, 1983
	10MG	A087985	001	Jan 18, 1983
	20MG	A087986	001	Jan 18, 1983
UPSHER SMITH	5MG	A087471	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISONTABLET; ORAL
PREDNISON

	20MG	A087470	001	
VALEANT PHARM INTL	5MG	A080237	001	
VANGARD	5MG	A087682	001	Jan 15, 1982
	20MG	A087701	001	Jan 15, 1982
VITARINE	5MG	A080334	001	
	5MG	A080506	001	
WATSON LABS	5MG	A085084	002	
	10MG	A087773	001	Jul 13, 1982
	20MG	A086813	001	
	50MG	A086867	001	
	50MG	A087772	001	Jul 13, 1982
WHITEWORTH TOWN PLSN	2.5MG	A084913	001	
	5MG	A080343	001	
	10MG	A089028	001	Jul 24, 1986
	20MG	A084913	002	

SERVISONE

LEDERLE

5MG A080223 001

PREGABALIN

CAPSULE; ORAL

PREGABALIN

ACI	25MG	A215755	001	Apr 26, 2023
	50MG	A215755	002	Apr 26, 2023
	75MG	A215755	003	Apr 26, 2023
	100MG	A215755	004	Apr 26, 2023
	150MG	A215755	005	Apr 26, 2023
	200MG	A215755	006	Apr 26, 2023
	225MG	A215755	007	Apr 26, 2023
	300MG	A215755	008	Apr 26, 2023
LUPIN LTD	25MG	A091040	001	May 03, 2022
	50MG	A091040	002	May 03, 2022
	75MG	A091040	003	May 03, 2022
	100MG	A091040	004	May 03, 2022
	150MG	A091040	005	May 03, 2022
	200MG	A091040	006	May 03, 2022
	225MG	A091040	007	May 03, 2022
	300MG	A091040	008	May 03, 2022
MYLAN	25MG	A091228	001	Sep 20, 2019
	50MG	A091228	002	Sep 20, 2019
	75MG	A091228	003	Sep 20, 2019
	100MG	A091228	004	Sep 20, 2019
	150MG	A091228	005	Sep 20, 2019
	200MG	A091228	006	Sep 20, 2019
	225MG	A091228	007	Sep 20, 2019
	300MG	A091228	008	Sep 20, 2019
ZYDUS PHARMS	25MG	A206752	001	Dec 09, 2022
	50MG	A206752	002	Dec 09, 2022
	75MG	A206752	003	Dec 09, 2022
	100MG	A206752	004	Dec 09, 2022
	150MG	A206752	005	Dec 09, 2022
	200MG	A206752	006	Dec 09, 2022
	225MG	A206752	007	Dec 09, 2022
	300MG	A206752	008	Dec 09, 2022

TABLET, EXTENDED RELEASE; ORAL

PREGABALIN

ALVOGEN	82.5MG	A211687	001	Jul 06, 2021
	165MG	A211687	002	Jul 06, 2021
	330MG	A211687	003	Jul 06, 2021
MYLAN	82.5MG	A211948	001	Apr 13, 2021
	165MG	A211967	001	Nov 04, 2021
	330MG	A211431	001	Jul 02, 2021
SCIEGEN PHARMS INC	82.5MG	A215675	001	Sep 14, 2022
	165MG	A215675	002	Sep 14, 2022
	330MG	A215675	003	Sep 14, 2022
ZYDUS PHARMS	82.5MG	A215577	001	Aug 26, 2022
	165MG	A215577	002	Aug 26, 2022
	330MG	A215577	003	Aug 26, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST

+	ASTRAZENECA	1% **	N014763	004
+		2% **	N014763	005
+		3% **	N014763	003

CITANEST PLAIN

+	ASTRAZENECA	4% **	N014763	007
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CITANEST PLAIN DENTAL

+	DENTSPLY PHARM	4%	N021382	001
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PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE PHOSPHATE

ALVOGEN	EQ 15MG BASE	A203924	001	Feb 03, 2014
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PRIMIDONE

SUSPENSION; ORAL

MYSOLINE

+	NURO PHARMA	250MG/5ML	N010401	001
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TABLET; ORAL

PRIMIDONE

DR REDDYS LABS LTD	50MG	A040862	001	Oct 03, 2008
	250MG	A040862	002	Oct 03, 2008
HIKMA INTL PHARMS	50MG	A040667	001	Jul 27, 2006
	250MG	A040667	002	Jul 27, 2006
IMPAX LABS	50MG	A040717	001	Feb 12, 2008
	250MG	A040717	002	Feb 12, 2008
WATSON LABS	250MG	A085052	001	

PROBENECID

TABLET; ORAL

BENEMID

+	MERCK	500MG **	N007898	004
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PROBENECID

AUROBINDO PHARMA USA	500MG	A084211	002	
IVAX SUB TEVA PHARMS	500MG	A083740	001	May 09, 1984
LEDERLE	500MG	A086917	001	
WATSON LABS	500MG	A086150	002	Apr 23, 1982

PROBUCOL

TABLET; ORAL

LORELCO

SANOFI AVENTIS US	250MG	N017535	001	
	500MG	N017535	002	Jul 06, 1988

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

ANI PHARMS	250MG	A089219	001	Jul 01, 1986
	375MG	A089219	002	Jul 01, 1986
	500MG	A089219	003	Jul 01, 1986
ASCOT	250MG	A087542	001	Jan 08, 1982
	375MG	A087697	001	Mar 01, 1983
	500MG	A087543	001	Jan 08, 1982
IVAX SUB TEVA PHARMS	250MG	A084604	001	
	375MG	A084595	001	
	500MG	A084606	001	
LANNETT	250MG	A083693	001	
	500MG	A084696	001	
LEDERLE	250MG	A086942	001	
	375MG	A086952	001	
	500MG	A086943	001	
ROXANE	250MG	A088989	001	Apr 26, 1985
	500MG	A088990	001	Apr 26, 1985
VANGARD	250MG	A087643	001	Jun 01, 1982
	500MG	A087875	001	Jun 01, 1982
WATSON LABS	250MG	A083287	001	
	250MG	A083795	001	
	250MG	A085167	001	
	375MG	A084403	001	
	375MG	A087020	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

500MG	A084280	001
500MG	A084357	001
500MG	A087021	001

PROCAN

PARKE DAVIS	250MG	A085804	001
	375MG	A087502	001
	500MG	A085079	001

PROCAPAN

PANRAY	250MG	A083553	002
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PRONESTYL

+	APOTHECON	250MG	**	N007335	001
+		375MG	**	N007335	004
+		500MG	**	N007335	003

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

ABRAXIS PHARM	100MG/ML	A089415	001	Nov 17, 1986
	500MG/ML	A089416	001	Nov 17, 1986
HOSPIRA	500MG/ML	A089537	001	Aug 25, 1987
INTL MEDICATION	500MG/ML	A088637	001	Jul 31, 1984
PHARMAFAIR	100MG/ML	A088824	001	Nov 20, 1985
	500MG/ML	A088830	001	Nov 20, 1985
SMITH AND NEPHEW	100MG/ML	A088530	001	Mar 04, 1985
	500MG/ML	A088531	001	Mar 04, 1985
SOLOPAK	500MG/ML	A088532	001	Mar 04, 1985
WARNER CHILCOTT	100MG/ML	A089528	001	May 03, 1988
	500MG/ML	A089529	001	May 03, 1988
WATSON LABS	100MG/ML	A087079	001	
	500MG/ML	A087080	001	
WEST-WARD PHARMS INT	100MG/ML	A089029	001	Apr 17, 1986
	500MG/ML	A089030	001	Apr 17, 1986

PRONESTYL

+	APOTHECON	100MG/ML	**	N007335	002
+		500MG/ML	**	N007335	005

TABLET; ORAL

PRONESTYL

APOTHECON	250MG	N017371	001
	375MG	N017371	002
	500MG	N017371	003

TABLET, EXTENDED RELEASE; ORAL

PROCAINAMIDE HYDROCHLORIDE

ANI PHARMS	250MG	A088958	001	Dec 02, 1985
	250MG	A089369	001	Aug 14, 1987
	500MG	A088959	001	Dec 02, 1985
	500MG	A088974	001	Jul 22, 1985
	500MG	A089369	002	Jan 09, 1987
	750MG	A089369	003	Aug 14, 1987
	750MG	A089438	001	Mar 23, 1987
	1GM	A040111	001	Dec 13, 1996
INWOOD LABS	500MG	A089840	001	Mar 06, 1989
SANDOZ	500MG	A089284	001	Jun 23, 1986
WATSON LABS	250MG	A088533	001	Dec 03, 1984
	250MG	A089026	001	Oct 22, 1985
	500MG	A088534	001	Dec 03, 1984
	500MG	A089027	001	Oct 22, 1985
	750MG	A088535	001	Nov 03, 1984
	750MG	A089042	001	Oct 22, 1985
	1GM	A089520	001	Jan 15, 1987

PROCAN SR

+	PARKE DAVIS	250MG	A086468	001	
+	PARKEDALE	500MG	A086065	001	
+		750MG	A087510	001	Apr 01, 1982
		1GM	A088489	001	Jan 16, 1985

PROCANBID

KING PHARMS	500MG	N020545	001	Jan 31, 1996
	1GM	N020545	002	Jan 31, 1996

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINAMIDE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PRONESTYL-SR

APOTHECON

500MG

A087361 001

PROCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

NOVOCAIN

HOSPIRA

1%

A085362 003

2%

A085362 004

10%

A086797 001

PROCAINE HYDROCHLORIDE

ABRAXIS PHARM

1%

A080384 002

1%

A080421 001

2%

A080384 003

2%

A080421 002

BEL MAR

1%

A080711 001

2%

A080756 001

ELKINS SINN

1%

A083315 001

2%

A083315 002

GD SEARLE LLC

1%

A086202 001

2%

A086202 002

HOSPIRA

1%

A080416 001

2%

A080416 002

MILES

1%

A080415 001

2%

A080415 002

WATSON LABS

1%

A080658 001

1%

A083535 001

2%

A080658 002

2%

A083535 002

PROCAINE HYDROCHLORIDE; TETRACYCLINE HYDROCHLORIDE

INJECTABLE;INJECTION

ACHROMYCIN

LEDERLE

40MG/VIAL;100MG/VIAL

N050276 001

40MG/VIAL;250MG/VIAL

N050276 003

TETRACYN

PFIZER

40MG/VIAL;100MG/VIAL

A060285 002

40MG/VIAL;250MG/VIAL

A060285 003

PROCAINE MERETHOXYLLINE; THEOPHYLLINE

INJECTABLE;INJECTION

DICURIN PROCAINE

LILLY

100MG/ML;50MG/ML

N008869 001

PROCHLORPERAZINE

SUPPOSITORY;RECTAL

COMPAZINE

GLAXOSMITHKLINE

2.5MG **

N011127 003

5MG **

N011127 001

25MG **

N011127 002

PROCHLORPERAZINE

ABLE

2.5MG

A040407 001

Jul 11, 2001

5MG

A040407 002

Jul 11, 2001

25MG

A040407 003

Jul 11, 2001

PROCHLORPERAZINE EDISYLATE

CONCENTRATE;ORAL

COMPAZINE

+

GLAXOSMITHKLINE

EQ 10MG BASE/ML

N011276 001

PROCHLORPERAZINE

ALPHARMA US PHARMS

EQ 10MG BASE/ML

A087153 001

Jun 08, 1982

PROCHLORPERAZINE EDISYLATE

MORTON GROVE

EQ 10MG BASE/ML

A088598 001

Oct 25, 1984

INJECTABLE;INJECTION

COMPAZINE

+

GLAXOSMITHKLINE

EQ 5MG BASE/ML **

N010742 002

PROCHLORPERAZINE

BAXTER HLTHCARE

EQ 5MG BASE/ML

A087759 001

Oct 01, 1982

PROCHLORPERAZINE EDISYLATE

HIKMA

EQ 5MG BASE/ML

A089523 001

May 03, 1988

HOSPIRA

EQ 5MG BASE/ML

A089703 001

Apr 07, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

MARSAM PHARMS LLC	EQ 5MG BASE/ML	A089675 001	Dec 05, 1988
SMITH AND NEPHEW	EQ 5MG BASE/ML	A089251 001	Dec 04, 1986
TEVA PARENTERAL	EQ 5MG BASE/ML	A040505 001	May 30, 2003
WATSON LABS	EQ 5MG BASE/ML	A089530 001	Jul 08, 1987
	EQ 5MG BASE/ML	A089605 001	Jul 08, 1987
	EQ 5MG BASE/ML	A089606 001	Jul 08, 1987
WYETH AYERST	EQ 5MG BASE/ML	A086348 001	

SYRUP; ORAL

COMPAZINE

GLAXOSMITHKLINE	EQ 5MG BASE/5ML	N011188 001	
PROCHLORPERAZINE EDISYLATE			
ALPHARMA US PHARMS	EQ 5MG BASE/5ML	A087154 001	Sep 01, 1982
MORTON GROVE	EQ 5MG BASE/5ML	A088597 001	Oct 25, 1984

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPAZINE

GLAXOSMITHKLINE	EQ 10MG BASE	N011000 001	
	EQ 10MG BASE	N021019 001	Oct 06, 1999
	EQ 15MG BASE	N011000 002	
	EQ 15MG BASE	N021019 002	Oct 06, 1999
	EQ 30MG BASE	N011000 003	
	EQ 75MG BASE	N011000 004	

TABLET; ORAL

COMPAZINE

+ GLAXOSMITHKLINE	EQ 5MG BASE **	N010571 001	
+	EQ 10MG BASE **	N010571 002	
+	EQ 25MG BASE **	N010571 003	

PROCHLORPERAZINE

WATSON LABS	EQ 5MG BASE	A085580 001	
	EQ 10MG BASE	A085178 001	
	EQ 25MG BASE	A085579 001	

PROCHLORPERAZINE MALEATE

CHARTWELL RX	EQ 5MG BASE	A040101 001	Jul 19, 1996
	EQ 10MG BASE	A040101 002	Jul 19, 1996
	EQ 25MG BASE	A040101 003	Jul 19, 1996
DURAMED PHARMS BARR	EQ 5MG BASE	A040207 001	May 01, 1997
	EQ 5MG BASE	A089484 001	Jan 20, 1987
	EQ 10MG BASE	A040207 002	May 01, 1997
	EQ 10MG BASE	A089485 001	Jan 20, 1987
	EQ 25MG BASE	A089486 001	Jan 20, 1987
IVAX SUB TEVA PHARMS	EQ 5MG BASE	A040162 001	Jan 20, 1998
	EQ 10MG BASE	A040162 002	Jan 20, 1998
MYLAN	EQ 5MG BASE	A040185 002	Oct 28, 1996
	EQ 10MG BASE	A040185 001	Oct 28, 1996
TEVA PHARMS	EQ 5MG BASE	A040120 001	Jul 11, 1996
	EQ 10MG BASE	A040120 002	Jul 11, 1996

PROCYCLIDINE HYDROCHLORIDE

TABLET; ORAL

KEMADRIN

MONARCH PHARMS	2MG	N009818 005	
	5MG	N009818 003	

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

TEVA PHARMS	100MG	A202121 001	Feb 29, 2012
	200MG	A202121 002	Feb 29, 2012

PROMETRIUM

VIRTUS	300MG	N019781 003	Oct 15, 1999
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INJECTABLE; INJECTION

PROGESTERONE

+ ACTAVIS LABS UT INC	50MG/ML **	N017362 002	
AM REGENT	50MG/ML	A090845 001	Jun 22, 2009
LILLY	25MG/ML	N009238 002	
	50MG/ML	N009238 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROGESTERONE

INSERT, EXTENDED RELEASE; INTRAUTERINE

PROGESTASERT

ALZA

38MG

N017553 001

SYSTEM; VAGINAL

MILPROSA

+

FERRING PHARMS INC

1.78GM

N201110 001 Apr 29, 2020

PROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

SPARINE

WYETH AYERST

30MG/ML

N010942 001

100MG/ML

N010942 004

INJECTABLE; INJECTION

PROMAZINE HYDROCHLORIDE

WATSON LABS

25MG/ML

A084510 001

50MG/ML

A084517 001

SPARINE

HIKMA

25MG/ML

N010349 008

50MG/ML

N010349 006

SYRUP; ORAL

SPARINE

WYETH AYERST

10MG/5ML

N010942 003

TABLET; ORAL

SPARINE

WYETH AYERST

10MG

N010348 006

25MG

N010348 001

50MG

N010348 002

100MG

N010348 003

200MG

N010348 004

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PHENERGAN

WYETH AYERST

25MG/ML

N008857 002

50MG/ML

N008857 003

PROMETHAZINE HYDROCHLORIDE

ABBOTT

25MG/ML

A084223 001

50MG/ML

A084222 001

AKORN

25MG/ML

A083955 002

50MG/ML

A083955 001

AM REGENT

25MG/ML

A040515 001 Mar 19, 2003

BEDFORD LABS

25MG/ML

A040524 001 Mar 17, 2004

50MG/ML

A040524 002 Mar 17, 2004

HOSPIRA

25MG/ML

A040372 001 Jun 08, 2000

50MG/ML

A040372 002 Jun 08, 2000

50MG/ML

A083838 002

MARSAM PHARMS LLC

25MG/ML

A089463 001 May 02, 1988

50MG/ML

A089477 001 May 02, 1988

MYLAN INSTITUTIONAL

25MG/ML

A040471 001 Nov 21, 2002

SANDOZ

25MG/ML

A040593 001 Nov 08, 2006

50MG/ML

A040593 002 Nov 08, 2006

TEVA PHARMS USA

25MG/ML **

A040454 001 Aug 22, 2002

50MG/ML **

A040454 002 Aug 22, 2002

WATSON LABS

25MG/ML

A083532 001

25MG/ML

A084591 001

50MG/ML

A080629 002

50MG/ML

A083532 002

WOCKHARDT

25MG/ML

A040785 001 Sep 26, 2008

50MG/ML

A040785 002 Sep 26, 2008

ZIPAN-25

ALTANA

25MG/ML

A083997 001

ZIPAN-50

ALTANA

50MG/ML

A083997 002

SUPPOSITORY; RECTAL

PHENERGAN

+

MYLAN

12.5MG **

N010926 002

+

25MG **

N010926 001

+

50MG **

N011689 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMETHAZINE HYDROCHLORIDE

SUPPOSITORY;RECTAL

PROMETHACON

POLYMEDICA	25MG	A084901	001	
	50MG	A084902	001	

PROMETHAZINE HYDROCHLORIDE

ABLE	12.5MG	A040504	001	Apr 11, 2003
	25MG	A040504	002	Apr 11, 2003
	50MG	A040449	001	Feb 27, 2003
WATSON LABS INC	12.5MG	A040479	001	Jun 24, 2003
	25MG	A040479	002	Jun 24, 2003

SYRUP;ORAL

MYMETHAZINE FORTIS

USL PHARMA	25MG/5ML	A087996	001	Jan 18, 1983
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PROMETH FORTIS

ALPHARMA US PHARMS	25MG/5ML	A084772	001	
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PROMETH PLAIN

ACTAVIS MID ATLANTIC	6.25MG/5ML	A085953	001	
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PROMETHAZINE

CENCI	6.25MG/5ML	A089013	001	Sep 20, 1985
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PROMETHAZINE HYDROCHLORIDE

AMNEAL PHARMS	6.25MG/5ML	A040882	001	Dec 30, 2009
KV PHARM	6.25MG/5ML	A085388	001	
	25MG/5ML	A085385	001	
PHARM ASSOC	6.25MG/5ML	A040643	001	Apr 26, 2006
	6.25MG/5ML	A087518	001	
WHITEWORTH TOWN PLSN	6.25MG/5ML	A086395	001	

PROMETHAZINE HYDROCHLORIDE PLAIN

+ ANI PHARMS	6.25MG/5ML **	N008381	004	Apr 18, 1984
+	25MG/5ML **	N008381	003	

TABLET;ORAL

PHENERGAN

+ DELCOR ASSET CORP	12.5MG **	N007935	002	
+	25MG **	N007935	003	
+	50MG **	N007935	004	

PROMETHAZINE HYDROCHLORIDE

ABBOTT	12.5MG	A084160	001	
	25MG	A084166	001	
	50MG	A084539	001	
ABLE	12.5MG	A040558	001	Jul 01, 2004
	25MG	A040558	002	Jul 01, 2004
	50MG	A040558	003	Jul 01, 2004
AUROBINDO PHARMA USA	12.5MG	A091054	001	Aug 30, 2011
	25MG	A091054	002	Aug 30, 2011
	50MG	A091054	003	Aug 30, 2011
CHARTWELL MOLECULAR	12.5MG	A080949	001	
	25MG	A080949	002	
	50MG	A080949	003	
IMPAX LABS	12.5MG	A040791	002	Feb 12, 2008
	25MG	A040791	003	Feb 12, 2008
	25MG	A084214	002	Jul 07, 1982
	50MG	A040791	001	May 20, 2008
INVATECH	12.5MG	A084233	001	
	25MG	A085146	001	
	50MG	A085146	002	
IVAX SUB TEVA PHARMS	12.5MG	A083604	001	
	25MG	A083603	001	
	50MG	A083613	001	
PVT FORM	12.5MG	A083214	001	
	25MG	A083658	001	
SANDOZ	12.5MG	A084176	002	May 22, 2009
SUN PHARM INDS INC	12.5MG	A040863	001	Dec 30, 2008
	25MG	A040863	002	Dec 30, 2008
	50MG	A040863	003	Dec 30, 2008
SUN PHARM INDUSTRIES	12.5MG	A084555	001	
	25MG	A084554	001	
	50MG	A084557	001	
TABLICAPS	12.5MG	A084080	001	
	25MG	A084027	001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMETHAZINE HYDROCHLORIDE

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

TEVA	25MG	A089109 001	Sep 10, 1985
WATSON LABS	12.5MG	A083401 001	
	12.5MG	A083712 001	
	12.5MG	A085986 001	
	25MG	A083204 001	
	25MG	A085684 001	
	50MG	A083403 001	
	50MG	A085664 001	
REMSD			
BRISTOL MYERS SQUIBB	25MG	A083176 002	
	50MG	A083176 001	

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPAFENONE HYDROCHLORIDE

MYLAN	225MG	A203803 001	Apr 29, 2016
	325MG	A203803 002	Apr 29, 2016
	425MG	A203803 003	Apr 29, 2016
RYTHMOL SR			
+ GLAXOSMITHKLINE LLC	225MG	N021416 001	Sep 04, 2003
+	325MG	N021416 002	Sep 04, 2003
+	425MG	N021416 003	Sep 04, 2003
TABLET; ORAL			
PROPAFENONE HYDROCHLORIDE			
NESHER PHARMS	150MG	A076193 001	Feb 07, 2002
	225MG	A076193 002	Feb 07, 2002
	300MG	A076193 003	Feb 07, 2002
RYTHMOL			
+ GLAXOSMITHKLINE LLC	150MG **	N019151 001	Nov 27, 1989
+	225MG **	N019151 003	Nov 20, 1992
+	300MG **	N019151 002	Nov 27, 1989

PROPANTHELINE BROMIDE

INJECTABLE; INJECTION

PRO-BANTHINE

GD SEARLE LLC	30MG/VIAL	N008843 001	
TABLET; ORAL			
PRO-BANTHINE			
+ SHIRE	7.5MG **	N008732 003	
+	15MG **	N008732 002	
PROPANTHELINE BROMIDE			
ASCOT	15MG	A087663 001	Oct 25, 1982
HEATHER	15MG	A085780 001	
HIKMA	7.5MG	A080927 001	
	15MG	A080927 002	
IMPAX LABS	15MG	A084541 002	
MYLAN	15MG	A083706 001	
PAR PHARM	15MG	A088377 001	Dec 08, 1983
PVT FORM	15MG	A080977 001	
SANDOZ	15MG	A080928 001	
TABLICAPS	15MG	A084428 001	
WATSON LABS	15MG	A083029 002	
	15MG	A083151 001	

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

KAINAIR

PHARMAFAIR	0.5%	A088087 001	Jun 07, 1983
OPHTHAINE			
+ APOTHECON	0.5% **	N008883 001	
OPHTHETIC			
+ ALLERGAN	0.5% **	N012583 001	
PARACAINE			
OPTOPICS	0.5%	A087681 001	Aug 05, 1982
PROPARACAINE HYDROCHLORIDE			
SOLA BARNES HIND	0.5%	A084144 001	
	0.5%	A084151 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPIOLACTONE

SOLUTION; IRRIGATION

BETAPRONE

FOREST LABS

N/A

N011657 001

PROPIOMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

LARGON

HIKMA

20MG/ML

N012382 002

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

FRESENIUS KABI USA

10MG/ML

N019627 001 Oct 02, 1989

PROPOFOL

TEVA PARENTERAL

10MG/ML

A075392 001 Sep 19, 2000

WATSON LABS INC

10MG/ML

A205307 001 Dec 22, 2015

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON

XANODYNE PHARM

32MG

N010997 001

65MG

N010997 003

DOLENE

HERITAGE PHARMS INC

65MG

A080530 001

KESSO-GESIC

MK LABS

65MG

A083544 001

PROPHENE 65

HALSEY

65MG

A083538 002

PROPOXYPHENE HYDROCHLORIDE

ALRA

65MG

A083184 001

IMPAX LABS

65MG

A083317 001

IVAX SUB TEVA PHARMS

32MG

A083597 001

MUTUAL PHARM

65MG

A083186 001

MYLAN

32MG

A083528 001

65MG

A040569 001 Dec 16, 2004

65MG

A083299 001

NEXGEN PHARMA INC

65MG

A083185 001

PAR PHARM

65MG

A080269 001

PUREPAC PHARM

65MG

A083278 001

PVT FORM

32MG

A083464 001

65MG

A083113 001

ROXANE

32MG

A083089 001

65MG

A083089 002

SANDOZ

32MG

A084014 001

65MG

A083125 002

65MG

A083688 001

65MG

A083870 002

65MG

A086495 001

TEVA

65MG

A088615 001 Oct 22, 1984

VALEANT PHARM INTL

65MG

A080783 001

VINTAGE PHARMS

65MG

A040908 001 Jul 17, 2009

WATSON LABS

65MG

A080908 002

65MG

A085190 001

WEST WARD

65MG

A083501 001

WHITEWORTH TOWN PLSN

65MG

A084551 001

PROPOXYPHENE HYDROCHLORIDE 65

WARNER CHILCOTT

65MG

A083786 001

PROPOXYPHENE NAPSYLATE

SUSPENSION; ORAL

DARVON-N

AAIPHARMA LLC

50MG/5ML

N016861 001

TABLET; ORAL

DARVON-N

XANODYNE PHARM

100MG

N016862 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

PROPRANOLOL HYDROCHLORIDE

INWOOD LABS	60MG	A072499	001	Apr 11, 1989
	80MG	A072500	001	Apr 11, 1989
	120MG	A072501	001	Apr 11, 1989
	160MG	A072502	001	Apr 11, 1989
MYLAN	60MG	A078022	001	Feb 15, 2007
	80MG	A078022	002	Feb 15, 2007
	120MG	A078022	003	Feb 15, 2007
	160MG	A078022	004	Feb 15, 2007
UPSHER SMITH LABS	60MG	A078311	001	Mar 06, 2009
	80MG	A078311	002	Mar 06, 2009
	120MG	A078311	003	Mar 06, 2009
	160MG	A078311	004	Mar 06, 2009

CONCENTRATE;ORAL

PROPRANOLOL HYDROCHLORIDE INTENSOL

ROXANE	80MG/ML	A071388	001	May 15, 1987
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INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

+ BAXTER HLTHCARE CORP	1MG/ML **	N016419	001	
CHARTWELL INJECTABLE	1MG/ML	A075792	001	Aug 29, 2000
FOSUN PHARMA	1MG/ML	A076400	001	Feb 26, 2003
SMITH AND NEPHEW	1MG/ML	A070135	001	Apr 15, 1986
	1MG/ML	A070137	001	Apr 15, 1986
SOLOPAK	1MG/ML	A070136	001	Apr 15, 1986

SOLUTION;ORAL

PROPRANOLOL HYDROCHLORIDE

PAI HOLDINGS PHARM	20MG/5ML	A071984	001	Mar 03, 1989
	40MG/5ML	A071985	001	Mar 03, 1989

SUSPENSION;ORAL

INDERAL

WYETH AYERST	10MG/ML	N019536	001	Dec 12, 1986
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TABLET;ORAL

INDERAL

+ WYETH PHARMS	10MG **	N016418	001	
+	20MG **	N016418	003	
+	40MG **	N016418	002	
+	60MG **	N016418	009	Oct 18, 1982
+	80MG **	N016418	004	
+	90MG **	N016418	010	Oct 18, 1982

PROPRANOLOL HYDROCHLORIDE

ANI PHARMS	60MG	A071791	001	Jul 15, 1987
	90MG	A071977	001	Apr 06, 1988
CHARTWELL RX	10MG	A070663	001	Jun 13, 1986
	20MG	A070664	001	Jun 13, 1986
	40MG	A070665	001	Jun 13, 1986
	60MG	A070666	001	Oct 10, 1986
	80MG	A070667	001	Jun 13, 1986
DAVA PHARMS INC	10MG	A070125	001	Jul 30, 1985
	20MG	A070126	001	Jul 30, 1985
	40MG	A070127	001	Jul 30, 1985
	60MG	A071495	001	Dec 31, 1987
	80MG	A070128	001	Jul 30, 1985
	90MG	A071496	001	Dec 31, 1987
DURAMED PHARMS BARR	10MG	A070306	001	Sep 09, 1985
	20MG	A070307	001	Sep 09, 1985
	40MG	A070308	001	Sep 09, 1985
	60MG	A070309	001	Oct 01, 1986
	80MG	A070310	001	Sep 09, 1985
	90MG	A071327	001	Oct 01, 1986
INTERPHARM	10MG	A071368	001	May 05, 1987
	20MG	A071369	001	May 05, 1987
	40MG	A071370	001	May 05, 1987
	80MG	A071371	001	May 05, 1987
IVAX SUB TEVA PHARMS	10MG	A072063	001	Jul 29, 1988
	20MG	A072066	001	Jul 29, 1988
	40MG	A072067	001	Jul 29, 1988
	60MG	A072068	001	Jul 29, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

	80MG	A072069 001	Jul 29, 1988
LEDERLE	10MG	A072117 001	Jun 23, 1988
	20MG	A072118 001	Jun 23, 1988
	40MG	A072119 001	Jun 23, 1988
	80MG	A072120 001	Jun 23, 1988
MYLAN	60MG	A072275 001	Jun 09, 1989
PUREPAC PHARM	10MG	A070814 001	Nov 03, 1986
	20MG	A070815 001	Nov 03, 1986
	40MG	A070816 001	Nov 03, 1986
	60MG	A070817 001	Nov 03, 1986
	80MG	A070757 001	Nov 03, 1986
ROXANE	10MG	A070516 001	Jul 07, 1986
	20MG	A070517 001	Jul 07, 1986
	40MG	A070518 001	Jul 07, 1986
	60MG	A070519 001	Sep 24, 1986
	80MG	A070520 001	Jul 07, 1986
	90MG	A070521 001	Sep 24, 1986
SANDOZ	10MG	A071658 001	Jul 05, 1988
	20MG	A071687 001	Jul 05, 1988
	40MG	A071688 001	Jul 05, 1988
	60MG	A072197 001	Jul 05, 1988
	80MG	A071689 001	Jul 05, 1988
	90MG	A072198 001	Jul 05, 1988
SCHERING	10MG	A070120 001	Aug 06, 1985
	20MG	A070121 001	Aug 06, 1985
	40MG	A070122 001	Aug 06, 1985
	60MG	A070123 001	Oct 29, 1986
	80MG	A070124 001	Aug 06, 1985
STRIDES PHARMA	90MG	A071288 001	Oct 22, 1986
SUPERPHARM	10MG	A071515 001	Jun 08, 1988
	20MG	A071516 001	Jun 08, 1988
	40MG	A071517 001	Jun 08, 1988
	80MG	A071518 001	Jun 08, 1988
TEVA	10MG	A070232 001	Oct 07, 1987
	20MG	A070233 001	Jun 23, 1986
	40MG	A070234 001	Jun 23, 1986
WARNER CHILCOTT	10MG	A070438 001	Sep 15, 1986
	20MG	A070439 001	Sep 15, 1986
	40MG	A070440 001	Sep 15, 1986
	60MG	A070441 001	Sep 24, 1986
	80MG	A070442 001	Sep 15, 1986
WATSON LABS	10MG	A070140 001	Jul 30, 1985
	10MG	A070378 001	Mar 19, 1987
	20MG	A070141 001	Jul 30, 1985
	20MG	A070379 001	Mar 19, 1987
	40MG	A070142 001	Jul 30, 1985
	40MG	A070380 001	Mar 19, 1987
	60MG	A070143 001	Jan 15, 1987
	60MG	A070178 002	Apr 23, 2018
	60MG	A070381 001	Mar 19, 1987
	60MG	A071098 001	Oct 06, 1986
	80MG	A070144 001	Jul 30, 1985
	80MG	A070382 001	Mar 19, 1987
	80MG	A070551 001	Jul 10, 1986
	90MG	A071183 001	Oct 06, 1986
	90MG	A071792 001	Jul 15, 1987
WATSON LABS TEVA	10MG	A070548 001	Jul 10, 1986
	20MG	A070549 001	Apr 11, 1986
	40MG	A070550 001	Apr 11, 1986

PROPYLIODONE

SUSPENSION; INTRATRACHEAL

DIONOSIL AQUEOUS

GLAXOSMITHKLINE 50% N009309 001

DIONOSIL OILY

GLAXOSMITHKLINE 60% N009309 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

ABBOTT	50MG	A084075	001
ANABOLIC	50MG	A080285	001
ANI PHARMS	50MG	A080215	001
CHARTWELL MOLECULAR	50MG	A080016	001
CHARTWELL RX	50MG	A084543	001
HALSEY	50MG	A080015	001
IMPAX LABS	50MG	A080159	001
LILLY	50MG	N006213	001
SUN PHARM INDUSTRIES	50MG	A083982	001
TABLICAPS	50MG	A080840	001
WATSON LABS	50MG	A080932	001
	50MG	A085201	001

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

HIKMA	10MG/ML	A089474	001	Nov 05, 1986
	10MG/ML	A089475	001	Nov 05, 1986
+ LILLY	10MG/ML **	N006460	002	
PHARMACIA AND UPJOHN	50MG/VIAL	N007413	001	
	250MG/VIAL	N007413	002	Aug 02, 1984

PROTEIN HYDROLYSATE

INJECTABLE; INJECTION

AMINOSOL 5%

ABBVIE	5%	N005932	012	Jan 31, 1985
HYPROTIGEN 5%				
B BRAUN	5%	N006170	003	Jan 10, 1984

PROTIRELIN

INJECTABLE; INJECTION

THYPINONE

ABBOTT	0.5MG/ML	N017638	001	
THYREL TRH				
FERRING	0.5MG/ML	N018087	001	

PROTOKYLLOL HYDROCHLORIDE

TABLET; ORAL

VENTAIRE

SANOFI AVENTIS US	2MG	A083459	001	
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PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

VIVACTIL

CHARTWELL RX	5MG	A073644	001	Aug 24, 1995
	10MG	A073645	001	Aug 24, 1995
TEVA WOMENS	5MG **	N016012	001	
	10MG **	N016012	002	

PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

NOVAFED

SANOFI AVENTIS US	120MG	N017603	001	
SUDAFED 12 HOUR				
+ GLAXOSMITHKLINE	120MG **	N017941	002	

TABLET, EXTENDED RELEASE; ORAL

SUDAFED 12 HOUR

MCNEIL CONS	120MG	A073585	001	Oct 31, 1991
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PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACTIFED

GLAXOSMITHKLINE	120MG; 5MG	N018996	001	Jun 17, 1985
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TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES

KV PHARM	120MG; 5MG	A071798	001	Mar 16, 1989
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SYRUP; ORAL

ACTAHIST

CENCI	30MG/5ML; 1.25MG/5ML	A088344	001	Feb 09, 1984
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

HISTAFED

CENCI 30MG/5ML;1.25MG/5ML A088283 001 Apr 20, 1984

MYFED

USL PHARMA 30MG/5ML;1.25MG/5ML A088116 001 Mar 04, 1983

TRILITRON

NEWTRON PHARMS 30MG/5ML;1.25MG/5ML A088474 001 Feb 12, 1985

TABLET; ORAL

ALLERFED

PVT FORM 60MG;2.5MG A088860 001 Jan 31, 1985

CORPHED

FOSUN PHARMA 60MG;2.5MG A088602 001 Apr 11, 1985

PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE

SANDOZ 60MG;2.5MG A088193 001 May 17, 1983

TRILITRON

NEWTRON PHARMS 60MG;2.5MG A088515 001 Jan 09, 1985

TRIPHED

TEVA 60MG;2.5MG A088630 001 May 17, 1984

TRIPROLIDINE AND PSEUDOEPHEDRINE

WATSON LABS 60MG;2.5MG A088318 002 Jan 13, 1984

WEST WARD 60MG;2.5MG A088117 001 Apr 19, 1983

TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS 60MG;2.5MG A085273 001 Dec 12, 1984

SUPERPHARM 60MG;2.5MG A088578 001 Feb 21, 1985

TABLET, EXTENDED RELEASE; ORAL

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES

KV PHARM 120MG;5MG A072758 001 Nov 25, 1991

PSEUDOEPHEDRINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

PSEUDO-12

UCB INC EQ 60MG HYDROCHLORIDE/5ML N019401 001 Jun 19, 1987

PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

AFRINOL

+ SCHERING PLOUGH 120MG N018191 001

PYRIDOSTIGMINE BROMIDE

TABLET; ORAL

PYRIDOSTIGMINE BROMIDE

ANI PHARMS 30MG A040512 002 Jul 20, 2005

60MG A040512 001 Oct 08, 2003

IMPAX LABS INC 60MG A040457 001 Dec 26, 2002

SOLVAY 30MG A089572 001 Nov 27, 1990

US ARMY 30MG N020414 001 Feb 05, 2003

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEXA-BETALIN

LILLY 100MG/ML A080854 001

PYRIDOXINE HYDROCHLORIDE

AKORN 100MG/ML A087967 001 Oct 01, 1982

BEL MAR 100MG/ML A080761 001

DELL LABS 50MG/ML A083771 001

100MG/ML A083772 001

DR REDDYS 100MG/ML A080572 001

ELKINS SINN 100MG/ML A080581 001

LUITPOLD 100MG/ML A080669 001

MYLAN INSTITUTIONAL 100MG/ML A204879 001 Jul 14, 2016

WATSON LABS 100MG/ML A083760 001

PYRILAMINE MALEATE

TABLET; ORAL

PYRILAMINE MALEATE

IMPAX LABS 25MG A080808 001

WATSON LABS 25MG A085231 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PYRIMETHAMINE; SULFADOXINE

TABLET; ORAL

FANSIDAR

ROCHE

25MG; 500MG

N018557 001

PYRITHIONE ZINC

LOTION; TOPICAL

HEAD & SHOULDERS CONDITIONER

WARNER CHILCOTT

0.3%

N019412 002 Mar 10, 1986

PYRVINIUM PAMOATE

SUSPENSION; ORAL

POVAN

PARKE DAVIS

EQ 50MG BASE/5ML

N011964 001

TABLET; ORAL

POVAN

PARKE DAVIS

EQ 50MG BASE

N012485 002

QUAZEPAM

TABLET; ORAL

DORAL

GALT PHARMS

7.5MG

N018708 003 Feb 26, 1987

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

ACTAVIS GRP PTC

EQ 25MG BASE

A201762 001 Feb 27, 2013

EQ 50MG BASE

A201762 002 Feb 27, 2013

EQ 100MG BASE

A201762 003 Feb 27, 2013

EQ 150MG BASE

A201762 004 Feb 27, 2013

EQ 200MG BASE

A201762 005 Feb 27, 2013

EQ 300MG BASE

A201762 006 Feb 27, 2013

EQ 400MG BASE

A201762 007 Feb 27, 2013

ALEMBIC

EQ 25MG BASE

A203390 001 Oct 28, 2014

EQ 50MG BASE

A203390 002 Oct 28, 2014

EQ 100MG BASE

A203390 003 Oct 28, 2014

EQ 200MG BASE

A203390 004 Oct 28, 2014

EQ 300MG BASE

A203390 005 Oct 28, 2014

EQ 400MG BASE

A203390 006 Oct 28, 2014

JUBILANT GENERICS

EQ 25MG BASE

A203150 001 Nov 26, 2013

MYLAN PHARMS INC

EQ 25MG BASE

A090323 001 Mar 27, 2012

TORRENT PHARMS LTD

EQ 25MG BASE

A200363 001 Mar 27, 2012

EQ 50MG BASE

A200363 002 Mar 27, 2012

EQ 100MG BASE

A200363 003 Mar 27, 2012

EQ 200MG BASE

A200363 004 Mar 27, 2012

EQ 300MG BASE

A200363 005 Mar 27, 2012

EQ 400MG BASE

A200363 006 Mar 27, 2012

SEROQUEL

+ ASTRAZENECA

EQ 150MG BASE **

N020639 004 Dec 20, 1998

TABLET, EXTENDED RELEASE; ORAL

QUETIAPINE FUMARATE

AMNEAL PHARMS

EQ 400MG BASE

A211405 001 Oct 26, 2018

ANCHEN PHARMS

EQ 150MG BASE

A090757 001 Dec 01, 2017

EQ 200MG BASE

A090757 002 Dec 01, 2017

EQ 300MG BASE

A090757 003 Dec 01, 2017

EQ 400MG BASE

A090757 004 Dec 01, 2017

RISING

EQ 50MG BASE

A202228 001 Feb 02, 2021

EQ 150MG BASE

A202228 002 Feb 02, 2021

EQ 200MG BASE

A202228 003 Feb 02, 2021

EQ 300MG BASE

A202228 004 Feb 02, 2021

EQ 400MG BASE

A202228 005 Feb 02, 2021

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCUPRIL

+ PFIZER PHARMS

EQ 5MG BASE

N019885 001 Nov 19, 1991

+

EQ 10MG BASE

N019885 002 Nov 19, 1991

+

EQ 20MG BASE

N019885 003 Nov 19, 1991

+

EQ 40MG BASE

N019885 004 Nov 19, 1991

QUINAPRIL HYDROCHLORIDE

ACTAVIS ELIZABETH

EQ 5MG BASE

A076459 001 Dec 22, 2004

EQ 10MG BASE

A076459 002 Dec 22, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

QUINAPRIL HYDROCHLORIDE

	EQ 20MG BASE	A076459 003	Dec 22, 2004
	EQ 40MG BASE	A076459 004	Dec 22, 2004
ACTAVIS LABS FL INC	EQ 5MG BASE	A076049 001	Jan 14, 2005
	EQ 10MG BASE	A076049 002	Jan 14, 2005
	EQ 20MG BASE	A076049 003	Jan 14, 2005
	EQ 40MG BASE	A076049 004	Jan 14, 2005
ANI PHARMS	EQ 5MG BASE	A075504 001	Aug 24, 2007
	EQ 10MG BASE	A075504 002	Aug 24, 2007
	EQ 20MG BASE	A075504 003	Aug 24, 2007
	EQ 40MG BASE	A075504 004	Aug 24, 2007
APOTEX INC	EQ 5MG BASE	A076240 001	Jan 26, 2006
	EQ 10MG BASE	A076240 002	Jan 26, 2006
	EQ 20MG BASE	A076240 003	Jan 26, 2006
	EQ 40MG BASE	A076240 004	Jan 26, 2006
CHARTWELL RX	EQ 5MG BASE	A076803 001	Mar 02, 2005
	EQ 10MG BASE	A076803 002	Mar 02, 2005
	EQ 20MG BASE	A076803 003	Mar 02, 2005
	EQ 40MG BASE	A076803 004	Mar 02, 2005
MYLAN	EQ 5MG BASE	A076036 001	Jan 28, 2005
	EQ 5MG BASE	A076694 001	Dec 23, 2004
	EQ 10MG BASE	A076036 002	Jan 28, 2005
	EQ 10MG BASE	A076694 002	Dec 23, 2004
	EQ 20MG BASE	A076036 003	Jan 28, 2005
	EQ 20MG BASE	A076694 003	Dec 23, 2004
	EQ 40MG BASE	A076036 004	Jan 28, 2005
	EQ 40MG BASE	A076694 004	Dec 23, 2004
SUN PHARM INDS LTD	EQ 5MG BASE	A076607 001	Dec 15, 2004
	EQ 5MG BASE	A090800 001	Jun 18, 2009
	EQ 10MG BASE	A076607 002	Dec 15, 2004
	EQ 10MG BASE	A090800 002	Jun 18, 2009
	EQ 20MG BASE	A076607 003	Dec 15, 2004
	EQ 20MG BASE	A090800 003	Jun 18, 2009
	EQ 40MG BASE	A076607 004	Dec 15, 2004
	EQ 40MG BASE	A090800 004	Jun 18, 2009

QUINESTROL

TABLET;ORAL

ESTROVIS

PARKE DAVIS	0.1MG	N016768 002	
	0.2MG	N016768 003	

QUINETHAZONE

TABLET;ORAL

HYDROMOX

LEDERLE	50MG	N013264 001	
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QUINETHAZONE; RESERPINE

TABLET;ORAL

HYDROMOX R

LEDERLE	50MG;0.125MG	N013927 001	
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QUINIDINE GLUCONATE

INJECTABLE; INJECTION

QUINIDINE GLUCONATE

+ LILLY

80MG/ML N007529 002 Feb 10, 1989

TABLET;ORAL

QUINACT

BAYER HLTHCARE	266MG	A085978 001	
	400MG	A086099 001	

TABLET, EXTENDED RELEASE;ORAL

DURAQUIN

WARNER CHILCOTT	330MG	N017917 001	
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QUINAGLUTE

+ BAYER HLTHCARE

324MG ** N016647 001

QUINALAN

CHARTWELL MOLECULAR	324MG	A088081 001	Feb 10, 1986
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QUINATIME

WATSON LABS	324MG	A087448 001	
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE GLUCONATE

ANI PHARMS	324MG	A087810 001	Sep 29, 1982
ASCOT	324MG	A088582 001	Jun 17, 1985
CYCLE	324MG	A088431 001	Jan 06, 1984
HALSEY	324MG	A089476 001	Apr 10, 1987
RISING	324MG	A089894 001	Dec 15, 1988
SUPERPHARM	324MG	A089164 001	Nov 21, 1985
WATSON LABS	324MG	A087785 001	Jan 24, 1983

QUINIDINE POLYGALACTURONATE

TABLET;ORAL

CARDIOQUIN

PHARM RES ASSOC	275MG	N011642 002	
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QUINIDINE SULFATE

CAPSULE;ORAL

CIN-QUIN

SOLVAY	200MG	A085296 001	
	300MG	A085297 001	

QUINIDINE SULFATE

LILLY	200MG	A085103 001	
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TABLET;ORAL

CIN-QUIN

+ SOLVAY	100MG	A085299 001	
+	200MG	A084932 001	
	300MG	A085298 001	

QUINIDINE SULFATE

BARR	200MG	A084177 001	
CHARTWELL MOLECULAR	200MG	A083743 001	
CONTRACT PHARMACAL	200MG	A083808 001	
CYCLE	200MG	A083640 001	
	300MG	A085632 001	
DAVA PHARMS INC	200MG	A087011 001	
ELKINS SINN	200MG	A083622 001	
EVERYLIFE	200MG	A083439 001	
HALSEY	200MG	A083583 001	
HIKMA	200MG	A083862 001	
IMPAX LABS	200MG	A083347 001	
IVAX SUB TEVA PHARMS	200MG	A084549 001	
KING PHARMS	200MG	A085175 001	
KV PHARM	200MG	A085276 001	
LEDERLE	200MG	A086176 001	
LILLY	200MG	A085038 001	
PERRIGO	200MG	A085322 001	
PHARMAVITE	200MG	A084627 001	
PUREPAC PHARM	200MG	A084003 001	
SANDOZ	200MG	A084631 001	
	200MG	A084914 001	
	300MG	A089839 001	Sep 29, 1988
SCHERER LABS	200MG	A085068 001	
SUN PHARM INDUSTRIES	100MG	A081029 001	Apr 14, 1989
	200MG	A081030 001	Apr 14, 1989
	300MG	A081031 001	Apr 14, 1989
SUPERPHARM	200MG	A088973 001	Apr 10, 1985
USL PHARMA	200MG	A087837 001	Apr 14, 1982
VALEANT PHARM INTL	200MG	A083393 001	
VANGARD	200MG	A087909 001	Jul 13, 1982
VINTAGE PHARMS	200MG	A083963 001	
WARNER CHILCOTT	200MG	A083879 001	
WATSON LABS	100MG	A085584 001	
	200MG	A083288 001	
	200MG	A085140 002	
	300MG	A085583 001	
WHITEWORTH TOWN PLSN	200MG	A085444 001	

QUINORA

KEY PHARMS	200MG	A083576 001	
+ SCHERING	300MG	A085222 001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINIDINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDEX

WYETH PHARMS INC

300MG

N012796 002

QUINIDINE SULFATE

COSETTE

300MG

A040045 001 Jun 30, 1994

QUININE SULFATE

CAPSULE;ORAL

QUININE SULFATE

AUROBINDO PHARMA USA

324MG

A202581 001 Dec 14, 2012

RABEPRAZOLE SODIUM

CAPSULE, DELAYED RELEASE;ORAL

ACIPHEX SPRINKLE

+ AYTU

5MG

N204736 001 Mar 26, 2013

+

10MG

N204736 002 Mar 26, 2013

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

+ WOODWARD

10MG **

N020973 001 May 29, 2002

RABEPRAZOLE SODIUM

MYLAN

20MG

A076885 001 Nov 08, 2013

TEVA PHARMS USA

20MG

A076822 001 Nov 08, 2013

RAMIPRIL

CAPSULE;ORAL

RAMIPRIL

ACTAVIS ELIZABETH

1.25MG

A077513 001 Jun 18, 2008

2.5MG

A077513 002 Jun 18, 2008

5MG

A077513 003 Jun 18, 2008

10MG

A077513 004 Jun 18, 2008

CHARTWELL RX

1.25MG

A077514 001 Jun 18, 2008

2.5MG

A077514 002 Jun 18, 2008

5MG

A077514 003 Jun 18, 2008

10MG

A077514 004 Jun 18, 2008

CIPLA

1.25MG

A077004 001 Aug 07, 2008

2.5MG

A077004 002 Aug 07, 2008

5MG

A077004 003 Aug 07, 2008

10MG

A077004 004 Aug 07, 2008

RANBAXY LABS LTD

5MG

A078849 001 Mar 06, 2009

10MG

A078849 002 Mar 06, 2009

TEVA PHARMS

1.25MG

A077470 001 Jun 18, 2008

2.5MG

A077470 002 Jun 18, 2008

5MG

A077470 003 Jun 18, 2008

10MG

A077470 004 Jun 18, 2008

WATSON LABS

5MG

A076549 003 Oct 24, 2005

TABLET;ORAL

ALTACE

+ KING PFIZER

1.25MG **

N022021 001 Feb 27, 2007

+

2.5MG **

N022021 002 Feb 27, 2007

+

5MG **

N022021 003 Feb 27, 2007

+

10MG **

N022021 004 Feb 27, 2007

RAMIPRIL

APOTEX

1.25MG

A091069 001 Dec 02, 2015

2.5MG

A091069 002 Dec 02, 2015

5MG

A091069 003 Dec 02, 2015

10MG

A091069 004 Dec 02, 2015

MYLAN PHARMS INC

1.25MG

A090650 001 Jun 30, 2011

2.5MG

A090650 002 Jun 30, 2011

5MG

A090650 003 Jun 30, 2011

10MG

A090650 004 Jun 30, 2011

ZYDUS PHARMS USA INC

1.25MG

A090697 001 Sep 24, 2009

2.5MG

A090697 002 Sep 24, 2009

5MG

A090697 003 Sep 24, 2009

10MG

A090697 004 Sep 24, 2009

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANITIDINE BISMUTH CITRATE

TABLET; ORAL

TRITEC

GLAXOSMITHKLINE 400MG N020559 001 Aug 08, 1996

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

AJANTA PHARMA LTD EQ 150MG BASE A209859 001 Sep 27, 2018

EQ 300MG BASE A209859 002 Sep 27, 2018

APPCO EQ 150MG BASE A211893 001 Apr 05, 2019

EQ 300MG BASE A211893 002 Apr 05, 2019

AUROBINDO PHARMA EQ 150MG BASE A211058 001 Jul 16, 2018

EQ 300MG BASE A211058 002 Jul 16, 2018

MYLAN EQ 150MG BASE A075564 001 Oct 27, 2000

EQ 300MG BASE A075564 002 Oct 27, 2000

NOVITIUM PHARMA EQ 150MG BASE A210681 001 Nov 23, 2018

EQ 300MG BASE A210681 002 Nov 23, 2018

TEVA EQ 150MG BASE A075557 001 Oct 31, 2003

EQ 300MG BASE A075557 002 Oct 31, 2003

ZANTAC 150

+ GLAXOSMITHKLINE EQ 150MG BASE ** N020095 001 Mar 08, 1994

ZANTAC 300

+ GLAXOSMITHKLINE EQ 300MG BASE ** N020095 002 Mar 08, 1994

GRANULE, EFFERVESCENT; ORAL

ZANTAC 150

GLAXO GRP LTD EQ 150MG BASE/PACKET N020251 002 Mar 31, 1994

INJECTABLE; INJECTION

RANITIDINE HYDROCHLORIDE

BEDFORD EQ 25MG BASE/ML A074764 001 Nov 19, 2004

HIKMA EQ 25MG BASE/ML A074777 001 Mar 02, 2005

EQ 25MG BASE/ML A077458 001 Feb 16, 2006

MYLAN LABS LTD EQ 25MG BASE/ML A079076 001 Jun 09, 2016

ZYDUS PHARMS USA INC EQ 25MG BASE/ML A091534 001 Feb 22, 2013

ZANTAC

+ PAI HOLDINGS PHARM EQ 25MG BASE/ML N019090 001 Oct 19, 1984

ZANTAC IN PLASTIC CONTAINER

PAI HOLDINGS PHARM EQ 1MG BASE/ML N019593 002 Sep 27, 1991

EQ 50MG BASE/100ML N019593 001 Dec 17, 1986

SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC EQ 15MG BASE/ML A076124 001 Feb 21, 2007

AMNEAL PHARMS EQ 15MG BASE/ML A078312 001 Sep 02, 2008

APOTEX INC EQ 15MG BASE/ML A077602 001 Sep 17, 2007

AUROBINDO PHARMA EQ 15MG BASE/ML A090623 001 Jul 28, 2010

EPIC PHARMA LLC EQ 15MG BASE/ML A091078 001 Mar 22, 2011

LANNETT CO INC EQ 15MG BASE/ML A078890 001 Jul 01, 2010

NOSTRUM LABS INC EQ 15MG BASE/ML A078684 001 Aug 27, 2009

EQ 15MG BASE/ML A091091 001 Sep 20, 2011

RANBAXY EQ 15MG BASE/ML A078448 001 Dec 13, 2007

TARO EQ 15MG BASE/ML A077476 001 Jun 13, 2011

TOLMAR EQ 15MG BASE/ML A090054 001 Nov 15, 2010

TORRENT EQ 15MG BASE/ML A090102 001 May 26, 2009

WOCKHARDT EQ 15MG BASE/ML A079211 001 May 26, 2009

EQ 15MG BASE/ML A079212 001 Feb 23, 2009

ZANTAC

+ GLAXO GRP LTD EQ 15MG BASE/ML N019675 001 Dec 30, 1988

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

AMNEAL PHARMS NY EQ 150MG BASE A077824 001 Oct 13, 2006

EQ 300MG BASE A077824 002 Oct 13, 2006

ANI PHARMS EQ 75MG BASE A075212 001 Jan 14, 2000

EQ 75MG BASE A075296 001 Jan 14, 2000

EQ 150MG BASE A074488 001 Jul 31, 1997

EQ 150MG BASE A077426 001 Dec 19, 2005

EQ 300MG BASE A074488 002 Jul 31, 1997

EQ 300MG BASE A077426 002 Dec 19, 2005

APOTEX EQ 75MG BASE A075167 001 May 04, 2000

APOTEX INC EQ 150MG BASE A200172 001 May 31, 2012

AUROBINDO PHARMA EQ 75MG BASE A207579 001 Nov 13, 2017

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANITIDINE HYDROCHLORIDE

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

	EQ 150MG BASE	A207578 001 Nov 13, 2017
BOEHRINGER INGELHEIM	EQ 150MG BASE	A074662 001 Aug 29, 1997
	EQ 300MG BASE	A074662 002 Aug 29, 1997
CONTRACT PHARMACAL	EQ 75MG BASE	A075094 001 Jun 21, 1999
GRANULES	EQ 150MG BASE	A210243 001 Aug 20, 2018
	EQ 150MG BASE	A210243 002 Aug 20, 2018
HERITAGE PHARMA AVET	EQ 150MG BASE	A075165 001 Sep 30, 1998
	EQ 300MG BASE	A075165 002 Sep 30, 1998
MYLAN	EQ 75MG BASE	A075497 001 Jan 14, 2000
	EQ 150MG BASE	A074023 001 Aug 22, 1997
	EQ 150MG BASE	A074552 001 Jul 30, 1998
	EQ 300MG BASE	A074023 002 Aug 22, 1997
	EQ 300MG BASE	A074552 002 Jul 30, 1998
NOSTRUM LABS INC	EQ 150MG BASE	A203694 001 Nov 30, 2017
	EQ 300MG BASE	A203694 002 Nov 30, 2017
PERRIGO	EQ 75MG BASE	A076195 001 Aug 30, 2002
PERRIGO R AND D	EQ 150MG BASE	A091429 001 May 11, 2011
	EQ 150MG BASE	A091429 002 May 11, 2011
RANBAXY	EQ 75MG BASE	A075254 001 Jan 14, 2000
	EQ 150MG BASE	A075000 001 Jan 30, 1998
	EQ 300MG BASE	A075000 002 Jan 30, 1998
SANDOZ	EQ 75MG BASE	A075519 001 Sep 26, 2002
STRIDES PHARMA	EQ 75MG BASE	A201745 001 Feb 29, 2012
	EQ 75MG BASE	A209160 001 Mar 05, 2018
	EQ 150MG BASE	A200536 001 Jun 28, 2011
	EQ 150MG BASE	A205512 001 Aug 22, 2016
	EQ 150MG BASE	A209161 001 Feb 22, 2018
	EQ 150MG BASE	A210010 001 Aug 01, 2018
	EQ 300MG BASE	A205512 002 Aug 22, 2016
	EQ 300MG BASE	A210010 002 Aug 01, 2018
SUN PHARM INDS LTD	EQ 75MG BASE	A075132 001 Jan 14, 2000
	EQ 150MG BASE	A075439 001 Apr 19, 2000
	EQ 300MG BASE	A075439 002 Apr 19, 2000
THINQ PHARM-CRO PVT	EQ 150MG BASE	A210228 001 Aug 30, 2019
WATSON LABS	EQ 150MG BASE	A074864 001 Oct 20, 1997
	EQ 300MG BASE	A074864 002 Oct 20, 1997
WOCKHARDT	EQ 75MG BASE	A076760 001 Feb 24, 2006
	EQ 75MG BASE	A078884 001 Jul 31, 2008
	EQ 150MG BASE	A078653 001 Nov 26, 2007
	EQ 150MG BASE	A078701 001 Nov 12, 2009
	EQ 300MG BASE	A078701 002 Dec 11, 2009
WOCKHARDT LTD	EQ 150MG BASE	A075208 001 Dec 17, 1998
	EQ 300MG BASE	A075208 002 Dec 17, 1998
ZANTAC 150		
+ CHATTEM SANOFI	EQ 150MG BASE	N021698 001 Aug 31, 2004
+	EQ 150MG BASE	N021698 002 Mar 13, 2007
+ GLAXO GRP LTD	EQ 150MG BASE **	N018703 001 Jun 09, 1983
ZANTAC 300		
+ GLAXO GRP LTD	EQ 300MG BASE **	N018703 002 Dec 09, 1985
ZANTAC 75		
+ CHATTEM SANOFI	EQ 75MG BASE	N020520 001 Dec 19, 1995
TABLET, EFFERVESCENT;ORAL		
ZANTAC 150		
GLAXO GRP LTD	EQ 150MG BASE	N020251 001 Mar 31, 1994
ZANTAC 25		
GLAXO GRP LTD	EQ 25MG BASE	N020251 003 Apr 01, 2004
ZANTAC 75		
+ CHATTEM SANOFI	EQ 75MG BASE **	N020745 001 Feb 26, 1998

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANEXA

+ MENARINI INTL	500MG	N021526 002 Jan 27, 2006
+	1GM	N021526 001 Feb 12, 2007

RANOLAZINE

ACCORD HLTHCARE	500MG	A212930 001 May 18, 2021
	1GM	A212930 002 May 18, 2021

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANOLAZINE

AMNEAL	500MG	A207690 001	Mar 11, 2021
	1GM	A207690 002	Mar 11, 2021
ANI PHARMS	500MG	A210482 001	Oct 29, 2019
	1GM	A210482 002	Oct 29, 2019
CIPLA	500MG	A211291 001	May 28, 2019
	1GM	A211291 002	May 28, 2019

RAPACURONIUM BROMIDE

INJECTABLE; INJECTION

RAPLON

ORGANON USA INC	100MG/VIAL	N020984 001	Aug 18, 1999
	200MG/VIAL	N020984 002	Aug 18, 1999

RASAGILINE MESYLATE

TABLET; ORAL

RASAGILINE MESYLATE

APOTEX INC	EQ 0.5MG BASE	A201950 001	Sep 12, 2013
	EQ 1MG BASE	A201950 002	Sep 12, 2013
CARNEGIE	EQ 0.5MG BASE	A201942 001	Nov 18, 2021
	EQ 1MG BASE	A201942 002	Nov 18, 2021
WATSON LABS INC	EQ 0.5MG BASE	A201823 001	Jul 01, 2013
	EQ 1MG BASE	A201823 002	Jul 01, 2013

RAUWOLFIA SERPENTINA ROOT

TABLET; ORAL

HIWOLFIA

BOWMAN PHARMS	50MG	N009276 003	
	50MG	N009276 005	
	100MG	N009276 004	

HYSERPIN

PHYS PRODS VA	50MG	N010581 001	
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KONGLUCOID

PANRAY	50MG	N009278 001	
	100MG	N009278 002	

RAUDIXIN

APOTHECON	50MG	N008842 001	
	100MG	N008842 002	

RAUSERPIN

FERNDALE LABS	50MG	N009926 002	
	100MG	N009926 004	

RAUVAL

PAL PAK	50MG	N009108 002	
	100MG	N009108 004	

RAUWOLFIA SERPENTINA

BUNDY	50MG	N009477 001	
	100MG	N009477 002	

HALSEY	50MG	A080498 001	
	100MG	A080498 002	

IMPAX LABS	50MG	N009273 001	
	100MG	N009273 002	

IVAX SUB TEVA PHARMS	50MG	N011521 001	
	100MG	N011521 002	

PUREPAC PHARM	50MG	A080842 001	
	100MG	A080842 002	

PVT FORM	50MG	A080583 001	
	100MG	A080583 002	

SOLVAY	50MG	A080500 001	
	100MG	A080500 002	

TABLICAPS	50MG	A083867 001	
	100MG	A083444 001	

VALEANT PHARM INTL	50MG	N009668 001	
	100MG	N009668 002	

WATSON LABS	50MG	A080907 001	
	100MG	A080914 001	

WOLFINA

FOREST PHARMS	50MG	N009255 008	
	100MG	N009255 006	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

REPAGLINIDE

TABLET; ORAL

PRANDIN

+	GEMINI LABS LLC	0.5MG **	N020741 001	Dec 22, 1997
+		1MG **	N020741 002	Dec 22, 1997
+		2MG **	N020741 003	Dec 22, 1997

REPAGLINIDE

	ACTAVIS TOTOWA	0.5MG	A090008 001	Jan 22, 2014
		1MG	A090008 002	Jan 22, 2014
		2MG	A090008 003	Jan 22, 2014
	KENTON	0.5MG	A091517 001	Apr 24, 2015
		1MG	A091517 002	Apr 24, 2015
		2MG	A091517 003	Apr 24, 2015
	MYLAN	0.5MG	A090252 001	Aug 23, 2013
		1MG	A090252 002	Jan 22, 2014
		2MG	A090252 003	Jan 22, 2014

RESCINNAMINE

CAPSULE; ORAL

CINNASIL

	PANRAY	0.5MG	A084736 001	
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TABLET; ORAL

MODERIL

	PFIZER	0.25MG	N010686 003	
		0.5MG	N010686 006	

RESERPINE

ELIXIR; ORAL

SERPASIL

	NOVARTIS	0.2MG/4ML	N009115 005	
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INJECTABLE; INJECTION

SANDRIL

	LILLY	2.5MG/ML	N010012 001	
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SERPASIL

	NOVARTIS	2.5MG/ML	N009434 002	
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TABLET; ORAL

HISERPIA

	BOWMAN PHARMS	0.1MG	N009631 002	
		0.25MG	N009631 004	

RAU-SED

	BRISTOL MYERS SQUIBB	0.1MG	N009357 001	
		0.25MG	N009357 004	
		0.5MG	N009357 006	
		1MG	N009357 008	

RESERPINE

	BARR	0.25MG	A080721 002	
	BELL PHARMA	0.1MG	A083058 001	
		0.25MG	A083058 002	
	BUNDY	0.1MG	N009663 001	
		0.25MG	N009663 003	
	CYCLE	0.1MG	N009859 001	
		0.25MG	N009859 002	
	ELKINS SINN	0.1MG	A083145 001	
		0.25MG	A083145 002	
	EVERYLIFE	0.1MG	N010441 001	
		0.25MG	N010441 002	
		0.5MG	N010441 003	
		1MG	N010441 004	
	HALSEY	0.1MG	A080457 002	
		0.25MG	A080457 001	
		1MG	A080457 003	
	HIKMA INTL PHARMS	0.1MG	A080975 001	
		0.25MG	A080975 002	
		1MG	A080975 003	
	IMPAX LABS	0.1MG	N009627 001	
		0.25MG	N009627 002	
	IVAX SUB TEVA PHARMS	0.1MG	N011185 001	
		0.25MG	N011185 002	
	MARSHALL PHARMA	0.1MG	A080492 001	
		0.25MG	A080492 002	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RESERPINE

TABLET; ORAL

RESERPINE

MK LABS	0.1MG	A080525 002	
	0.25MG	A080525 001	
MYLAN	1MG	A084974 001	
PHARMAVITE	0.25MG	A084663 001	
PUREPAC PHARM	0.1MG	A080753 002	
	0.25MG	A080753 001	
PVT FORM	0.1MG	A086117 001	
	0.25MG	A080582 001	
	0.25MG	A085775 001	
	1MG	A080582 002	
REXALL	0.25MG	A080637 001	
+ SANDOZ	0.1MG	N009838 001	
+	0.25MG	N009838 002	
SOLVAY	0.25MG	A080446 001	
TABLICAPS	0.25MG	A085207 001	
TEVA	0.1MG	A089020 001	Mar 07, 1985
	0.25MG	A089019 001	Mar 07, 1985
VALEANT PHARM INTL	0.1MG	N009667 001	
	0.25MG	N009667 002	
WATSON LABS	0.1MG	A080679 001	
	0.25MG	A080393 001	
	0.25MG	A085401 001	
	1MG	A080749 001	
WHITEWORTH TOWN PLSN	0.1MG	A080723 001	
	0.25MG	A080723 002	
	1MG	A080723 003	
SANDRIL			
LILLY	0.1MG	N009376 004	
	0.25MG	N009376 001	
SERPALAN			
LANNETT	0.1MG	N010124 001	
	0.25MG	N010124 002	
SERPANRAY			
PANRAY	0.1MG	N009391 001	
	0.25MG	N009391 002	
	1MG	N009391 004	
SERPASIL			
NOVARTIS	0.1MG	N009115 001	
	0.25MG	N009115 003	
	1MG	N009115 004	
SERPATE			
VALE	0.1MG	N009453 001	
	0.25MG	N009453 002	
SERPIVITE			
VITARINE	0.25MG	N009645 002	

RESERPINE; TRICHLORMETHIAZIDE

TABLET; ORAL

METATENSIN #2

SANOFI AVENTIS US	0.1MG;2MG	N012972 001	
METATENSIN #4			
SANOFI AVENTIS US	0.1MG;4MG	N012972 002	
NAQUIVAL			
SCHERING	0.1MG;4MG	N012265 003	
TRICHLORMETHIAZIDE W/ RESERPINE			
WATSON LABS	0.1MG;4MG	A085248 001	

RIBAVIRIN

CAPSULE; ORAL

REBETOL

MERCK SHARP DOHME	200MG**Indicated for use and comarketed with Interferon ALFA-2B, Recombinant (INTRON A), as Rebetrone Combination Therapy**	N020903 001	Jun 03, 1998
+	200MG **	N020903 002	Jul 25, 2001
RIBASPHERE			
CHARTWELL RX	200MG	A076203 001	Apr 06, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RIBAVIRIN

CAPSULE; ORAL

RIBAVIRIN

CHARTWELL RX

200MG

A076192 001 Apr 06, 2004

TEVA

200MG

A076277 001 Oct 04, 2004

SOLUTION; ORAL

REBETOL

+ SCHERING

40MG/ML

N021546 001 Jul 29, 2003

TABLET; ORAL

COPEGUS

+ ROCHE

200MG **

N021511 001 Dec 03, 2002

+

400MG **

N021511 002 Jun 21, 2005

RIBAVIRIN

BEXIMCO PHARMS USA

200MG

A202546 001 Aug 12, 2014

400MG

A202546 002 Aug 12, 2014

500MG

A202546 003 Aug 12, 2014

600MG

A202546 004 Aug 12, 2014

CHARTWELL RX

200MG

A077456 001 Dec 05, 2005

400MG

A077456 002 Dec 05, 2005

600MG

A077456 003 Dec 05, 2005

HERITAGE PHARMA AVET

200MG

A077053 001 Dec 05, 2005

ZYDUS PHARMS USA

400MG

A077094 002 Mar 16, 2007

500MG

A077094 004 Apr 18, 2008

600MG

A077094 003 Mar 16, 2007

RIFAMPIN

CAPSULE; ORAL

RIFADIN

SANOFI AVENTIS US

150MG

A062303 001

+

300MG

N050420 001

INJECTABLE; INJECTION

RIFAMPIN

AVET LIFESCIENCES

600MG/VIAL

A204101 001 Aug 18, 2014

WATSON PHARMS TEVA

600MG/VIAL

A206736 001 Jan 19, 2016

RILUZOLE

TABLET; ORAL

RILUZOLE

APOTEX CORP

50MG

A091300 001 Jun 18, 2013

DAITO PHARMS CO LTD

50MG

A204430 001 Oct 16, 2018

RIMANTADINE HYDROCHLORIDE

SYRUP; ORAL

FLUMADINE

FOREST LABS

50MG/5ML

N019650 001 Sep 17, 1993

TABLET; ORAL

FLUMADINE

+ SUN PHARM INDS INC

100MG

N019649 001 Sep 17, 1993

RIMANTADINE HYDROCHLORIDE

CHARTWELL RX

100MG

A076375 001 Jan 14, 2003

IMPAX LABS INC

100MG

A075916 001 Nov 02, 2001

RIMEXOLONE

SUSPENSION/DROPS; OPHTHALMIC

VEXOL

HARROW EYE

1%

N020474 001 Dec 30, 1994

RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL

+ APIL

5MG

N020835 002 Apr 14, 2000

+

30MG

N020835 001 Mar 27, 1998

+

75MG **

N020835 004 Apr 16, 2007

RISEDRONATE SODIUM

HANGZHOU BINJIANG

35MG

A207516 001 Feb 15, 2019

MYLAN

5MG

A200477 001 Nov 30, 2015

30MG

A200477 002 Nov 30, 2015

35MG

A200477 003 Nov 30, 2015

75MG

A200477 004 Jun 10, 2014

150MG

A200477 005 Jun 10, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RISEDRONATE SODIUM

TABLET, DELAYED RELEASE;ORAL

RISEDRONATE SODIUM

IMPAX LABS INC	35MG
ZYDUS PHARMS	35MG

A205066	001	Jun 29, 2018
A203822	001	Sep 11, 2018

RISPERIDONE

SOLUTION;ORAL

RISPERIDONE

ANI PHARMS	1MG/ML
APOZEAL PHARMS	1MG/ML
PRECISION DOSE	1MG/ML
WOCKHARDT	1MG/ML

A076440	001	Jan 30, 2009
A078909	001	Jul 29, 2009
A076797	001	Jun 28, 2010
A078744	001	Oct 08, 2009

TABLET;ORAL

RISPERDAL

JANSSEN PHARMS	5MG
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N020272	005	Dec 29, 1993
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RISPERIDONE

DASH PHARMS	0.25MG
	0.5MG
	1MG
	2MG
	3MG
	4MG

A076288	001	Sep 15, 2008
A076288	002	Sep 15, 2008
A076288	003	Sep 15, 2008
A076288	004	Sep 15, 2008
A076288	005	Sep 15, 2008
A076288	006	Sep 15, 2008

HERITAGE PHARMA AVET	0.25MG
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A076228	001	Jun 30, 2008
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	0.25MG
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A077769	001	Oct 16, 2008
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	0.5MG
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A076228	002	Jun 30, 2008
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	0.5MG
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A077769	002	Oct 16, 2008
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	1MG
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A076228	003	Jun 30, 2008
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	1MG
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A077769	003	Oct 16, 2008
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	2MG
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A076228	004	Jun 30, 2008
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	2MG
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A077769	004	Oct 16, 2008
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	3MG
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A076228	005	Jun 30, 2008
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	3MG
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A077769	005	Oct 16, 2008
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	4MG
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A076228	006	Jun 30, 2008
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	4MG
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A077769	006	Oct 16, 2008
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JUBILANT CADISTA	0.25MG
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A078828	001	Mar 23, 2009
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	0.5MG
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A078828	002	Mar 23, 2009
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	1MG
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A078828	003	Mar 23, 2009
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	2MG
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A078828	004	Mar 23, 2009
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	3MG
--	-----

A078828	005	Mar 23, 2009
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	4MG
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A078828	006	Mar 23, 2009
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RATIOPHARM	0.25MG
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A077784	001	Jun 08, 2010
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	0.5MG
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A077784	002	Jun 08, 2010
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	1MG
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A077784	003	Jun 08, 2010
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	2MG
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A077784	004	Jun 08, 2010
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	3MG
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A077784	005	Jun 08, 2010
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	4MG
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A077784	006	Jun 08, 2010
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SUN PHARM INDS INC	0.25MG
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A078036	001	Mar 10, 2014
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	0.5MG
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A078036	002	Mar 10, 2014
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	1MG
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A078036	003	Mar 10, 2014
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	2MG
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A078036	004	Mar 10, 2014
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	3MG
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A078036	005	Mar 10, 2014
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	4MG
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A078036	006	Mar 10, 2014
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SYNTHON PHARMS	0.25MG
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A078187	001	Oct 22, 2009
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	0.5MG
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A078187	002	Oct 22, 2009
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	1MG
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A078187	003	Oct 22, 2009
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	2MG
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A078187	004	Oct 22, 2009
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	3MG
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A078187	005	Oct 22, 2009
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	4MG
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A078187	006	Oct 22, 2009
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WATSON LABS	0.25MG
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A077860	001	Dec 05, 2008
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	0.5MG
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A077860	002	Dec 05, 2008
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	1MG
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A077860	003	Dec 05, 2008
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	2MG
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A077860	004	Dec 05, 2008
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	3MG
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A077860	005	Dec 05, 2008
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	4MG
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A077860	006	Dec 05, 2008
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WEST WARD PHARMS	0.25MG
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A078740	001	May 29, 2009
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	0.5MG
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A078740	002	May 29, 2009
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	1MG
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A078740	003	May 29, 2009
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	2MG
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A078740	004	May 29, 2009
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	3MG
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A078740	005	May 29, 2009
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RISPERIDONE

TABLET; ORAL

RISPERIDONE

4MG

A078740 006 May 29, 2009

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

+	JANSSEN PHARMS	0.5MG **
+		1MG **
+		2MG **
+		3MG **
+		4MG **

N021444	001	Apr 02, 2003
N021444	002	Apr 02, 2003
N021444	003	Apr 02, 2003
N021444	004	Dec 23, 2004
N021444	005	Dec 23, 2004

RISPERIDONE

ACTAVIS LABS FL INC

0.5MG
1MG
2MG
3MG
4MG

A076996	001	Apr 19, 2011
A076996	002	Apr 19, 2011
A076996	003	Apr 19, 2011
A076996	004	Apr 19, 2011
A076996	005	Apr 19, 2011

CHARTWELL RX

0.5MG
1MG
2MG

A076908	001	Mar 12, 2012
A076908	002	Mar 12, 2012
A076908	003	Mar 12, 2012

DASH PHARMS

0.25MG
0.5MG
1MG
2MG
3MG
4MG

A091537	006	Feb 12, 2013
A091537	001	Mar 30, 2011
A091537	002	Mar 30, 2011
A091537	003	Mar 30, 2011
A091537	004	Mar 30, 2011
A091537	005	Mar 30, 2011

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION

RITODRINE HYDROCHLORIDE

ABRAXIS PHARM

10MG/ML
15MG/ML

A071188	001	Jul 23, 1987
A071189	001	Jul 23, 1987
A071618	001	Feb 28, 1991
A071619	001	Feb 28, 1991

HOSPIRA

10MG/ML
15MG/ML

RITODRINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA

30MG/100ML

A071438 001 Jan 22, 1991

YUTOPAR

ASTRAZENECA

10MG/ML
15MG/ML

N018580	001	
N018580	002	Sep 27, 1984

TABLET; ORAL

YUTOPAR

ASTRAZENECA

10MG

N018555 001

RITONAVIR

CAPSULE; ORAL

NORVIR

ABBOTT

100MG

N020680 001 Mar 01, 1996

+ ABBVIE

100MG **

N020945 001 Jun 29, 1999

RITONAVIR

HIKMA

100MG

A205801 001 Dec 03, 2020

SOLUTION; ORAL

NORVIR

+ ABBVIE

80MG/ML

N020659 001 Mar 01, 1996

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

EXELON

+ NOVARTIS

EQ 1.5MG BASE **

N020823 003 Apr 21, 2000

+

EQ 3MG BASE **

N020823 004 Apr 21, 2000

+

EQ 4.5MG BASE **

N020823 005 Apr 21, 2000

+

EQ 6MG BASE **

N020823 006 Apr 21, 2000

RIVASTIGMINE TARTRATE

APOTEX INC

EQ 1.5MG BASE
EQ 3MG BASE
EQ 4.5MG BASE
EQ 6MG BASE

A091072	001	May 16, 2013
A091072	002	May 16, 2013
A091072	003	May 16, 2013
A091072	004	May 16, 2013

SOLUTION; ORAL

EXELON

NOVARTIS

EQ 2MG BASE/ML

N021025 001 Apr 21, 2000

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RIZATRIPTAN BENZOATE

TABLET; ORAL

MAXALT

+	ORGANON LLC	EQ 5MG BASE **	N020864	001	Jun 29, 1998
	RIZATRIPTAN BENZOATE				
	APOTEX INC	EQ 5MG BASE	A202244	001	Dec 31, 2012
		EQ 10MG BASE	A202244	002	Dec 31, 2012
	AVET LIFESCIENCES	EQ 5MG BASE	A204090	001	Nov 26, 2013
		EQ 10MG BASE	A204090	002	Nov 26, 2013
	JUBILANT GENERICS	EQ 5MG BASE	A203252	001	Dec 31, 2014
		EQ 10MG BASE	A203252	002	Dec 31, 2014
	SANDOZ	EQ 5MG BASE	A079230	001	Dec 31, 2012
		EQ 10MG BASE	A079230	002	Dec 31, 2012
	UNICHEM	EQ 5MG BASE	A207836	001	Mar 07, 2017
		EQ 10MG BASE	A207836	002	Mar 07, 2017

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

+	ORGANON	EQ 5MG BASE **	N020865	001	Jun 29, 1998
	RIZATRIPTAN BENZOATE				
	APOTEX INC	EQ 5MG BASE	A202477	001	Jul 01, 2013
		EQ 10MG BASE	A202477	002	Jul 01, 2013
	JUBILANT GENERICS	EQ 5MG BASE	A203334	001	Oct 16, 2015
		EQ 10MG BASE	A203334	002	Oct 16, 2015
	MYLAN PHARMS INC	EQ 5MG BASE	A078173	001	Dec 31, 2012
		EQ 10MG BASE	A078173	002	Dec 31, 2012
	SANDOZ	EQ 5MG BASE	A078739	001	Jul 01, 2013
		EQ 10MG BASE	A078739	002	Jul 01, 2013
	UNICHEM	EQ 5MG BASE	A207835	001	Mar 07, 2017
		EQ 10MG BASE	A207835	002	Mar 07, 2017

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ROCURONIUM BROMIDE

	TEVA PHARMS	50MG/5ML (10MG/ML)	A078717	001	Nov 26, 2008
		100MG/10ML (10MG/ML)	A078717	002	Nov 26, 2008
	ZEMURON				
+	ORGANON USA INC	50MG/5ML (10MG/ML) **	N020214	001	Mar 17, 1994
+		10MG/ML (10MG/ML) **	N020214	002	Mar 17, 1994
+		100MG/10ML (10MG/ML) **	N020214	003	Mar 17, 1994

ROFECOXIB

SUSPENSION; ORAL

VIOXX

	MERCK	12.5MG/5ML	N021052	001	May 20, 1999
		25MG/5ML	N021052	002	May 20, 1999

TABLET; ORAL

VIOXX

	MERCK	12.5MG	N021042	001	May 20, 1999
		25MG	N021042	002	May 20, 1999
		50MG	N021042	003	Feb 25, 2000

ROFLUMILAST

TABLET; ORAL

ROFLUMILAST

	BRECKENRIDGE	500MCG	A208236	001	Oct 03, 2018
	TORRENT	500MCG	A208272	001	Aug 06, 2018

ROLAPITANT HYDROCHLORIDE

EMULSION; INTRAVENOUS

VARUBI

+	TERSERA	EQ 166.5MG BASE/92.5ML (EQ 1.8MG BASE/ML)	N208399	001	Oct 25, 2017
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ROMIDEPSIN

SOLUTION; INTRAVENOUS

ROMIDEPSIN

+	TEVA PHARMS USA INC	10MG/2ML (5MG/ML)	N208574	001	Mar 13, 2020
+		27.5MG/5.5ML (5MG/ML)	N208574	002	Mar 13, 2020

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

REQUIP

+	GLAXOSMITHKLINE LLC	EQ 0.25MG BASE **	N020658 001	Sep 19, 1997
+		EQ 0.5MG BASE **	N020658 002	Sep 19, 1997
+		EQ 1MG BASE **	N020658 003	Sep 19, 1997
+		EQ 2MG BASE **	N020658 004	Sep 19, 1997
+		EQ 3MG BASE **	N020658 006	Jan 27, 1999
+		EQ 4MG BASE **	N020658 007	Jan 27, 1999
+		EQ 5MG BASE **	N020658 005	Sep 19, 1997

ROPINIROLE HYDROCHLORIDE

COSETTE

		EQ 0.25MG BASE	A077460 001	May 05, 2008
		EQ 0.5MG BASE	A077460 002	May 05, 2008
		EQ 1MG BASE	A077460 003	May 05, 2008
		EQ 2MG BASE	A077460 004	May 05, 2008
		EQ 3MG BASE	A077460 005	May 05, 2008
		EQ 4MG BASE	A077460 006	May 05, 2008
		EQ 5MG BASE	A077460 007	May 19, 2008

EPIC PHARMA LLC

		EQ 0.25MG BASE	A078230 001	May 20, 2008
		EQ 0.5MG BASE	A078230 002	May 20, 2008
		EQ 1MG BASE	A078230 003	May 20, 2008
		EQ 2MG BASE	A078230 004	May 20, 2008
		EQ 3MG BASE	A078230 005	May 20, 2008
		EQ 4MG BASE	A078230 006	May 20, 2008
		EQ 5MG BASE	A078230 007	May 20, 2008

HIKMA

		EQ 0.25MG BASE	A077852 001	May 05, 2008
		EQ 0.5MG BASE	A077852 002	May 05, 2008
		EQ 1MG BASE	A077852 003	May 05, 2008
		EQ 2MG BASE	A077852 004	May 05, 2008
		EQ 3MG BASE	A077852 005	May 05, 2008
		EQ 4MG BASE	A077852 006	May 05, 2008
		EQ 5MG BASE	A077852 007	May 19, 2008

MYLAN

		EQ 0.25MG BASE	A078881 001	May 05, 2008
		EQ 0.5MG BASE	A078881 002	May 05, 2008
		EQ 1MG BASE	A078881 003	May 05, 2008
		EQ 2MG BASE	A078881 004	May 05, 2008
		EQ 3MG BASE	A078881 005	May 05, 2008
		EQ 4MG BASE	A078881 006	May 05, 2008
		EQ 5MG BASE	A078881 007	May 19, 2008

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

+	GLAXOSMITHKLINE LLC	EQ 2MG BASE **	N022008 001	Jun 13, 2008
+		EQ 3MG BASE **	N022008 002	Jun 13, 2008
+		EQ 4MG BASE **	N022008 003	Jun 13, 2008
+		EQ 6MG BASE **	N022008 006	Apr 10, 2009
+		EQ 8MG BASE **	N022008 004	Jun 13, 2008
+		EQ 12MG BASE **	N022008 005	Oct 31, 2008

ROPINIROLE HYDROCHLORIDE

MYLAN PHARMS INC

		EQ 2MG BASE	A200462 001	Oct 15, 2012
		EQ 3MG BASE	A200462 002	Oct 15, 2012
		EQ 4MG BASE	A200462 003	Oct 15, 2012
		EQ 6MG BASE	A200462 004	Oct 15, 2012
		EQ 8MG BASE	A200462 005	Oct 15, 2012
		EQ 12MG BASE	A200462 006	Oct 15, 2012

WATSON LABS INC

		EQ 2MG BASE	A200431 001	Jun 06, 2012
		EQ 4MG BASE	A200431 002	Jun 06, 2012
		EQ 6MG BASE	A200431 003	Jun 06, 2012
		EQ 8MG BASE	A200431 004	Jun 06, 2012
		EQ 12MG BASE	A200431 005	Jun 06, 2012

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

NAROPIN

+	FRESENIUS KABI USA	50MG/10ML (5MG/ML) **	N020533 013	May 01, 1998
+		75MG/10ML (7.5MG/ML) **	N020533 012	Sep 24, 1996

ROPIVACAINE HYDROCHLORIDE

RISING

		40MG/20ML (2MG/ML)	A090318 001	Sep 23, 2014
		150MG/30ML (5MG/ML)	A090318 002	Sep 23, 2014
		150MG/20ML (7.5MG/ML)	A090318 003	Sep 23, 2014
		200MG/20ML (10MG/ML)	A090318 004	Sep 23, 2014

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ROSE BENGAL SODIUM I-131

INJECTABLE; INJECTION

ROBENGATOPE

BRACCO

0.5mCi/VIAL

N016224 001

1mCi/VIAL

N016224 002

2mCi/VIAL

N016224 003

SODIUM ROSE BENGAL I 131

SORIN

0.5mCi/ML

N017318 001

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

+ WOODWARD

EQ 2MG BASE

N021071 002 May 25, 1999

+

EQ 4MG BASE

N021071 003 May 25, 1999

+

EQ 8MG BASE **

N021071 004 May 25, 1999

ROSIGLITAZONE MALEATE

ANI PHARMS

EQ 2MG BASE

A076747 001 Jan 25, 2013

EQ 4MG BASE

A076747 002 Jan 25, 2013

EQ 8MG BASE

A076747 003 Jan 25, 2013

ROSUVASTATIN CALCIUM

CAPSULE; ORAL

EZALLOR SPRINKLE

+ SUN PHARM

EQ 5MG BASE

N208647 001 Dec 18, 2018

+

EQ 10MG BASE

N208647 002 Dec 18, 2018

+

EQ 20MG BASE

N208647 003 Dec 18, 2018

+

EQ 40MG BASE

N208647 004 Dec 18, 2018

TABLET; ORAL

ROSUVASTATIN CALCIUM

AMNEAL PHARMS CO

EQ 5MG BASE

A208850 001 Oct 16, 2018

EQ 10MG BASE

A208850 002 Oct 16, 2018

EQ 20MG BASE

A208850 003 Oct 16, 2018

EQ 40MG BASE

A208850 004 Oct 16, 2018

APOTEX

EQ 5MG BASE

A079145 001 Jul 19, 2016

EQ 10MG BASE

A079145 002 Jul 19, 2016

EQ 20MG BASE

A079145 003 Jul 19, 2016

EQ 40MG BASE

A079145 004 Jul 19, 2016

INVENTIA

EQ 5MG BASE

A207653 001 Feb 05, 2021

EQ 10MG BASE

A207653 002 Feb 05, 2021

EQ 20MG BASE

A207653 003 Feb 05, 2021

EQ 40MG BASE

A207653 004 Feb 05, 2021

STRIDES PHARMA

EQ 5MG BASE

A079161 001 Jul 19, 2016

EQ 10MG BASE

A079161 002 Jul 19, 2016

EQ 20MG BASE

A079161 003 Jul 19, 2016

EQ 40MG BASE

A079161 004 Jul 19, 2016

SUNSHINE

EQ 5MG BASE

A210667 001 Apr 01, 2020

EQ 10MG BASE

A210667 002 Apr 01, 2020

EQ 20MG BASE

A210667 003 Apr 01, 2020

EQ 40MG BASE

A210667 004 Apr 01, 2020

TEVA PHARMS USA

EQ 5MG BASE

A079166 001 Jul 19, 2016

EQ 10MG BASE

A079166 002 Jul 19, 2016

EQ 20MG BASE

A079166 003 Jul 19, 2016

EQ 40MG BASE

A079166 004 Jul 19, 2016

ZHEJIANG JINGXIN

EQ 5MG BASE

A206513 001 Mar 01, 2019

EQ 10MG BASE

A206513 002 Mar 01, 2019

EQ 20MG BASE

A206513 003 Mar 01, 2019

EQ 40MG BASE

A206513 004 Mar 01, 2019

RUFINAMIDE

TABLET; ORAL

BANZEL

+ EISAI INC

100MG **

N021911 001 Nov 14, 2008

SAFFLOWER OIL

INJECTABLE; INJECTION

LIPOSYN 10%

ABBOTT

10% (10GM/100ML)

N018203 001

LIPOSYN 20%

ABBOTT

20% (20GM/100ML)

N018614 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SAFFLOWER OIL; SOYBEAN OIL

INJECTABLE; INJECTION

LIPOSYN II 10%

HOSPIRA

5%;5% (5GM/100ML)

N018997 001 Aug 27, 1984

LIPOSYN II 20%

HOSPIRA

10%;10% (10GM/100ML)

N018991 001 Aug 27, 1984

SAFINAMIDE MESYLATE

TABLET; ORAL

SAFINAMIDE MESYLATE

AUROBINDO PHARMA

EQ 50MG BASE

A215902 001 Jun 14, 2023

EQ 100MG BASE

A215902 002 Jun 14, 2023

SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

SEREVENT

GLAXOSMITHKLINE

EQ 0.021MG BASE/INH

N020236 001 Feb 04, 1994

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION

QUADRAMET

+

LANTHEUS MEDICAL

50mCi/ML

N020570 001 Mar 28, 1997

SAQUINAVIR

CAPSULE; ORAL

FORTOVASE

+

HOFFMANN LA ROCHE

200MG **

N020828 001 Nov 07, 1997

SAQUINAVIR MESYLATE

CAPSULE; ORAL

INVIRASE

+

HOFFMANN LA ROCHE

EQ 200MG BASE

N020628 001 Dec 06, 1995

TABLET; ORAL

INVIRASE

+

HOFFMANN-LA ROCHE

EQ 500MG BASE

N021785 001 Dec 17, 2004

SARALASIN ACETATE

INJECTABLE; INJECTION

SARENIN

PROCTER AND GAMBLE

EQ 0.6MG BASE/ML

N018009 001

SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

ONGLYZA

+

ASTRAZENECA AB

EQ 2.5MG BASE

N022350 001 Jul 31, 2009

+

EQ 5MG BASE

N022350 002 Jul 31, 2009

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECOBARBITAL SODIUM

ANABOLIC

100MG

A084422 001

BARR

100MG

A084225 001

EVERYLIFE

100MG

A085895 001

HALSEY

100MG

A084676 001

IVAX PHARMS

100MG

A085869 001

KV PHARM

100MG

A085285 001

LANNETT

50MG

A085909 001

100MG

A085903 001

PARKE DAVIS

100MG

A084762 001

PERRIGO

100MG

A084561 001

PUREPAC PHARM

100MG

A085867 001

VALEANT PHARM INTL

100MG

A085477 001

VITARINE

100MG

A085898 001

100MG

A086273 001

WATSON LABS

100MG

A085792 001

WEST WARD

100MG

A084926 001

WHITWORTH TOWN PLSN

100MG

A085798 001

WYETH AYERST

100MG

A086390 001

SECONAL SODIUM

VALEANT PHARMS NORTH

50MG

A086101 001 Oct 03, 1983

100MG

A086101 002 Oct 03, 1983

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SECOBARBITAL SODIUM

INJECTABLE; INJECTION

SECOBARBITAL SODIUM

ELKINS SINN 100MG/VIAL A083281 001

WYETH AYERST 50MG/ML A083262 001

SECONAL SODIUM

LILLY 50MG/ML ** N007392 002

SUPPOSITORY; RECTAL

SECONAL SODIUM

LILLY 30MG A086530 001

60MG A086530 002

120MG A086530 003

200MG A086530 004

SECRETIN

INJECTABLE; INJECTION

SECRETIN-FERRING

FERRING 75CU/VIAL N018290 001

SECRETIN SYNTHETIC PORCINE

FOR SOLUTION; INTRAVENOUS

SECREFLO

CHIRHOCLIN 16MCG/VIAL N021136 001 Apr 04, 2002

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

ELDEPRYL

+ SOMERSET 5MG ** N020647 001 May 15, 1996

SELEGILINE HYDROCHLORIDE

LANNETT CO INC 5MG A075145 001 Sep 15, 2003

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

BAYSHORE PHARMS LLC 5MG A074912 001 Apr 30, 1998

CHARTWELL MOLECULES 5MG A074565 001 Aug 02, 1996

5MG A074641 001 Aug 02, 1996

COSETTE 5MG A074744 001 Jan 27, 1997

5MG A074756 001 Nov 25, 1998

DASH PHARMS NATCO 5MG A074866 001 Nov 26, 1997

G AND W LABS INC 5MG A074537 001 Aug 02, 1996

+ SOMERSET 5MG ** N019334 001 Jun 05, 1989

SELENIUM SULFIDE

LOTION; SHAMPOO; TOPICAL

EXSEL

ALLERGAN HERBERT 2.5% A083892 001

SELENIUM SULFIDE

ACTAVIS MID ATLANTIC 2.5% A084394 001

COSETTE 2.5% A086209 001

IVAX PHARMS 2.5% A085777 001

PHARMOBEDIANT CNSLTG 2.5% A088228 001 Sep 01, 1983

SELSUN

+ CHATTEM 2.5% ** N007936 001

SELENOMETHIONINE SE-75

INJECTABLE; INJECTION

SELENOMETHIONINE SE 75

GE HEALTHCARE 250uCi/ML N017257 001

MALLINCKRODT 100uCi/ML N017098 001

PHARMALUCENCE 500uCi/ML N017322 001

SETHOTOPE

BRACCO 85-550uCi/ML N017047 001

SELEXIPAG

TABLET; ORAL

SELEXIPAG

ZYDUS 0.2MG A214302 001 Dec 21, 2022

0.4MG A214302 002 Dec 21, 2022

0.6MG A214302 003 Dec 21, 2022

0.8MG A214302 004 Dec 21, 2022

1MG A214302 005 Dec 21, 2022

1.2MG A214302 006 Dec 21, 2022

1.4MG A214302 007 Dec 21, 2022

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SELEXIPAGTABLET; ORAL
SELEXIPAG

1.6MG

A214302 008 Dec 21, 2022

SERMORELIN ACETATEINJECTABLE; INJECTION
GEREF

+ EMD SERONO

EQ 0.05MG BASE/AMP **

N019863 001 Dec 28, 1990

+ EMD SERONO INC

EQ 0.5MG BASE/VIAL **

N020443 001 Sep 26, 1997

+

EQ 1MG BASE/VIAL **

N020443 002 Sep 26, 1997

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

RANBAXY LABS LTD

EQ 20MG BASE/ML

A078053 001 Feb 05, 2007

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

ANDA REPOSITORY

EQ 25MG BASE

A077818 001 Feb 06, 2007

EQ 50MG BASE

A077818 002 Feb 06, 2007

EQ 100MG BASE

A077818 003 Feb 06, 2007

CHARTWELL MOLECULAR

EQ 25MG BASE

A077162 001 Feb 06, 2007

EQ 50MG BASE

A077162 002 Feb 06, 2007

EQ 100MG BASE

A077162 003 Feb 06, 2007

FOSUN PHARMA

EQ 25MG BASE

A077713 001 Feb 06, 2007

EQ 50MG BASE

A077713 002 Feb 06, 2007

EQ 100MG BASE

A077713 003 Feb 06, 2007

HERITAGE PHARMA AVET

EQ 25MG BASE

A077299 001 Feb 06, 2007

EQ 25MG BASE

A077345 001 Feb 06, 2007

EQ 25MG BASE

A077663 001 Feb 06, 2007

EQ 50MG BASE

A077299 002 Feb 06, 2007

EQ 50MG BASE

A077345 002 Feb 06, 2007

EQ 50MG BASE

A077663 002 Feb 06, 2007

EQ 100MG BASE

A077299 003 Feb 06, 2007

EQ 100MG BASE

A077345 003 Feb 06, 2007

EQ 100MG BASE

A077663 003 Feb 06, 2007

HIKMA PHARMS

EQ 25MG BASE

A077864 001 Aug 10, 2009

EQ 50MG BASE

A077864 002 Aug 10, 2009

EQ 100MG BASE

A077864 003 Aug 10, 2009

IVAX SUB TEVA PHARMS

EQ 25MG BASE

A075719 003 Jun 30, 2006

EQ 50MG BASE

A075719 001 Jun 30, 2006

EQ 100MG BASE

A075719 002 Jun 30, 2006

MYLAN

EQ 25MG BASE

A076671 001 Feb 06, 2007

EQ 50MG BASE

A076671 002 Feb 06, 2007

EQ 100MG BASE

A076671 003 Feb 06, 2007

MYLAN PHARMS INC

EQ 25MG BASE

A076540 001 Mar 20, 2007

EQ 25MG BASE

A078626 001 Jan 31, 2008

EQ 50MG BASE

A076540 002 Mar 20, 2007

EQ 50MG BASE

A078626 002 Jan 31, 2008

EQ 100MG BASE

A076540 003 Mar 20, 2007

EQ 100MG BASE

A078626 003 Jan 31, 2008

SUN PHARM INDS (IN)

EQ 25MG BASE

A078108 001 Feb 06, 2007

EQ 50MG BASE

A078108 002 Feb 06, 2007

EQ 100MG BASE

A078108 003 Feb 06, 2007

SUN PHARM INDS LTD

EQ 25MG BASE

A077977 001 Feb 06, 2007

EQ 50MG BASE

A077977 002 Feb 06, 2007

EQ 100MG BASE

A077977 003 Feb 06, 2007

EQ 150MG BASE

A077977 004 Feb 06, 2007

EQ 200MG BASE

A077977 005 Feb 06, 2007

TORRENT PHARMS

EQ 25MG BASE

A077765 001 Feb 06, 2007

EQ 50MG BASE

A077765 002 Feb 06, 2007

EQ 100MG BASE

A077765 003 Feb 06, 2007

ZYDUS

EQ 25MG BASE

A077106 001 Feb 06, 2007

EQ 50MG BASE

A077106 002 Feb 06, 2007

EQ 100MG BASE

A077106 003 Feb 06, 2007

ZOLOFT

+ VIATRIS

EQ 150MG BASE **

N019839 003 Dec 30, 1991

+

EQ 200MG BASE **

N019839 004 Dec 30, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SEVELAMER CARBONATE

FOR SUSPENSION;ORAL

SEVELAMER CARBONATE

LUPIN LTD

800MG/PACKET

A201513 001 Dec 23, 2021

2.4GM/PACKET

A201513 002 Dec 23, 2021

TABLET;ORAL

SEVELAMER CARBONATE

IMPAX LABS INC

800MG

A090975 001 Oct 23, 2017

LUPIN LTD

800MG

A204600 001 Jan 14, 2021

STRIDES PHARMA

800MG

A201069 001 Aug 05, 2020

SEVELAMER HYDROCHLORIDE

CAPSULE;ORAL

RENAGEL

GENZYME

403MG

N020926 001 Oct 30, 1998

TABLET;ORAL

SEVELAMER HYDROCHLORIDE

RISING

400MG

A201068 001 Dec 14, 2020

800MG

A201068 002 Dec 14, 2020

SIBUTRAMINE HYDROCHLORIDE

CAPSULE;ORAL

MERIDIA

ABBOTT

5MG

N020632 001 Nov 22, 1997

10MG

N020632 002 Nov 22, 1997

15MG

N020632 003 Nov 22, 1997

SILDENAFIL CITRATE

FOR SUSPENSION;ORAL

SILDENAFIL CITRATE

TRIS PHARMA INC

EQ 10MG BASE/ML

A212312 001 Nov 17, 2021

TABLET;ORAL

SILDENAFIL CITRATE

ACTAVIS GRP PTC

EQ 20MG BASE

A200149 001 Feb 25, 2013

APOTEX CORP

EQ 20MG BASE

A091379 001 Nov 06, 2012

PERRIGO R AND D

EQ 25MG BASE

A205791 001 Apr 23, 2020

EQ 50MG BASE

A205791 002 Apr 23, 2020

WATSON LABS INC

EQ 25MG BASE

A202506 001 Nov 25, 2020

EQ 50MG BASE

A202506 002 Nov 25, 2020

EQ 100MG BASE

A202506 003 Nov 25, 2020

SILODOSIN

CAPSULE;ORAL

SILODOSIN

ALEMBIC

4MG

A211731 001 Nov 22, 2019

8MG

A211731 002 Nov 22, 2019

ZYDUS PHARMS

4MG

A204816 001 Dec 08, 2022

8MG

A204816 002 Dec 08, 2022

SILVER SULFADIAZINE

CREAM;TOPICAL

SSD AF

DR REDDYS LA

1%

N018578 003 Jul 11, 1990

DRESSING;TOPICAL

SILDAFLO

FRANKLIN PHARMS

1%

N019608 001 Nov 30, 1989

SIMEPREVIR SODIUM

CAPSULE;ORAL

OLYSIO

+

JANSSEN PRODS

EQ 150MG BASE

N205123 001 Nov 22, 2013

SIMETHICONE-CELLULOSE

SUSPENSION;ORAL

SONORX

BRACCO

7.5MG/ML

N020773 001 Oct 29, 1998

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SIMVASTATIN

TABLET;ORAL

SIMVASTATIN

CHARTWELL RX	5MG	A077766 001	Dec 20, 2006
	10MG	A077766 002	Dec 20, 2006
	20MG	A077766 003	Dec 20, 2006
	40MG	A077766 004	Dec 20, 2006
	80MG	A077766 005	Dec 20, 2006
IVAX SUB TEVA PHARMS	5MG	A076052 001	Jun 23, 2006
	10MG	A076052 002	Jun 23, 2006
	20MG	A076052 003	Jun 23, 2006
	40MG	A076052 004	Jun 23, 2006
	80MG	A076052 005	Dec 20, 2006
MYLAN PHARMS INC	5MG	A090868 001	Jun 08, 2010
	10MG	A090868 002	Jun 08, 2010
	20MG	A090868 003	Jun 08, 2010
	40MG	A090868 004	Jun 08, 2010
	80MG	A090868 005	Jun 08, 2010
SUN PHARM INDS LTD	5MG	A076285 001	Dec 20, 2006
	10MG	A076285 002	Dec 20, 2006
	20MG	A076285 003	Dec 20, 2006
	40MG	A076285 004	Dec 20, 2006
	80MG	A076285 005	Jun 23, 2006
WATSON LABS TEVA	5MG	A076685 001	Dec 20, 2006
	10MG	A076685 002	Dec 20, 2006
	20MG	A076685 003	Dec 20, 2006
	40MG	A076685 004	Dec 20, 2006
	80MG	A076685 005	Dec 20, 2006

ZOCOR

+ ORGANON 80MG **

N019766 005 Jul 10, 1998

TABLET, ORALLY DISINTEGRATING;ORAL

SIMVASTATIN

+ SYNTHON PHARMS	10MG	N021961 001	Oct 09, 2007
	20MG	N021961 002	Oct 09, 2007
	40MG	N021961 003	Oct 09, 2007
	80MG	N021961 004	Oct 09, 2007

SIMVASTATIN; SITAGLIPTIN PHOSPHATE

TABLET;ORAL

JUVISYNC

+ MERCK SHARP DOHME	10MG;EQ 50MG BASE **	N202343 004	Sep 18, 2012
	10MG;EQ 100MG BASE **	N202343 001	Oct 07, 2011
	20MG;EQ 50MG BASE **	N202343 005	Sep 18, 2012
	20MG;EQ 100MG BASE **	N202343 002	Oct 07, 2011
	40MG;EQ 50MG BASE **	N202343 006	Sep 18, 2012
	40MG;EQ 100MG BASE **	N202343 003	Oct 07, 2011

SIROLIMUS

SOLUTION;ORAL

SIROLIMUS

TORRENT	1MG/ML	A215016 001	Dec 27, 2021
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TABLET;ORAL

RAPAMUNE

+ PF PRISM CV	5MG **	N021110 003	Feb 23, 2004
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SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION;ORAL

UCEPHAN

B BRAUN	100MG/ML;100MG/ML	N019530 001	Dec 23, 1987
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SODIUM BICARBONATE

INJECTABLE;INJECTION

SODIUM BICARBONATE

HOSPIRA	0.9MEQ/ML	A077394 001	Nov 09, 2005
	1MEQ/ML	A077394 002	Nov 09, 2005

SODIUM BICARBONATE IN PLASTIC CONTAINER

+ ABBOTT	0.9MEQ/ML **	N019443 001	Jun 03, 1986
	1MEQ/ML **	N019443 002	Jun 03, 1986

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM BICARBONATE; TARTARIC ACIDGRANULE, EFFERVESCENT; ORAL
BAROS

MALLINCKRODT INC 460MG/GM; 420MG/GM N018509 001 Aug 07, 1985

SODIUM CHLORIDEAEROSOL, METERED; INHALATION
BRONCHO SALINE

+ BLAIREX 0.9% N019912 001 Sep 03, 1992

INJECTABLE; INJECTION
SODIUM CHLORIDE

ABBOTT 20GM/100ML N017013 001

B BRAUN 20GM/100ML N017038 001

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN 450MG/100ML N018184 001

MILES 450MG/100ML N018503 001

SODIUM CHLORIDE 0.9%

+ MEDEFIL INC 9MG/ML (9MG/ML) N202832 001 Jan 06, 2012

+ 18MG/2ML (9MG/ML) N202832 002 Jan 06, 2012

+ 22.5MG/2.5ML (9MG/ML) N202832 003 Jan 06, 2012

+ 27MG/3ML (9MG/ML) N202832 004 Jan 06, 2012

+ 45MG/5ML (9MG/ML) N202832 005 Jan 06, 2012

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+ LIEBEL-FLARSHEIM 450MG/50ML (9MG/ML) N021569 001 Jul 27, 2006

SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER

+ ABRAXIS PHARM 234MG/ML ** N019329 001 Apr 22, 1987

SOLUTION; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABRAXIS PHARM 90MG/10ML (9MG/ML) A088909 002 May 15, 1985

270MG/30ML (9MG/ML) A088909 001 Feb 07, 1985

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABBOTT 90MG/10ML (9MG/ML) N019218 001 Jul 13, 1984

180MG/20ML (9MG/ML) N019218 002 Apr 30, 1985

+ HOSPIRA 18MG/2ML (9MG/ML) N018803 004 Jan 22, 2015

+ 27MG/3ML (9MG/ML) N018803 005 Jan 22, 2015

+ 45MG/5ML (9MG/ML) N018803 006 Jan 22, 2015

+ ICU MEDICAL INC 45MG/5ML (9MG/ML) N019217 002 Nov 18, 1998

+ 90MG/10ML (9MG/ML) N019217 003 Nov 18, 1998

+ 360MG/40ML (9MG/ML) N019217 004 Apr 01, 1999

JUBILANT CADISTA 27MG/3ML (9MG/ML) A203352 002 May 18, 2016

90MG/10ML (9MG/ML) A203352 001 May 18, 2016

MILES 450MG/50ML (9MG/ML) N018502 002

900MG/100ML N018502 001

SODIUM CHLORIDE 0.9% IN PLASTIC THERMOJECT KIT FOR CARDIAC OUTPUT USE

+ ICU MEDICAL INC 90MG/10ML (9MG/ML) N019217 001 Jul 13, 1984

SOLUTION; INTRAVENOUS

SODIUM CHLORIDE 14.6%

+ HOSPIRA 50MEQ/20ML (2.5MEQ/ML) ** N018897 001 Jul 20, 1984

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

BAXTER HLTHCARE 450MG/100ML N017864 001

450MG/100ML N018497 001 Feb 19, 1982

HOSPIRA 450MG/100ML N017670 001

450MG/100ML N018380 001

SODIUM CHLORIDE IN PLASTIC CONTAINER

MILES 900MG/100ML N018247 001

SODIUM CHROMATE CR-51

INJECTABLE; INJECTION

CHROMITOPE SODIUM

BRACCO 2mCi/VIAL N013993 002

200uCi/ML N013993 001

SODIUM CHROMATE CR 51

CURIUM 100uCi/ML N016708 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

FLUORINE F-18

+ GE HEALTHCARE

2mCi/ML **

N017042 001

SODIUM FLUORIDE F 18

+ NIH NCI DCTD

10-200mCi/ML **

N022494 001 Jan 26, 2011

SODIUM FLUORIDE F-18

DECATUR

10-200mCi/ML

A204464 001 Oct 21, 2014

SHERTECH LABS LLC

10-200mCi/ML

A204315 001 Sep 22, 2014

UCSF RODIOPHARM

10-200mCi/ML

A204437 001 Mar 13, 2014

UIHC PET IMAGING

10-200mCi/ML

A204462 001 Nov 17, 2015

UNIV TX MD ANDERSON

10-200mCi/ML

A203247 001 Dec 23, 2013

SODIUM FLUORIDE; TRICLOSAN

PASTE; DENTAL

COLGATE TOTAL

+ COLGATE PALMOLIVE

0.24%; 0.3%

N020231 001 Jul 11, 1997

SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

CARDINAL HEALTH 418

400uCi

N018671 003 May 27, 1982

GE HEALTHCARE

100uCi

N017630 001

200uCi

N017630 003 Jan 08, 1993

SOLUTION; ORAL

SODIUM IODIDE I 123

GE HEALTHCARE

2mCi/ML **

N017630 002

SODIUM IODIDE I-131

CAPSULE; ORAL

IODOTOPE

BRACCO

1-130mCi

N010929 001

1-150mCi

N010929 003

SODIUM IODIDE I 131

CIS

50uCi

N017316 001

100uCi

N017316 002

CURIUM

0.8-100mCi

N016515 002

+

0.8-100mCi

N016517 001

15-100uCi

N016517 002

JUBILANT

2-200mCi

N021305 004 Nov 18, 2004

SOLUTION; ORAL

HICON

JUBILANT

1-250mCi/0.25ML

N021305 002 Jan 24, 2003

1-500mCi/0.5ML

N021305 003 Jan 24, 2003

1-1000mCi/ML

N021305 005 Apr 04, 2006

IODOTOPE

BRACCO

7-106mCi/BOT

N010929 002

SODIUM IODIDE I 131

CIS

50mCi/ML

N017315 001

+

CURIUM

3.5-150mCi/VIAL

N016515 001

SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER

B BRAUN

1.87GM/100ML

N018186 001

BAXTER HLTHCARE

1.87GM/100ML

N016692 001

HOSPIRA

1.87GM/100ML

N018249 001

SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER

B BRAUN

1.87GM/100ML

N020004 001 Apr 21, 1992

SODIUM LACTATE IN PLASTIC CONTAINER

+ HOSPIRA

5MEQ/ML

N018947 001 Sep 05, 1984

SODIUM MONOFLUOROPHOSPHATE

GEL; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS

1.2%

N019518 002 Aug 06, 1986

PASTE; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS

1.2%

N019518 001 Jun 03, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NIPRIDE

ROCHE	50MG/VIAL	N017546	001	
NITROPRESS				
ABBOTT	50MG/VIAL	A071555	001	Nov 16, 1987
+ ABBVIE	50MG/VIAL **	N018450	001	
HOSPIRA	50MG/VIAL	A070566	001	Jun 09, 1986
VPNA	25MG/ML	A071961	001	Aug 01, 1988

SODIUM NITROPRUSSIDE

ABRAXIS PHARM	50MG/VIAL	A070031	001	Jan 17, 1985
AKORN	25MG/ML	A208635	001	May 04, 2017
AMPHASTAR PHARMS INC	25MG/ML	A209832	001	Dec 18, 2017
AVET LIFESCIENCES	25MG/ML	A208923	001	Nov 08, 2022
+ BAXTER HLTHCARE	50MG/VIAL **	N018581	001	Jul 28, 1982
CHARTWELL RX	25MG/ML	A209834	001	Jun 26, 2018
CIPLA	25MG/ML	A210855	001	Jul 16, 2018
EUGIA PHARMA	25MG/ML	A211934	001	Dec 10, 2020
SUN PHARM	25MG/ML	A210467	001	Nov 26, 2018
TEVA PARENTERAL	25MG/ML	A073465	001	Mar 30, 1992

SOLUTION; INTRAVENOUS

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%				
+ EXELA PHARMA	10MG/50ML (0.2MG/ML) **	N209387	002	Dec 07, 2017

SODIUM OXYBATE

SOLUTION; ORAL

SODIUM OXYBATE

HIKMA	0.5GM/ML	A202090	001	Jan 17, 2017
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SODIUM PHENYLBUTYRATE

TABLET; ORAL

SODIUM PHENYLBUTYRATE

ALVOGEN	500MG	A090910	001	Nov 18, 2011
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SODIUM PHOSPHATE P-32

SOLUTION; INJECTION, ORAL

PHOSPHOTOPE

BRACCO	1-8mCi/VIAL	N010927	001	
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SODIUM PHOSPHATE P 32

MALLINCKRODT	0.67mCi/ML	N011777	001	
	1.5mCi/VIAL	N011777	002	

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

OSMOPREP

+ SALIX PHARMS	0.398GM;1.102GM	N021892	001	Mar 16, 2006
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VISICOL

SALIX PHARMS	0.398GM;1.102GM	N021097	001	Sep 21, 2000
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SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KAYEXALATE

+ CONCORDIA	453.6GM/BOT **	N011287	001	
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SODIUM POLYSTYRENE SULFONATE

CITRUSPHARMA	454GM/BOT	A040909	001	Dec 03, 2008
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+ PAI HOLDINGS PHARM	453.6GM/BOT	A088786	001	Sep 11, 1984
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SUSPENSION; ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

ANI PHARMS	15GM/60ML	A090590	001	May 13, 2011
HIKMA	15GM/60ML	A089049	001	Nov 17, 1986
MORTON GROVE	15GM/60ML	A088717	001	Sep 11, 1984
ROXANE	15GM/60ML	A088453	001	Nov 17, 1983

SODIUM SUCCINATE

INJECTABLE; INJECTION

SODIUM SUCCINATE

ELKINS SINN	30%	A080516	001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SOTRADECOL

+	ELKINS SINN	1% **	N005970 004
+		3% **	N005970 005

SODIUM THIOSULFATE

INJECTABLE; INJECTION

SODIUM THIOSULFATE

+	US ARMY	250MG/ML	N020166 001	Feb 14, 1992
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SOFOSBUVIR

TABLET; ORAL

SOFOSBUVIR

	TEVA PHARMS USA INC	400MG	A211353 001	Jan 27, 2022
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SOLIFENACIN SUCCINATE

TABLET; ORAL

SOLIFENACIN SUCCINATE

	ACCORD HLTHCARE	5MG	A207477 001	Jan 04, 2022
		10MG	A207477 002	Jan 04, 2022
	AJANTA PHARMA LTD	5MG	A205483 001	May 20, 2019
		10MG	A205483 002	May 20, 2019
	AMNEAL PHARMS CO	5MG	A209719 001	May 20, 2019
		10MG	A209719 002	May 20, 2019
	BRECKENRIDGE	5MG	A209818 001	May 20, 2019
		10MG	A209818 002	May 20, 2019
	CIPLA	5MG	A209839 001	May 20, 2019
		10MG	A209839 002	May 20, 2019
	LANNETT CO INC	5MG	A211622 001	Jun 06, 2023
		10MG	A211622 002	Jun 06, 2023
	SUNSHINE	5MG	A213346 001	Apr 13, 2020
		10MG	A213346 002	Apr 13, 2020
	ZYDUS PHARMS	5MG	A207721 001	Oct 19, 2020
		10MG	A207721 002	Oct 19, 2020

SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	3GM/100ML	N018512 001	May 27, 1982
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SOTALOL HYDROCHLORIDE

TABLET; ORAL

BETAPACE

	LEGACY PHARMA	320MG	N019865 004	Oct 30, 1992
	BETAPACE AF			
	LEGACY PHARMA	40MG	N021151 006	Apr 02, 2003
		60MG	N021151 007	Apr 02, 2003
		100MG	N021151 005	Mar 14, 2003

SOTALOL HYDROCHLORIDE

	IMPAX PHARMS	80MG	A075663 001	Nov 07, 2000
		120MG	A075663 002	Nov 07, 2000
		160MG	A075663 003	Nov 07, 2000
		240MG	A075663 004	Nov 07, 2000
	MYLAN	80MG	A075725 001	Dec 19, 2000
		120MG	A075725 002	Dec 19, 2000
		160MG	A075725 003	Dec 19, 2000
		240MG	A075725 004	Dec 19, 2000
	NATCO PHARMA USA	80MG	A075237 001	May 01, 2000
		120MG	A075237 002	May 01, 2000
		160MG	A075237 003	May 01, 2000
		240MG	A075237 004	May 01, 2000
	SUN PHARM INDUSTRIES	80MG	A075515 001	Oct 15, 2001
		80MG	A076576 001	Apr 08, 2004
		120MG	A075515 004	Oct 15, 2001
		120MG	A076576 002	Apr 08, 2004
		160MG	A075515 002	Oct 15, 2001
		160MG	A076576 003	Apr 08, 2004
		240MG	A075515 003	Oct 15, 2001
	TEVA	80MG	A076883 001	Jul 26, 2004
		120MG	A076883 002	Jul 26, 2004
		160MG	A076883 003	Jul 26, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

UPSHER SMITH LABS

80MG

A075366 001 May 01, 2000

120MG

A075366 002 May 01, 2000

160MG

A075366 003 May 01, 2000

240MG

A075366 004 May 01, 2000

WATSON LABS

80MG

A075238 001 Jul 13, 2000

120MG

A075238 002 Jul 13, 2000

160MG

A075238 003 Jul 13, 2000

240MG

A075238 004 Jul 13, 2000

SOYBEAN OIL

INJECTABLE; INJECTION

INTRALIPID 10%

+ FRESENIUS

10%

N017643 001

LIPOSYN III 10%

HOSPIRA

10%

N018969 001 Sep 24, 1984

LIPOSYN III 20%

HOSPIRA

20%

N018970 001 Sep 25, 1984

LIPOSYN III 30%

HOSPIRA

30%

N020181 001 Jan 13, 1998

SOYACAL 10%

ALPHA THERA

10%

N018465 001 Jun 29, 1983

SOYACAL 20%

ALPHA THERA

20%

N018786 001 Jun 29, 1983

TRAVAMULSION 10%

BAXTER HLTHCARE

10%

N018660 001 Feb 26, 1982

TRAVAMULSION 20%

BAXTER HLTHCARE

20%

N018758 001 Feb 15, 1983

SPARFLOXACIN

TABLET; ORAL

ZAGAM

MYLAN

200MG

N020677 001 Dec 19, 1996

SPECTINOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

TROBICIN

PFIZER

EQ 2GM BASE/VIAL

N050347 001

EQ 4GM BASE/VIAL

N050347 002

SPIRAPRIL HYDROCHLORIDE

TABLET; ORAL

RENORMAX

SCHERING

3MG

N020240 001 Dec 29, 1994

6MG

N020240 002 Dec 29, 1994

12MG

N020240 003 Dec 29, 1994

24MG

N020240 004 Dec 29, 1994

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

ACTAVIS ELIZABETH

25MG

A040353 003 Mar 15, 2006

50MG

A040353 001 Jul 29, 1999

100MG

A040353 002 Jul 29, 1999

ASCOT

25MG

A087687 001 Oct 20, 1982

CHARTWELL RX

25MG

A086809 001

IVAX PHARMS

25MG

A087108 001

LEDERLE

25MG

A087634 001

MUTUAL PHARM

25MG

A087265 001

MYLAN

25MG

A087086 001

PUREPAC PHARM

25MG

A087998 001 Oct 14, 1983

25MG

A088053 001 Aug 25, 1983

SUPERPHARM

25MG

A089364 001 Nov 07, 1986

UPSHER SMITH

25MG

A087554 001

VANGARD

25MG

A087648 001 Feb 01, 1982

WARNER CHILCOTT

25MG

A087952 001 Nov 18, 1982

WATSON LABS

25MG

A086898 002 Mar 02, 1982

25MG

A087078 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

STANOZOLOL

TABLET; ORAL

WINSTROL

+ LUNDBECK INC 2MG N012885 001 May 14, 1984

STAVUDINE

CAPSULE; ORAL

STAVUDINE

AUROBINDO PHARMA 15MG A077672 003 Dec 29, 2008

20MG A077672 004 Dec 29, 2008

30MG A077672 001 Dec 29, 2008

40MG A077672 002 Dec 29, 2008

HETERO LABS LTD III 15MG A078957 001 Dec 29, 2008

20MG A078957 002 Dec 29, 2008

30MG A078957 003 Dec 29, 2008

40MG A078957 004 Dec 29, 2008

MYLAN 15MG A079069 001 Dec 29, 2008

20MG A079069 002 Dec 29, 2008

30MG A079069 003 Dec 29, 2008

40MG A079069 004 Dec 29, 2008

MYLAN LABS LTD 30MG A078775 001 Jan 05, 2009

40MG A078775 002 Jan 05, 2009

ZERIT

BRISTOL 5MG N020412 001 Jun 24, 1994

+ 15MG ** N020412 002 Jun 24, 1994

+ 20MG ** N020412 003 Jun 24, 1994

+ 30MG ** N020412 004 Jun 24, 1994

+ 40MG ** N020412 005 Jun 24, 1994

CAPSULE, EXTENDED RELEASE; ORAL

ZERIT XR

BRISTOL MYERS SQUIBB 37.5MG N021453 001 Dec 31, 2002

50MG N021453 002 Dec 31, 2002

75MG N021453 003 Dec 31, 2002

100MG N021453 004 Dec 31, 2002

FOR SOLUTION; ORAL

STAVUDINE

AUROBINDO PHARMA 1MG/ML A077774 001 Dec 29, 2008

CIPLA LTD 1MG/ML A078030 001 Mar 20, 2009

ZERIT

+ BRISTOL 1MG/ML ** N020413 001 Sep 06, 1996

STERILE WATER FOR INJECTION

LIQUID; N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

ABRAXIS PHARM 100% A089099 001 Dec 29, 1987

100% A089100 001 Dec 29, 1987

STERILE WATER FOR INJECTION

HIKMA 100% (20ML) A206369 002 Sep 02, 2015

+ HOSPIRA 100% (1ML) N018801 001 Oct 27, 1982

+ 100% (5.2ML) N018801 006 May 02, 2023

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

B BRAUN 100% N019077 001 Mar 02, 1984

STERILE WATER FOR IRRIGATION

LIQUID; IRRIGATION

STERILE WATER IN PLASTIC CONTAINER

MILES 100% N018246 001

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

COPANOS EQ 500MG BASE/ML A060684 001

LILLY EQ 1GM BASE/VIAL A060107 001

EQ 1GM BASE/2ML A060404 001

EQ 5GM BASE/VIAL A060107 002

PFIZER EQ 1GM BASE/VIAL ** A060076 001

EQ 1GM BASE/2.5ML A060111 001

EQ 5GM BASE/VIAL ** A060076 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

ANECTINE

SANDOZ	50MG/ML	N008453	003	
	500MG/VIAL	N008453	001	
	1GM/VIAL	N008453	004	
QUELICIN PRESERVATIVE FREE				
+ HOSPIRA	20MG/ML **	N008845	001	
+	50MG/ML **	N008845	002	
+	100MG/ML **	N008845	004	
SUCCINYLCHOLINE CHLORIDE				
AMPHASTAR PHARMS INC	20MG/ML	A213432	001	Jun 08, 2020
BRECKENRIDGE	20MG/ML	A212638	001	Oct 09, 2019
INTL MEDICATION	100MG/VIAL	A085400	001	Feb 04, 1982
MANKIND PHARMA	20MG/ML	A216127	001	Feb 02, 2023
ORGANON USA INC	20MG/ML	A080997	001	
SUCOSTRIN				
+ APOTHECON	20MG/ML	N008847	001	
+	100MG/ML	N008847	003	

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTANIL CITRATE

WATSON LABS	EQ 0.05MG BASE/ML	A074406	001	Dec 15, 1995
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SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

BLEPH-10

ALLERGAN	10%	A084015	001	
CETAMIDE				
ALCON	10%	A080021	001	
SODIUM SULAMYD				
+ SCHERING	10% **	N005963	002	
SULFAIR 10				
PHARMAFAIR	10%	A088000	001	Dec 22, 1982
SOLUTION/DROPS; OPHTHALMIC				
BLEPH-10				
+ ALLERGAN	10%	A080028	001	
BLEPH-30				
ALLERGAN	30%	A080028	002	
ISOPTO CETAMIDE				
ALCON	15%	A080020	002	
OCUSULF-10				
MIZA PHARMS USA	10%	A080660	001	
OCUSULF-30				
MIZA PHARMS USA	30%	A080660	002	
SODIUM SULAMYD				
+ SCHERING	10% **	N005963	001	
+	30% **	N005963	003	
SODIUM SULFACETAMIDE				
AKORN	10%	A083021	001	
	15%	A083021	002	
	30%	A083021	003	
SOLA BARNES HIND	10%	A084143	001	
	10%	A084145	001	
	30%	A084146	001	
	30%	A084147	001	
SULF-10				
NOVARTIS	10%	A080025	001	
SULF-15				
NOVARTIS	15%	A089047	001	Oct 31, 1995
SULFACEL-15				
OPTOPICS	15%	A080024	001	
SULFACETAMIDE SODIUM				
AKORN	10%	A040215	001	May 25, 1999
ALCON PHARMS LTD	30%	A089068	001	May 05, 1987
EPIC PHARMA LLC	30%	A040216	001	May 25, 1999
PHARMAFAIR	10%	A088947	001	May 17, 1985
SULFAIR 10				
PHARMAFAIR	10%	A087949	001	Dec 13, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFAIR FORTE

PHARMAFAIR 30% A088385 001 Oct 13, 1983

SULFAIR-15

PHARMAFAIR 15% A088186 001 May 25, 1983

SULTEN-10

BAUSCH AND LOMB 10% A087818 001 Feb 03, 1983

SULFACYTINE

TABLET;ORAL

RENOQUID

GLENWOOD 250MG N017569 001

SULFADIAZINE

TABLET;ORAL

SULFADIAZINE

ABBVIE 300MG N004125 005

CHARTWELL MOLECULAR 500MG A080084 001

EVERYLIFE 500MG A080088 001

+ IMPAX LABS 500MG A080081 001

LEDERLE 500MG N004054 001

+ LILLY 500MG N004122 002

SULFADIAZINE SODIUM

INJECTABLE;INJECTION

SULFADIAZINE SODIUM

LEDERLE 250MG/ML N004054 002

SULFADIAZINE; SULFAMERAZINE

SUSPENSION;ORAL

SULFONAMIDES DUPLEX

LILLY 250MG/5ML;250MG/5ML N006317 007

SULFAMETER

TABLET;ORAL

SULLA

BAYER HLTHCARE 500MG N016000 002

SULFAMETHIZOLE

TABLET;ORAL

MICROSUL

FOREST PHARMS 1GM A086012 001

PROKLAR

FOREST PHARMS 500MG A080273 001

THIOSULFIL

WYETH AYERST 250MG N008565 001

500MG N008565 004

SULFAMETHOXAZOLE

SUSPENSION;ORAL

GANTANOL

ROCHE 500MG/5ML N013664 002

TABLET;ORAL

GANTANOL

ROCHE 500MG N012715 002

GANTANOL-DS

ROCHE 1GM N012715 003

SULFAMETHOXAZOLE

ASCOT 500MG A087662 001 Oct 20, 1982

BARR 500MG A087189 001 Jul 25, 1983

HEATHER 500MG A086163 001

RISING 500MG A085844 001

WATSON LABS 500MG A085053 001

1GM A086000 001

UROBAK

SHIONOGI 500MG A087307 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

BACTRIM

+ SUN PHARM INDS INC 80MG/ML;16MG/ML ** N018374 001

SEPTRA

MONARCH PHARMS 80MG/ML;16MG/ML ** N018452 001

SULFAMETHOXAZOLE AND TRIMETHOPRIM

ABRAXIS PHARM 80MG/ML;16MG/ML A070223 001 Dec 29, 1987

BEDFORD 80MG/ML;16MG/ML A072383 001 Apr 29, 1992

HIKMA 80MG/ML;16MG/ML A070627 001 Dec 29, 1987

80MG/ML;16MG/ML A070628 001 Dec 29, 1987

HOSPIRA 80MG/ML;16MG/ML A073199 001 Sep 11, 1992

WATSON LABS 80MG/ML;16MG/ML A071556 001 Dec 29, 1987

SUSPENSION; ORAL

BACTRIM

+ SUN PHARM INDUSTRIES 200MG/5ML;40MG/5ML ** N017560 001

BACTRIM PEDIATRIC

SUN PHARM INDUSTRIES 200MG/5ML;40MG/5ML ** N017560 002

SEPTRA

MONARCH PHARMS 200MG/5ML;40MG/5ML ** N017598 001

SEPTRA GRAPE

MONARCH PHARMS 200MG/5ML;40MG/5ML ** N017598 002 Feb 12, 1986

SULFAMETHOXAZOLE AND TRIMETHOPRIM

ANI PHARMS 200MG/5ML;40MG/5ML A070028 001 Jun 02, 1987

LUPIN LTD 200MG/5ML;40MG/5ML A212699 001 Jan 05, 2021

TEVA 200MG/5ML;40MG/5ML N018812 001 Jan 28, 1983

200MG/5ML;40MG/5ML N018812 002 Jun 10, 1983

SULFATRIM

PHARM ASSOC 200MG/5ML;40MG/5ML N018615 002 Jan 07, 1983

SULMEPRIM

USL PHARMA 200MG/5ML;40MG/5ML A070063 001 Aug 01, 1986

SULMEPRIM PEDIATRIC

USL PHARMA 200MG/5ML;40MG/5ML A070064 001 Aug 01, 1986

TRIMETH/SULFA

ALPHARMA US PHARMS 200MG/5ML;40MG/5ML A072289 001 May 23, 1988

200MG/5ML;40MG/5ML A072398 001 May 23, 1988

NASKA 200MG/5ML;40MG/5ML A072399 001 May 23, 1988

TABLET; ORAL

COTRIM

TEVA 400MG;80MG A070034 001 May 16, 1985

COTRIM D.S.

TEVA 800MG;160MG A070048 001 Mar 18, 1985

SULFAMETHOPRIM

NOVEL LABS INC 400MG;80MG A070022 001 Feb 15, 1985

SULFAMETHOPRIM-DS

NOVEL LABS INC 800MG;160MG A070032 001 Feb 15, 1985

SULFAMETHOXAZOLE AND TRIMETHOPRIM

FOSUN PHARMA 400MG;80MG A070889 001 Nov 13, 1986

400MG;80MG N018598 003 May 19, 1982

800MG;160MG A070890 001 Nov 13, 1986

HEATHER 400MG;80MG N018946 001 Aug 10, 1984

800MG;160MG N018946 002 Aug 10, 1984

INTERPHARM 400MG;80MG A071299 001 Oct 27, 1987

800MG;160MG A071300 001 Oct 27, 1987

MARTEC USA LLC 400MG;80MG A072408 001 Dec 07, 1988

MUTUAL PHARM 400MG;80MG A070006 001 Nov 14, 1984

PLIVA 400MG;80MG A070215 001 Sep 10, 1985

800MG;160MG A070216 001 Sep 10, 1985

ROXANE 400MG;80MG A072768 001 Aug 30, 1991

TEVA 400MG;80MG N018242 001

800MG;160MG N018242 002

USL PHARMA 400MG;80MG A070203 001 Nov 08, 1985

800MG;160MG A070204 001 Nov 08, 1985

WATSON LABS 400MG;80MG A070002 001 Nov 07, 1984

400MG;80MG N018852 001 May 09, 1983

800MG;160MG A070000 001 Nov 07, 1984

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

FOSUN PHARMA 800MG;160MG N018598 004 May 19, 1982

HERITAGE PHARMA AVET 800MG;160MG A070037 001 Jun 02, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

MARTEC USA LLC	800MG;160MG	A072417	001	Dec 07, 1988
MUTUAL PHARM	800MG;160MG	A070007	001	Nov 14, 1984
ROXANE	800MG;160MG	A072769	001	Aug 30, 1991
WATSON LABS	800MG;160MG	N018854	001	May 09, 1983

SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH

HERITAGE PHARMA AVET	400MG;80MG	A070030	001	Jun 02, 1987
SULFATRIM-DS				
SUPERPHARM	800MG;160MG	A070066	001	Jun 24, 1985
SULFATRIM-SS				
SUPERPHARM	400MG;80MG	A070065	002	Jun 24, 1985
UROPLUS DS				
SHIONOGI	800MG;160MG	A071816	001	Sep 28, 1987
UROPLUS SS				
SHIONOGI	400MG;80MG	A071815	001	Sep 28, 1987

SULFANILAMIDE

CREAM; VAGINAL

AVC					
+	MYLAN SPECIALITY LP	15% **	N006530	003	Jan 27, 1987
SULFANILAMIDE					
	COSETTE	15%	A088718	001	Sep 19, 1985

SUPPOSITORY; VAGINAL

AVC					
	MYLAN SPECIALITY LP	1.05GM	N006530	004	Jan 27, 1987

SULFAPHENAZOLE

SUSPENSION; ORAL

SULFABID					
	PHARM RES ASSOC	500MG/5ML	N013093	001	

TABLET; ORAL

SULFABID					
	PURDUE FREDERICK	500MG	N013092	002	

SULFAPYRIDINE

TABLET; ORAL

SULFAPYRIDINE					
	LILLY	500MG	N000159	001	

SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE					
	PHARMACIA AND UPJOHN	250MG/5ML	N018605	001	

TABLET; ORAL

S.A.S.-500					
	SOLVAY	500MG	A083450	001	
SULFASALAZINE					
	EPIC PHARMA LLC	500MG	A086184	001	
	SUN PHARM INDUSTRIES	500MG	A089590	001	Oct 19, 1987
	SUPERPHARM	500MG	A089339	001	Oct 26, 1987
	WATSON LABS	500MG	A084964	001	
		500MG	A087197	001	

TABLET, DELAYED RELEASE; ORAL

SULFASALAZINE					
	WATSON LABS	500MG	A088052	001	May 24, 1983

SULFINPYRAZONE

CAPSULE; ORAL

ANTURANE					
+	NOVARTIS	200MG **	N011556	004	
SULFINPYRAZONE					
	BARR	200MG	A087666	001	Sep 17, 1982
	IVAX PHARMS	200MG	A087770	001	Nov 19, 1982
	PAR PHARM	200MG	A088934	001	Sep 06, 1985
	VANGARD	200MG	A088666	001	Feb 17, 1984

TABLET; ORAL

ANTURANE					
	NOVARTIS	100MG **	N011556	003	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFINPYRAZONE

TABLET; ORAL

SULFINPYRAZONE

BARR	100MG	A087665 001	Sep 17, 1982
IVAX PHARMS	100MG	A087769 001	Jun 01, 1982
PAR PHARM	100MG	A088933 001	Sep 06, 1985
WATSON LABS	100MG	A087667 001	May 26, 1982

SULFISOXAZOLE

TABLET; ORAL

GANTRISIN

ROCHE	500MG	N006525 001	
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SOSOL

MK LABS	500MG	A080036 001	
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SOXAZOLE

ALRA	500MG	A080366 001	
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SULFALAR

PARKE DAVIS	500MG	A084955 001	
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SULFISOXAZOLE

ANI PHARMS	500MG	A080142 001	
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BARR	500MG	A084031 001	
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HEATHER	500MG	A080189 001	
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IMPAX LABS	500MG	A080109 001	
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LANNETT	500MG	A080085 001	
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LEDERLE	500MG	A087649 001	
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PHARMERAL	500MG	A084385 001	
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PUREPAC PHARM	500MG	A080087 001	
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RISING	500MG	A085628 001	
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ROXANE	500MG	A080082 001	
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VALEANT PHARM INTL	500MG	A080268 002	
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VITARINE	500MG	A087332 001	
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WATSON LABS	500MG	A085534 001	
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WEST WARD	500MG	A080379 001	
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SULSOXIN

SOLVAY	500MG	A080040 001	
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SULFISOXAZOLE ACETYL

EMULSION; ORAL

LIPO GANTRISIN

ROCHE	EQ 1GM BASE/5ML	N009182 009	
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SUSPENSION; ORAL

GANTRISIN PEDIATRIC

ROCHE	EQ 500MG BASE/5ML	N009182 004	
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SYRUP; ORAL

GANTRISIN

ROCHE	EQ 500MG BASE/5ML	N009182 002	
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SULFISOXAZOLE DIOLAMINE

INJECTABLE; INJECTION

GANTRISIN

ROCHE	EQ 400MG BASE/ML	N006917 001	
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OINTMENT; OPHTHALMIC

GANTRISIN

ROCHE	EQ 4% BASE	N008414 002	
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SOLUTION/DROPS; OPHTHALMIC

GANTRISIN

ROCHE	EQ 4% BASE	N007757 002	
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SULFISOXAZOLE DIOLAMINE

SOLA BARNES HIND	EQ 4% BASE	A084148 001	
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SULFOXONE SODIUM

TABLET, DELAYED RELEASE; ORAL

DIASONE SODIUM

ABBVIE	165MG	N006044 003	
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SULFUR

POWDER; TOPICAL

BENSULFOID

POYTHRESS	33.32%	N002918 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULINDAC

TABLET; ORAL

CLINORIL

+	MERCK	150MG **	N017911	001	
+		200MG **	N017911	002	

SULINDAC

ANI PHARMS	150MG	A072973	002	Feb 28, 1992
	200MG	A072973	001	Feb 28, 1992
CHARTWELL RX	150MG	A072712	001	Aug 30, 1991
	200MG	A072713	001	Aug 30, 1991
EPIC PHARMA LLC	150MG	A073262	002	Sep 06, 1991
	200MG	A073262	001	Sep 06, 1991
RISING	150MG	A073039	002	Jun 22, 1993
	200MG	A073039	001	Jun 22, 1993

SUMATRIPTAN

SPRAY; NASAL

IMITREX

GLAXOSMITHKLINE	10MG/SPRAY	N020626	002	Aug 26, 1997
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SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

ALSUMA

MERIDIAN MEDCL	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N022377	001	Jun 29, 2010
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SUMATRIPTAN SUCCINATE

ANTARES PHARMA INC	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	A078319	001	Dec 10, 2015
	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A078319	002	Dec 10, 2015
BAXTER HLTHCARE CORP	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A207101	001	Jan 19, 2023
FRESENIUS KABI USA	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A079240	002	Sep 18, 2009
	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	A079240	001	Sep 18, 2009
MYLAN LABS LTD	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A203322	001	Apr 14, 2014
PAR STERILE PRODUCTS	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A077871	001	Jul 09, 2009
SANDOZ	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	A078067	002	Feb 06, 2009
	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A078067	001	Feb 06, 2009
STERISCIENCE	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090314	001	Jun 10, 2010
	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090641	001	Jul 28, 2010
TEVA PARENTERAL	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	A078318	001	Feb 06, 2009
	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A078318	002	Feb 06, 2009
TEVA PHARMS USA	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A077907	001	Feb 06, 2009
ZYDUS	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	A090310	001	Aug 11, 2010

SUMAVEL DOSEPRO

+	ENDO VENTURES LTD	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	N022239	002	Nov 26, 2013
+		EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N022239	001	Jul 15, 2009

SYSTEM; IONTOPHORESIS

ZECUITY

+	TEVA BRANDED PHARM	EQ 6.5MG BASE/4HR	N202278	001	Jan 17, 2013
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TABLET; ORAL

SUMATRIPTAN SUCCINATE

FOSUN PHARMA	EQ 25MG BASE	A076976	001	Aug 10, 2009
	EQ 50MG BASE	A076976	002	Aug 10, 2009
	EQ 100MG BASE	A076976	003	Aug 10, 2009
HIKMA PHARMS	EQ 25MG BASE	A078298	001	May 21, 2013
	EQ 50MG BASE	A078298	002	May 21, 2013
	EQ 100MG BASE	A078298	003	May 21, 2013
MYLAN	EQ 25MG BASE	A077163	001	Nov 02, 2009
	EQ 50MG BASE	A077163	002	Nov 02, 2009
	EQ 100MG BASE	A077163	003	Nov 02, 2009
ROXANE	EQ 25MG BASE	A078241	001	Aug 10, 2009
	EQ 50MG BASE	A078241	002	Aug 10, 2009
	EQ 100MG BASE	A078241	003	Aug 10, 2009
SUN PHARM INDS LTD	EQ 25MG BASE	A076554	001	Aug 10, 2009
	EQ 50MG BASE	A076554	002	Aug 10, 2009
	EQ 100MG BASE	A076572	001	Feb 09, 2009
TEVA	EQ 25MG BASE	A076840	001	Feb 09, 2009
	EQ 50MG BASE	A076840	002	Feb 09, 2009
	EQ 100MG BASE	A076840	003	Feb 09, 2009

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SUPROFEN

SOLUTION/DROPS;OPHTHALMIC

PROFENAL

ALCON

1%

N019387 001 Dec 23, 1988

TACRINE HYDROCHLORIDE

CAPSULE;ORAL

COGNEX

SHIONOGI INC

EQ 10MG BASE

N020070 001 Sep 09, 1993

EQ 20MG BASE

N020070 002 Sep 09, 1993

EQ 30MG BASE

N020070 003 Sep 09, 1993

EQ 40MG BASE

N020070 004 Sep 09, 1993

TACROLIMUS

CAPSULE;ORAL

TACROLIMUS

CONCORD BIOTECH LTD

EQ 0.5MG BASE

A213112 001 Nov 10, 2020

HERITAGE PHARMA AVET

EQ 5MG BASE

A090402 001 Jul 01, 2010

INJECTABLE;INJECTION

TACROLIMUS

HOSPIRA

EQ 5MG BASE/ML

A203900 001 Aug 25, 2017

TADALAFIL

TABLET;ORAL

TADALAFIL

CHARTWELL RX

2.5MG

A210716 001 Dec 29, 2020

5MG

A210716 002 Dec 29, 2020

10MG

A210716 003 Dec 29, 2020

20MG

A210716 004 Dec 29, 2020

MYLAN

5MG

A206957 001 Apr 29, 2019

20MG

A200630 001 Aug 03, 2018

RISING

2.5MG

A206956 001 Apr 29, 2019

10MG

A206956 002 Apr 29, 2019

20MG

A206956 003 Apr 29, 2019

ZYDUS PHARMS

20MG

A212515 001 Jun 06, 2023

TALBUTAL

TABLET;ORAL

LOTUSATE

SANOFI AVENTIS US

120MG

N009410 005

TAMOXIFEN CITRATE

TABLET;ORAL

NOLVADEX

+ ASTRAZENECA

EQ 10MG BASE **

N017970 001

+

EQ 20MG BASE **

N017970 002 Mar 21, 1994

TAMOXIFEN CITRATE

ACTAVIS LABS FL INC

EQ 10MG BASE

A076179 001 Feb 20, 2003

EQ 20MG BASE

A076179 002 Feb 20, 2003

AEGIS PHARMS

EQ 10MG BASE

A076398 001 Mar 31, 2003

EQ 20MG BASE

A076398 002 Mar 31, 2003

IVAX SUB TEVA PHARMS

EQ 10MG BASE

A075740 001 Feb 20, 2003

EQ 20MG BASE

A075740 002 Feb 20, 2003

PHARMACHEMIE

EQ 10MG BASE

A074539 001 Mar 31, 2003

ROXANE

EQ 10MG BASE

A076027 001 Feb 20, 2003

EQ 20MG BASE

A076027 002 Feb 20, 2003

TEVA

EQ 10MG BASE

A074504 001 Apr 28, 2003

EQ 20MG BASE

A074504 002 Apr 28, 2003

TAMSULOSIN HYDROCHLORIDE

CAPSULE;ORAL

TAMSULOSIN HYDROCHLORIDE

ANCHEN PHARMS

0.4MG

A202010 001 Jan 04, 2013

ASCENT PHARMS INC

0.4MG

A214730 001 May 04, 2022

MYLAN

0.4MG

A090408 001 Apr 27, 2010

TAPENTADOL HYDROCHLORIDE

SOLUTION;ORAL

NUCYNTA

+ COLLEGIUM PHARM INC

EQ 20MG BASE/ML

N203794 001 Oct 15, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TAZAROTENE

CREAM; TOPICAL

TAZAROTENE

FOUGERA PHARMS INC 0.1%

A211175 001 Jan 28, 2019

TECHNETIUM TC-99M APCITIDE

INJECTABLE; INJECTION

ACUTECT

CIS BIO INTL SA N/A

N020887 001 Sep 14, 1998

TECHNETIUM TC-99M DEPREOTIDE

INJECTABLE; INJECTION

NEO TECT KIT

CIS BIO INTL SA N/A **

N021012 001 Aug 03, 1999

TECHNETIUM TC-99M DISOFENIN KIT

INJECTABLE; INJECTION

HEPATOLITE

SUN PHARM INDS INC N/A

N018467 001 Mar 16, 1982

TECHNETIUM TC-99M ETIDRONATE KIT

INJECTABLE; INJECTION

CINTICHEM TECHNETIUM 99M HEDSPA

GE HEALTHCARE N/A

N017653 001

MPI STANNOUS DIPHOSPHONATE

GE HEALTHCARE N/A

N017667 001

OSTEOSCAN

MALLINCKRODT N/A

N017454 001

TECHNETIUM TC 99M DIPHOSPHONATE-TIN KIT

GE HEALTHCARE N/A

N017562 001

TECHNETIUM TC-99M FERPENTETATE KIT

INJECTABLE; INJECTION

RENOTEC

BRACCO N/A

N017045 001

TECHNETIUM TC-99M GLUCEPTATE KIT

INJECTABLE; INJECTION

GLUCOSCAN

BRISTOL MYERS SQUIBB N/A

N017907 001

TECHNESCAN GLUCEPTATE

DRAXIMAGE N/A

N018272 001 Jan 27, 1982

TECHNETIUM TC-99M LIDOFENIN KIT

INJECTABLE; INJECTION

TECHNESCAN HIDA

DRAXIMAGE N/A

N018489 001 Oct 31, 1986

TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-10

JUBILANT N/A

N018035 001

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

AMERSCAN MDP KIT

GE HEALTHCARE N/A

N018335 001 Aug 05, 1982

OSTEOLITE

PHARMALUCENCE N/A

N017972 001

TECHNETIUM TC 99M MPI MDP

GE HEALTHCARE N/A

N018141 001

N/A

N018141 002 Jun 12, 1989

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

AN-DTPA

JUBILANT DRAXIMAGE N/A

N017714 001

MPI DTPA KIT - CHELATE

GE HEALTHCARE N/A

N017255 001

TECHNETIUM TC-99M PENTETATE KIT

GE HEALTHCARE N/A

N017264 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M POLYPHOSPHATE KIT

INJECTABLE;INJECTION

SODIUM POLYPHOSPHATE-TIN KIT

GE HEALTHCARE

N/A

N017664 001

TECHNETIUM TC-99M PYRO/TRIMETA PHOSPHATES KIT

INJECTABLE;INJECTION

PYROLITE

PHARMALUCENCE

N/A

N017684 001

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE;INJECTION

PHOSPHOTEC

BRACCO

N/A

N017680 001

TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE;INJECTION

RBC-SCAN

CADEMA

N/A

N020063 001 Jun 11, 1992

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE;INJECTION

MIRALUMA

LANTHEUS MEDCL

N/A

N019785 003 May 23, 1997

TECHNETIUM TC-99M SODIUM PERTECHNETATE

SOLUTION;INJECTION, ORAL

SODIUM PERTECHNETATE TC 99M

+ GE HEALTHCARE

2-100mCi/ML **

N017471 001

+ MALLINCKRODT

10-60mCi/ML **

N017725 001

PHARMALUCENCE

12mCi/ML

N017321 001

24mCi/ML

N017321 002

48mCi/ML

N017321 003

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION;INJECTION, ORAL

MINITEC

BRACCO

0.22-2.22 CI/GENERATOR

N017339 001

SOLUTION;INTRAVENOUS

TECHNELITE

LANTHEUS MEDCL

0.0083-2.7 CI/GENERATOR

N017771 001

ULTRA-TECHNEKOW FM

CURIUM

0.25-3 CI/GENERATOR

N017243 002

SOLUTION;INTRAVENOUS, INTRAVESICULAR, OPHTHALMIC

RADIOGENIX SYSTEM

+ NORTHSTAR MEDICAL

30-1153mCi/GENERATOR

N202158 001 Feb 08, 2018

SOLUTION;INTRAVENOUS, ORAL

TECHNETIUM TC 99M GENERATOR

+ GE HEALTHCARE

68-2703mCi/GENERATOR

N017693 002 Dec 13, 2013

830-16600mCi/GENERATOR

N017693 001

TECHNETIUM TC-99M SUCCIMER KIT

INJECTABLE;INJECTION

MPI DMSA KIDNEY REAGENT

+ GE HEALTHCARE

N/A **

N017944 001 May 18, 1982

TECHNETIUM TC-99M SULFUR COLLOID

SOLUTION;INJECTION, ORAL

TECHNETIUM TC 99M SULFUR COLLOID

GE HEALTHCARE

4mCi/ML

N017456 001

SOLUTION;ORAL

TECHNETIUM TC 99M SULFUR COLLOID

MALLINCKRODT

3mCi/ML

N017724 001

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION;INJECTION, ORAL

TECHNECOLL

MALLINCKRODT

N/A

N017059 001

TECHNETIUM TC 99M TSC

GE HEALTHCARE

N/A

N017784 001

TESULOID

BRACCO

N/A

N016923 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M TEBOROXIME KIT

INJECTABLE; INJECTION

CARDIOTEC

BRACCO

N/A

N019928 001 Dec 19, 1990

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

+ ALFASIGMA

EQ 2MG BASE

N021200 001 Jul 24, 2002

+

EQ 6MG BASE

N021200 002 Jul 24, 2002

TELAPREVIR

TABLET; ORAL

INCIVEK

VERTEX PHARMS

375MG

N201917 001 May 23, 2011

TELAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

VIBATIV

+ CUMBERLAND

EQ 250MG BASE/VIAL

N022110 001 Sep 11, 2009

TELBIVUDINE

SOLUTION; ORAL

TYZEKA

NOVARTIS

100MG/5ML

N022154 001 Apr 28, 2009

TABLET; ORAL

TYZEKA

+ NOVARTIS

600MG

N022011 001 Oct 25, 2006

TELITHROMYCIN

TABLET; ORAL

KETEK

SANOFI AVENTIS US

300MG

N021144 002 Feb 09, 2005

400MG

N021144 001 Apr 01, 2004

TELMISARTAN

TABLET; ORAL

TELMISARTAN

HISUN PHARM HANGZHOU

20MG

A207843 001 Feb 19, 2019

40MG

A207843 002 Feb 19, 2019

80MG

A207843 003 Feb 19, 2019

JUBILANT GENERICS

20MG

A204164 001 Aug 22, 2016

40MG

A204164 002 Aug 22, 2016

80MG

A204164 003 Aug 22, 2016

TORRENT

20MG

A203171 001 Jul 07, 2014

40MG

A203171 002 Jul 07, 2014

80MG

A203171 003 Jul 07, 2014

TEMAZEPAM

CAPSULE; ORAL

TEMAZ

QUANTUM PHARMICS

15MG

A070564 001 Oct 15, 1985

30MG

A070547 001 Oct 15, 1985

TEMAZEPAM

AUROBINDO PHARMA USA

7.5MG

A070920 002 May 21, 2010

15MG

A070920 004 Jul 07, 1986

22.5MG

A070920 003 Jun 12, 2009

30MG

A070920 001 Jul 10, 1986

DURAMED PHARMS BARR

15MG

A071708 001 Sep 29, 1988

30MG

A071709 001 Sep 29, 1988

SUN PHARM INDUSTRIES

15MG

A071174 001 Jul 10, 1986

22.5MG

A071175 002 Sep 14, 2009

30MG

A071175 001 Jul 10, 1986

USL PHARMA

15MG

A070489 001 Jul 07, 1986

30MG

A070490 001 Jul 07, 1986

WATSON LABS

15MG

A070383 001 Mar 23, 1987

15MG

A071446 001 May 21, 1993

30MG

A070384 001 Mar 23, 1987

30MG

A071447 001 May 21, 1993

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TEMOZOLOMIDE

CAPSULE; ORAL

TEMODAR

+	MERCK SHARP DOHME	5MG	N021029	001	Aug 11, 1999
+		20MG	N021029	002	Aug 11, 1999
+		100MG	N021029	003	Aug 11, 1999
+		140MG	N021029	005	Oct 19, 2006
+		180MG	N021029	006	Oct 19, 2006
+		250MG	N021029	004	Aug 11, 1999

TEMOZOLOMIDE

ANI PHARMS

		5MG	A203490	001	Jul 13, 2016
		20MG	A203490	002	Jul 13, 2016
		100MG	A203490	003	Jul 13, 2016
		140MG	A203490	004	Jul 13, 2016
		180MG	A203490	005	Jul 13, 2016
		250MG	A203490	006	Jul 13, 2016

APOTEX

		5MG	A204159	001	Jul 05, 2018
		20MG	A204159	002	Jul 05, 2018
		100MG	A204159	003	Jul 05, 2018
		140MG	A204159	004	Jul 05, 2018
		180MG	A204159	005	Jul 05, 2018
		250MG	A204159	006	Jul 05, 2018

CHARTWELL MOLECULAR

		5MG	A203898	001	Feb 10, 2016
		20MG	A203898	002	Feb 10, 2016
		100MG	A203898	003	Feb 10, 2016
		140MG	A203898	004	Feb 10, 2016
		180MG	A203898	005	Feb 10, 2016
		250MG	A203898	006	Feb 10, 2016

HERITAGE

		5MG	A078879	001	Mar 01, 2010
		20MG	A078879	002	Mar 01, 2010
		100MG	A078879	003	Mar 01, 2010
		140MG	A078879	005	Mar 01, 2010
		180MG	A078879	006	Mar 01, 2010
		250MG	A078879	004	Mar 01, 2010

MYLAN

		5MG	A205227	001	Jun 29, 2016
		20MG	A205227	002	Jun 29, 2016
		100MG	A205227	003	Jun 29, 2016
		140MG	A205227	004	Jun 29, 2016
		180MG	A205227	005	Jun 29, 2016
		250MG	A205227	006	Jun 29, 2016

WATSON LABS TEVA

		5MG	A203959	001	Apr 18, 2017
		20MG	A203959	002	Apr 18, 2017
		100MG	A203959	003	Apr 18, 2017
		140MG	A203959	004	Apr 18, 2017
		250MG	A203959	005	Apr 18, 2017

TENIPOSIDE

INJECTABLE; INJECTION

VUMON

+	HQ SPECLT PHARMA	10MG/ML	N020119	001	Jul 14, 1992
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TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

TENOFOVIR DISOPROXIL FUMARATE

CASI PHARMS INC

		300MG	A209550	001	Feb 26, 2018
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MYLAN

		150MG	A206569	001	Nov 27, 2018
		200MG	A206569	002	Nov 27, 2018
		250MG	A206569	003	Nov 27, 2018
		300MG	A206569	004	Nov 27, 2018

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

HYTRIN

+	ABBOTT	EQ 1MG BASE **	N020347	001	Dec 14, 1994
+		EQ 2MG BASE **	N020347	002	Dec 14, 1994
+		EQ 5MG BASE **	N020347	003	Dec 14, 1994
+		EQ 10MG BASE **	N020347	004	Dec 14, 1994

TERAZOSIN HYDROCHLORIDE

BIONPHARMA

		EQ 1MG BASE	A075667	001	Jul 28, 2000
		EQ 2MG BASE	A075667	002	Jul 28, 2000
		EQ 5MG BASE	A075667	003	Jul 28, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

TERAZOSIN HYDROCHLORIDE

	EQ 10MG BASE	A075667 004	Jul 28, 2000
HIKMA	EQ 1MG BASE	A075498 001	Apr 12, 2001
	EQ 2MG BASE	A075498 002	Apr 12, 2001
	EQ 5MG BASE	A075498 003	Apr 12, 2001
	EQ 10MG BASE	A075498 004	Apr 12, 2001
MYLAN	EQ 1MG BASE	A075140 002	Feb 11, 2000
	EQ 2MG BASE	A075140 003	Feb 11, 2000
	EQ 5MG BASE	A075140 001	Feb 11, 2000
	EQ 10MG BASE	A075140 004	Feb 11, 2000
MYLAN TECHNOLOGIES	EQ 1MG BASE	A075384 001	Dec 01, 2000
	EQ 2MG BASE	A075384 002	Dec 01, 2000
	EQ 5MG BASE	A075384 003	Dec 01, 2000
	EQ 10MG BASE	A075384 004	Dec 01, 2000
RANBAXY LABS LTD	EQ 1MG BASE	A076021 001	Aug 22, 2002
	EQ 2MG BASE	A076021 002	Aug 22, 2002
	EQ 5MG BASE	A076021 003	Aug 22, 2002
	EQ 10MG BASE	A076021 004	Aug 22, 2002

TABLET;ORAL

HYTRIN

ABBOTT	EQ 1MG BASE	N019057 001	Aug 07, 1987
	EQ 2MG BASE	N019057 002	Aug 07, 1987
	EQ 5MG BASE	N019057 003	Aug 07, 1987
	EQ 10MG BASE	N019057 004	Aug 07, 1987

TERAZOSIN HYDROCHLORIDE

CHARTWELL RX

	EQ 1MG BASE	A074657 001	Apr 28, 2000
	EQ 2MG BASE	A074657 002	Apr 28, 2000
	EQ 5MG BASE	A074657 003	Apr 28, 2000
	EQ 10MG BASE	A074657 004	Apr 28, 2000

IVAX SUB TEVA PHARMS

	EQ 1MG BASE	A074530 001	Apr 21, 2000
	EQ 2MG BASE	A074530 002	Apr 21, 2000
	EQ 5MG BASE	A074530 003	Apr 21, 2000
	EQ 10MG BASE	A074530 004	Apr 21, 2000

SANDOZ

	EQ 1MG BASE	A074315 001	Dec 31, 1998
	EQ 2MG BASE	A074315 002	Dec 31, 1998
	EQ 5MG BASE	A074315 003	Dec 31, 1998
	EQ 10MG BASE	A074315 004	Dec 31, 1998

TEVA

	EQ 1MG BASE	A074446 001	May 18, 2000
	EQ 2MG BASE	A074446 002	May 18, 2000
	EQ 5MG BASE	A074446 003	May 18, 2000
	EQ 10MG BASE	A074446 004	May 18, 2000

TERBINAFINE

GEL;TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS	1%	N020846 001	Apr 29, 1998
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TERBINAFINE HYDROCHLORIDE

CREAM;TOPICAL

LAMISIL

NOVARTIS	1% **	N020192 001	Dec 30, 1992
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GRANULE;ORAL

LAMISIL

+ NOVARTIS	EQ 125MG BASE/PACKET	N022071 001	Sep 28, 2007
+	EQ 187.5MG BASE/PACKET	N022071 002	Sep 28, 2007

SOLUTION;TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS	1%	N020749 001	Oct 17, 1997
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TABLET;ORAL

LAMISIL

+ NOVARTIS	EQ 250MG BASE **	N020539 001	May 10, 1996
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TERBINAFINE HYDROCHLORIDE

GEDEON RICHTER USA	EQ 250MG BASE	A077065 001	Jul 02, 2007
HERITAGE PHARMA AVET	EQ 250MG BASE	A076377 001	Jul 02, 2007
MYLAN	EQ 250MG BASE	A077136 001	Jul 02, 2007
	EQ 250MG BASE	A077195 001	Jul 02, 2007
ROXANE	EQ 250MG BASE	A077223 001	Jul 02, 2007
WOCKHARDT	EQ 250MG BASE	A078229 001	Jul 02, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERBUTALINE SULFATE

AEROSOL, METERED; INHALATION

BRETHAIRE

NOVARTIS 0.2MG/INH N018762 001 Aug 17, 1984

BRICANYL

SANOFI AVENTIS US 0.2MG/INH N018000 001 Mar 19, 1985

INJECTABLE; INJECTION

BRETHINE

+ PHARMACARE 1MG/ML ** N018571 001

BRICANYL

SANOFI AVENTIS US 1MG/ML N017466 001

TERBUTALINE SULFATE

AKORN 1MG/ML A078151 001 Jan 07, 2008

DR REDDYS 1MG/ML A076853 001 Jul 20, 2004

TABLET; ORAL

BRICANYL

SANOFI AVENTIS US 2.5MG N017618 001

5MG N017618 002

TERCONAZOLE

CREAM; VAGINAL

TERAZOL 3

+ JANSSEN PHARMS 0.8% ** N019964 001 Feb 21, 1991

TERAZOL 7

+ JANSSEN PHARMS 0.4% ** N019579 001 Dec 31, 1987

SUPPOSITORY; VAGINAL

TERAZOL 3

+ JANSSEN PHARMS 80MG ** N019641 001 May 24, 1988

TERCONAZOLE

FOUGERA PHARMS 80MG A076850 001 Jul 12, 2006

TERIFLUNOMIDE

TABLET; ORAL

TERIFLUNOMIDE

BRECKENRIDGE 7MG A209583 001 Sep 24, 2021

14MG A209583 002 Sep 24, 2021

WATSON LABS TEVA 7MG A209549 001 Jul 27, 2018

14MG A209549 002 Jul 27, 2018

TERIPARATIDE

SOLUTION; SUBCUTANEOUS

FORTEO

LILLY 0.75MG/3ML (0.25MG/ML) N021318 001 Nov 26, 2002

TERIPARATIDE ACETATE

INJECTABLE; INJECTION

PARATHAR

SANOFI AVENTIS US 200 UNITS/VIAL N019498 001 Dec 23, 1987

TESTOLACTONE

INJECTABLE; INJECTION

TESLAC

BRISTOL MYERS SQUIBB 100MG/ML N016119 001

TABLET; ORAL

TESLAC

BRISTOL MYERS SQUIBB 50MG N016118 001

250MG N016118 002

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

+ ABBVIE 2MG/24HR ** N020489 003 Oct 20, 2011

2.5MG/24HR N020489 001 Sep 29, 1995

+ 4MG/24HR ** N020489 004 Oct 20, 2011

5MG/24HR N020489 002 May 02, 1997

TESTODERM

ALZA 4MG/24HR N019762 001 Oct 12, 1993

6MG/24HR N019762 002 Oct 12, 1993

TESTODERM TTS

ALZA 5MG/24HR N020791 001 Dec 18, 1997

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TESTOSTERONE

GEL; TRANSDERMAL

ANDROGEL

+	BESINS HLTHCARE	1.62% (20.25MG/1.25GM PACKET) **	N022309	002	Sep 07, 2012
+		1.62% (40.5MG/2.5GM PACKET) **	N022309	003	Sep 07, 2012
+		25MG/2.5GM PACKET **	N021015	001	Feb 28, 2000
+		50MG/5GM PACKET **	N021015	002	Feb 28, 2000

TESTOSTERONE

ANI PHARMS	25MG/2.5GM PACKET	N202763	001	Feb 14, 2012
	50MG/5GM PACKET	N202763	002	Feb 14, 2012
PERRIGO ISRAEL	25MG/2.5GM PACKET	N203098	002	Jan 31, 2013
	50MG/5GM PACKET	N203098	003	Jan 31, 2013

GEL, METERED; TRANSDERMAL

ANDROGEL

+	BESINS HLTHCARE	12.5MG/1.25GM ACTUATION **	N021015	003	Sep 26, 2003
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TESTOSTERONE

ALEMBIC	1.62% (20.25MG/1.25GM ACTUATION)	A213922	001	Mar 03, 2021
PERRIGO ISRAEL	12.5MG/1.25GM ACTUATION	N203098	001	Jan 31, 2013

INJECTABLE; INJECTION

TESTOSTERONE

DR REDDYS	100MG/ML	A086417	001	Jul 07, 1983
WATSON LABS	25MG/ML	A086420	001	May 10, 1983
	50MG/ML	A086419	001	Aug 23, 1983

SOLUTION, METERED; TRANSDERMAL

AXIRON

+	ELI LILLY AND CO	30MG/1.5ML ACTUATION **	N022504	001	Nov 23, 2010
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TESTOSTERONE

ALEMBIC	30MG/1.5ML ACTUATION	A212882	001	Jun 14, 2021
APOTEX	30MG/1.5ML ACTUATION	A209181	001	Nov 25, 2020

TABLET, EXTENDED RELEASE; BUCCAL

STRIANT

+	AUXILIUM PHARMS LLC	30MG	N021543	001	Jun 19, 2003
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TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

PFIZER	50MG/ML	A085635	001	
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TESTOSTERONE CYPIONATE

RISING	200MG/ML	A040652	001	Dec 11, 2006
WATSON LABS	100MG/ML	A084401	001	
	100MG/ML	A086029	001	
	200MG/ML	A084401	002	

SOLUTION; INTRAMUSCULAR

TESTOSTERONE CYPIONATE

+	SLAYBACK PHARMA LLC	200MG/ML (200MG/ML)	N216318	001	Jun 02, 2022
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TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DELATESTRYL

ENDO PHARMS	200MG/ML	N009165	001	
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+		200MG/ML	N009165	003
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TESTOSTERONE ENANTHATE

RISING	200MG/ML	A040647	001	Oct 05, 2009
WATSON LABS	100MG/ML	A083667	001	
	100MG/ML	A085599	001	
	200MG/ML	A083667	002	

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION

TESTOSTERONE PROPIONATE

BEL MAR	25MG/ML	A080741	001	
	50MG/ML	A080742	001	
	100MG/ML	A080743	001	
ELKINS SINN	25MG/ML	A080276	001	
LILLY	50MG/ML	A080254	002	
WATSON LABS	25MG/ML	A080188	001	
	25MG/ML	A085490	001	
	50MG/ML	A080188	002	
	50MG/ML	A085490	002	
	100MG/ML	A080188	003	
	100MG/ML	A083595	003	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

AJANTA PHARMA LTD	12.5MG	A213621 001	Dec 04, 2020
	25MG	A213621 002	Dec 04, 2020
HIKMA	12.5MG	A209739 001	Apr 08, 2019
	25MG	A209739 002	Apr 08, 2019
SUN PHARM	12.5MG	A206129 001	Aug 17, 2015
	25MG	A206129 002	Aug 17, 2015

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

ACHROMYCIN V

+ AVET	250MG	N050278 003	
+	500MG	N050278 001	

BRISTACYCLINE

BRISTOL	250MG	A061658 001	
	250MG	A061888 001	
	500MG	A061658 002	
	500MG	A061888 002	

CYCLOPAR

WARNER CHILCOTT	250MG	A061725 001	
	250MG	A062175 001	
	250MG	A062332 001	
	500MG	A061725 002	
	500MG	A062332 002	

PANMYCIN

PHARMACIA AND UPJOHN	250MG	A060347 001	
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RETET

SOLVAY	250MG	A061443 001	
	500MG	A061443 002	

ROBITET

WYETH AYERST	250MG	A061734 001	
	500MG	A061734 002	

SUMYCIN

APOTHECON	100MG	A060429 002	
	125MG	A060429 004	
	250MG	A060429 001	
	500MG	A060429 003	

TETRACHEL

ANGUS	250MG	A060343 001	
	500MG	A060343 003	

TETRACYCLINE HYDROCHLORIDE

ABBOTT	250MG	A061802 001	
	500MG	A061802 002	
ANI PHARMS	250MG	A061471 001	
AUROBINDO PHARMA USA	250MG	A060783 001	
	500MG	A060783 002	
ELKINS SINN	250MG	A060059 001	
FERRANTE	125MG	A060173 001	
	250MG	A060173 002	
HEATHER	250MG	A061148 001	
	500MG	A061148 002	
HIKMA	250MG	A060768 001	
	500MG	A060768 002	
IMPAX LABS	100MG	A060469 002	
	250MG	A060469 001	
	500MG	A060469 003	
IVAX SUB TEVA PHARMS	250MG	A060704 001	
	500MG	A060704 002	
MAST MM	250MG	A062085 001	
PUREPAC PHARM	250MG	A060290 001	
	500MG	A060290 002	
PVT FORM	250MG	A062686 001	Jul 24, 1986
	500MG	A062686 002	Jul 24, 1986
ROXANE	500MG	A061214 002	
SUN PHARM INDUSTRIES	250MG	A060736 001	
	500MG	A060736 002	
SUPERPHARM	250MG	A062540 001	Mar 21, 1985
	500MG	A062540 002	Mar 21, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

VALEANT PHARM INTL	250MG	A060471 001
	500MG	A060471 002
WARNER CHILCOTT	250MG	A062300 001
	500MG	A062300 002
WATSON LABS	250MG	A062103 001
	250MG	A062343 001
	500MG	A062103 002
	500MG	A062343 002
WYETH AYERST	250MG	A061685 001
	500MG	A061685 002

TETRACYN

PFIPHARMECS	250MG	A060082 003
	500MG	A060082 004

FIBER, EXTENDED RELEASE; PERIODONTAL

ACTISITE

SCHIFF AND CO	12.7MG/FIBER	N050653 001	Mar 25, 1994
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FOR SOLUTION; TOPICAL

TOPICYCLINE

SHIRE	2.2MG/ML	N050493 001
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INJECTABLE; INJECTION

ACHROMYCIN

LEDERLE	250MG/VIAL	N050273 002
	500MG/VIAL	N050273 003

TETRACYN

PFIZER	250MG/VIAL	A060096 001
	500MG/VIAL	A060096 002

OINTMENT; OPHTHALMIC

ACHROMYCIN

STORZ	10MG/GM	N050266 001
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SUSPENSION; ORAL

ACHROMYCIN V

LEDERLE	125MG/5ML	N050263 002
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SUMYCIN

PAR PHARM	125MG/5ML	A060400 001
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TETRACYCLINE HYDROCHLORIDE

ALPHARMA US PHARMS	125MG/5ML	A060633 001
FERRANTE	125MG/5ML	A060174 001
PROTER	125MG/5ML	A060446 001
PUREPAC PHARM	125MG/5ML	A060291 001

TETRACYN

PFIPHARMECS	125MG/5ML	A060095 001
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TETRAMED

IVAX SUB TEVA PHARMS	125MG/5ML	A061468 001
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SUSPENSION/DROPS; OPHTHALMIC

ACHROMYCIN

STORZ	1%	N050268 001
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TABLET; ORAL

PANMYCIN

PHARMACIA AND UPJOHN	250MG	A061705 001
	500MG	A061705 002

SUMYCIN

STRIDES PHARMA	50MG	A061147 003
	100MG	A061147 002
	250MG	A061147 001
	500MG	A061147 004

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE; ORAL

TETREX

BRISTOL	EQ 100MG HYDROCHLORIDE	A061653 001
	EQ 250MG HYDROCHLORIDE	A061653 002
	EQ 250MG HYDROCHLORIDE	A061889 002
	EQ 250MG HYDROCHLORIDE	N050212 002
	EQ 500MG HYDROCHLORIDE	A061653 003
	EQ 500MG HYDROCHLORIDE	A061889 001
	EQ 500MG HYDROCHLORIDE	N050212 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THALIDOMIDE

CAPSULE; ORAL

THALIDOMIDE

NATCO

150MG

A213267 001 Apr 27, 2023

THALLOUS CHLORIDE TL-201

INJECTABLE; INJECTION

THALLOUS CHLORIDE TL 201

BRACCO

1mCi/ML

N018548 001 Dec 30, 1982

+ GE HEALTHCARE

1mCi/ML

N018110 002 Feb 27, 1996

+ LANTHEUS MEDCL

1mCi/ML

N017806 001

TRACE LIFE

1mCi/ML

A075569 001 Nov 21, 2001

INJECTABLE; INTRAVENOUS

THALLOUS CHLORIDE TL 201

CURIUM

2mCi/ML

A077698 001 Nov 09, 2006

+ LANTHEUS MEDCL

2mCi/ML

N017806 002 Oct 09, 1998

THEOPHYLLINE

CAPSULE; ORAL

BRONKODYL

SANOFI AVENTIS US

100MG

A085264 001

200MG

A085264 002

ELIXOPHYLLIN

FOREST LABS

100MG

A085545 001 Jul 31, 1984

200MG

A083921 001 Jul 31, 1984

SOMOPHYLLIN-T

FISONS

100MG

A087155 001 Feb 25, 1985

200MG

A087155 002 Feb 25, 1985

250MG

A087155 003 Feb 25, 1985

THEOPHYLLINE

KV PHARM

100MG

A085263 001

200MG

A085263 002

SCHERER RP

100MG

A084731 002 Nov 07, 1986

200MG

A084731 001 Nov 07, 1986

250MG

A084731 003 Nov 07, 1986

CAPSULE, EXTENDED RELEASE; ORAL

AEROLATE III

FLEMING PHARMS

65MG

A085075 003 Nov 24, 1986

AEROLATE JR

FLEMING PHARMS

130MG

A085075 002 Nov 24, 1986

AEROLATE SR

FLEMING PHARMS

260MG

A085075 001 Nov 24, 1986

ELIXOPHYLLIN SR

FOREST LABS

125MG

A086826 001 Jan 29, 1985

250MG

A086826 002 Jan 29, 1985

SLO-BID

SANOFI AVENTIS US

50MG

A088269 001 Jan 31, 1985

75MG

A089539 001 May 10, 1989

100MG

A087892 001 Jan 31, 1985

125MG

A089540 001 May 10, 1989

200MG

A087893 001 Jan 31, 1985

300MG

A087894 001 Jan 31, 1985

SLO-PHYLLIN

SANOFI AVENTIS US

60MG

A085206 001 May 24, 1982

+

125MG

A085203 001 May 24, 1982

250MG

A085205 001 May 24, 1982

SOMOPHYLLIN-CRT

GRAHAM DM

50MG

A087763 001 Feb 27, 1985

100MG

A087194 001

200MG

A088382 001 Feb 27, 1985

+

250MG

A087193 001

300MG

A088383 001 Feb 27, 1985

THEO-DUR

SCHERING

50MG

A088022 001 Sep 10, 1985

75MG

A088015 001 Sep 10, 1985

125MG

A088016 001 Sep 10, 1985

200MG

A087995 001 Sep 10, 1985

THEOBID

WHITBY

260MG

A085983 001 Mar 20, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE;ORAL

THEOBID JR.

WHITBY	130MG	A087854	001	Mar 20, 1985
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THEOCLEAR L.A.-130

SCHWARZ PHARMA	130MG	A086569	001	May 27, 1982
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THEOCLEAR L.A.-260

SCHWARZ PHARMA	260MG	A086569	002	May 27, 1982
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THEOPHYL-SR

ORTHO MCNEIL PHARM	125MG	A086480	001	Feb 08, 1985
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	250MG	A086471	001	Feb 08, 1985
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THEOPHYLLINE

CENT PHARMS	125MG	A088654	001	Feb 12, 1985
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	250MG	A088689	001	Feb 12, 1985
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HOSPIRA	100MG	A089976	001	Jan 04, 1995
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	200MG	A089977	001	Jan 04, 1995
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	300MG	A089932	001	Jan 04, 1995
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INWOOD LABS	100MG	A040052	001	Feb 14, 1994
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	125MG	A040052	002	Feb 14, 1994
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	200MG	A040052	003	Feb 14, 1994
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	300MG	A040052	004	Feb 14, 1994
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SANDOZ	260MG	A087462	001	May 11, 1982
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THEOPHYLLINE-SR

SCHERER RP	300MG	A088255	001	Jun 12, 1986
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THEOVENT

SCHERING	125MG	A087010	001	Jan 31, 1985
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	250MG	A087910	001	Jan 31, 1985
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ELIXIR;ORAL

ELIXOMIN

CENCI	80MG/15ML	A088303	001	Jan 25, 1984
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LANOPHYLLIN

LANNETT	80MG/15ML	A084578	001	
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THEOLIXIR

PANRAY	80MG/15ML	A084559	001	
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THEOPHYL-225

ORTHO MCNEIL PHARM	112.5MG/15ML	A086485	001	
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THEOPHYLLINE

ALPHARMA US PHARMS	80MG/15ML	A089223	001	May 27, 1988
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CENCI	80MG/15ML	A087679	001	Apr 15, 1982
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CHARTWELL RX	80MG/15ML	A085952	001	
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HALSEY	80MG/15ML	A085169	001	
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PHARM ASSOC	80MG/15ML	A086720	001	
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+ PRECISION DOSE	80MG/15ML	A085863	001	
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ROXANE	80MG/15ML	A084739	001	
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TARO	80MG/15ML	A089626	001	Oct 28, 1988
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WOCKHARDT	80MG/15ML	A086748	001	
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INJECTABLE;INJECTION

THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	40MG/100ML	N019083	001	Nov 07, 1984
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+ B BRAUN	40MG/100ML	N019826	001	Aug 14, 1992
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THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	80MG/100ML	N019083	002	Nov 07, 1984
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+ B BRAUN	80MG/100ML	N019826	002	Aug 14, 1992
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THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	160MG/100ML	N019083	003	Nov 07, 1984
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+ B BRAUN	160MG/100ML	N019826	003	Aug 14, 1992
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THEOPHYLLINE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	200MG/100ML	N019212	001	Nov 07, 1984
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	200MG/100ML	N019826	004	Aug 14, 1992
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THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER

+ B BRAUN	320MG/100ML	N019826	006	Aug 14, 1992
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THEOPHYLLINE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	4MG/ML	N019212	003	Nov 07, 1984
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	400MG/100ML	N019212	002	Nov 07, 1984
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	400MG/100ML	N019826	005	Aug 14, 1992
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THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4MG/ML	N018649	007	Jul 26, 1982
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	40MG/100ML	N018649	001	Jul 26, 1982
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	80MG/100ML	N018649	002	Jul 26, 1982
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

INJECTABLE; INJECTION

THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER				
	160MG/100ML	N018649	003	Jul 26, 1982
	200MG/100ML	N018649	004	Jul 26, 1982
	320MG/100ML	N018649	006	Nov 13, 1985
	400MG/100ML	N018649	005	Jul 26, 1982
THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER				
+	HOSPIRA INC	4MG/ML	N019211	007 Dec 14, 1984
+		40MG/100ML	N019211	001 Dec 14, 1984
		80MG/100ML	N019211	002 Dec 14, 1984
+		160MG/100ML	N019211	003 Dec 14, 1984
		200MG/100ML	N019211	004 Dec 14, 1984
+		320MG/100ML	N019211	006 Jan 20, 1988
		400MG/100ML	N019211	005 Dec 14, 1984

SOLUTION; ORAL

AEROLATE				
	FLEMING PHARMS	150MG/15ML	A089141	001 Dec 03, 1986
THEOLAIR				
	3M	80MG/15ML	A086107	001
THEOPHYLLINE				
+	ROXANE	80MG/15ML	A087449	001 Sep 15, 1983

SUSPENSION; ORAL

ELIXICON				
	FOREST LABS	100MG/5ML	A085502	001

SYRUP; ORAL

ACCURBRON				
	SANOFI AVENTIS US	150MG/15ML	A088746	001 Nov 22, 1985
AQUAPHYLLIN				
	FERNDAL LABS	80MG/15ML	A087917	001 Jan 18, 1983
SLO-PHYLLIN				
	SANOFI AVENTIS US	80MG/15ML	A085187	001
THEOCLEAR-80				
	CENT PHARMS	80MG/15ML	A087095	001 Mar 01, 1982
THEOPHYLLINE				
	ALPHARMA US PHARMS	80MG/15ML	A086001	001
+		150MG/15ML	A086545	001

TABLET; ORAL

QUIBRON-T				
	MONARCH PHARMS	300MG	A088656	001 Aug 22, 1985
SLO-PHYLLIN				
	SANOFI AVENTIS US	100MG	A085202	001
		200MG	A085204	001
THEOCLEAR-100				
	CENT PHARMS	100MG	A085353	002
THEOCLEAR-200				
	CENT PHARMS	200MG	A085353	001
THEOLAIR				
	MEDICIS	125MG	A086399	001
		250MG	A086399	002
THEOPHYL-225				
	ORTHO MCNEIL PHARM	225MG	A084726	001

TABLET, CHEWABLE; ORAL

THEOPHYL				
	ORTHO MCNEIL PHARM	100MG	A086506	001 Sep 12, 1985

TABLET, EXTENDED RELEASE; ORAL

DURAPHYL				
	FOREST LABS	100MG	A088503	001 Apr 03, 1985
		200MG	A088504	001 Apr 03, 1985
		300MG	A088505	001 Apr 03, 1985

LABID

	WARNER CHILCOTT	250MG	A087225	001
QUIBRON-T/SR				
	MONARCH PHARMS	300MG	A087563	001 Jun 21, 1983
SUSTAIRE				
	ROERIG	100MG	A085665	001
		300MG	A085665	002

T-PHYL

	PHARM RES ASSOC	200MG	A088253	001 Aug 17, 1983
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

TABLET, EXTENDED RELEASE;ORAL

THEO-DUR

+	SCHERING	100MG **	A085328	001	
+		200MG	A086998	001	
+		300MG **	A085328	002	
		450MG	A089131	001	Jun 25, 1986

THEOLAIR-SR

3M

		200MG	A088369	001	Jul 16, 1987
		250MG	A086363	002	Jul 16, 1987
		300MG	A088364	001	Jul 16, 1987
		500MG	A089132	001	Jul 16, 1987

THEOPHYLLINE

ABLE

		300MG	A040548	001	Apr 30, 2004
		400MG	A040543	001	Apr 27, 2004
		450MG	A040546	001	Apr 30, 2004
		600MG	A040539	001	Apr 27, 2004

HERITAGE PHARMA AVET

		100MG	A089807	001	Apr 30, 1990
		200MG	A089808	001	Apr 30, 1990

INWOOD LABS

		450MG	A040034	001	Apr 28, 1995
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UNI-DUR

SCHERING

		400MG	A089822	001	Jan 04, 1995
		600MG	A089823	001	Jan 04, 1995

THEOPHYLLINE SODIUM GLYCINATE

ELIXIR;ORAL

SYNOPHYLATE

CENT PHARMS

		EQ 165MG BASE/15ML	N006333	008	
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TABLET;ORAL

ASBRON

NOVARTIS

		EQ 150MG BASE	A085148	001	
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THIABENDAZOLE

SUSPENSION;ORAL

MINTEZOL

MERCK SHARP DOHME

		500MG/5ML	N016097	001	
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TABLET, CHEWABLE;ORAL

MINTEZOL

MERCK SHARP DOHME

		500MG	N016096	001	
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THIAMINE HYDROCHLORIDE

INJECTABLE;INJECTION

BETALIN S

+ LILLY

		100MG/ML	A080853	001	
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THIAMINE HYDROCHLORIDE

ABRAXIS PHARM

		100MG/ML	A080509	001	
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AKORN

		100MG/ML	A087968	001	Oct 01, 1982
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BEL MAR

		100MG/ML	A080718	001	
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		200MG/ML	A080712	001	
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DELL LABS

		100MG/ML	A083775	001	
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HOSPIRA

		100MG/ML	A040079	001	May 03, 1996
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LUITPOLD

		100MG/ML	A080667	001	
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PARKE DAVIS

		100MG/ML	A080770	001	
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WATSON LABS

		100MG/ML	A083534	001	
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		200MG/ML	A083534	002	
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WYETH AYERST

		100MG/ML	A080553	001	
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THIAMYLAL SODIUM

INJECTABLE;INJECTION

SURITAL

PARKEDALE

		1GM/VIAL	N007600	003	
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		5GM/VIAL	N007600	005	
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		10GM/VIAL	N007600	009	
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THIETHYLPERAZINE MALATE

INJECTABLE;INJECTION

TORECAN

NOVARTIS

		5MG/ML	N012754	002	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIETHYLPERAZINE MALEATE

SUPPOSITORY;RECTAL

TORECAN

NOVARTIS 10MG N013247 001

TABLET;ORAL

TORECAN

NOVARTIS 10MG N012753 001

THIOPENTAL SODIUM

SUSPENSION;RECTAL

PENTOTHAL

ABBOTT 400MG/GM N011679 001

THIORIDAZINE

SUSPENSION;ORAL

MELLARIL-S

NOVARTIS EQ 25MG HYDROCHLORIDE/5ML ** N017923 001

EQ 100MG HYDROCHLORIDE/5ML ** N017923 002

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE;ORAL

MELLARIL

NOVARTIS 30MG/ML ** N011808 012

100MG/ML ** N011808 018

THIORIDAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 100MG/ML A088229 001 Aug 23, 1983

ALPHARMA US PHARMS 30MG/ML A087766 001 Apr 26, 1983

ANI PHARMS 30MG/ML A089602 001 Nov 09, 1987

100MG/ML A089603 001 Nov 09, 1987

EPIC PHARMA LLC 30MG/ML A040125 001 Aug 16, 1996

100MG/ML A040126 001 Aug 16, 1996

PHARM ASSOC 30MG/ML A040187 001 Aug 28, 1997

100MG/ML A040213 001 May 29, 1998

SANDOZ 30MG/ML A088307 001 Nov 23, 1983

100MG/ML A088308 001 Nov 23, 1983

WOCKHARDT 30MG/ML A088258 001 Jul 25, 1983

100MG/ML A088227 001 Jul 05, 1983

THIORIDAZINE HYDROCHLORIDE INTENSOL

ROXANE 30MG/ML A088941 001 Dec 16, 1985

100MG/ML A088942 001 Dec 16, 1985

TABLET;ORAL

MELLARIL

+ NOVARTIS 10MG ** N011808 003

+ 15MG ** N011808 016

+ 25MG ** N011808 006

+ 50MG ** N011808 011

+ 100MG ** N011808 009

+ 150MG ** N011808 017

+ 200MG ** N011808 015

THIORIDAZINE HYDROCHLORIDE

ANI PHARMS 10MG A088270 001 Apr 14, 1983

10MG A088493 001 May 17, 1985

15MG A088271 001 Apr 14, 1983

25MG A088272 001 Apr 14, 1983

50MG A088194 001 Apr 14, 1983

100MG A088273 001 Oct 03, 1983

100MG A088456 001 May 17, 1985

CHARTWELL RX 10MG A088131 001 Aug 30, 1983

15MG A088132 001 Aug 30, 1983

25MG A088133 001 Aug 30, 1983

50MG A088134 001 Aug 30, 1983

100MG A088135 001 Nov 20, 1984

150MG A088136 001 Sep 17, 1986

200MG A088137 001 Sep 17, 1986

HERITAGE PHARMA AVET 10MG A088476 001 Nov 08, 1983

25MG A088478 001 Nov 08, 1983

50MG A088479 001 Nov 08, 1983

100MG A088736 001 Jul 24, 1984

MUTUAL PHARM 10MG A088375 001 Nov 18, 1983

25MG A087264 001 Nov 18, 1983

50MG A088370 001 Nov 18, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

	100MG	A088379	001	Nov 16, 1983
MYLAN	10MG	A088332	001	Jun 27, 1983
	25MG	A088333	001	Jun 27, 1983
	50MG	A088334	001	Jun 27, 1983
	100MG	A088335	001	Nov 18, 1983
PAR PHARM	10MG	A088351	001	Dec 05, 1983
	15MG	A088352	001	Dec 05, 1983
	25MG	A088336	001	Dec 05, 1983
	50MG	A088322	001	Dec 05, 1983
	100MG	A088480	001	Dec 29, 1983
	150MG	A089764	001	Feb 09, 1988
	200MG	A089765	001	Feb 09, 1988
ROXANE	10MG	A088663	001	Mar 15, 1984
	25MG	A088664	001	Mar 15, 1984
	50MG	A088665	001	Mar 15, 1984
	100MG	A089048	001	Feb 26, 1985
SUN PHARM INDUSTRIES	10MG	A089953	004	Aug 01, 1986
	15MG	A088461	001	Nov 18, 1983
	25MG	A089953	003	Aug 01, 1986
	50MG	A089953	002	Aug 01, 1986
	100MG	A089953	001	Oct 07, 1988
	150MG	A088737	001	Sep 26, 1984
	200MG	A088738	001	Oct 16, 1984
SUPERPHARM	10MG	A089103	001	Jul 02, 1985
	25MG	A089104	001	Jul 02, 1985
	50MG	A089105	001	Jul 02, 1985
WATSON LABS	10MG	A088412	001	Sep 12, 1983
	10MG	A088561	001	May 11, 1984
	15MG	A088345	001	Jul 28, 1983
	15MG	A088562	001	May 11, 1984
	25MG	A088296	001	Jul 28, 1983
	25MG	A088755	001	Jul 24, 1984
	50MG	A088323	001	Jul 28, 1983
	50MG	A088563	001	May 11, 1984
	100MG	A088284	001	Aug 25, 1983
	100MG	A088564	001	May 11, 1984
	150MG	A088410	001	Mar 05, 1984
	150MG	A088869	001	Jun 28, 1985
	200MG	A088381	001	Mar 14, 1984
WATSON LABS TEVA	15MG	A088477	001	Nov 08, 1983
	25MG	A088567	001	May 11, 1984
	200MG	A088872	001	Apr 26, 1985
WEST WARD	10MG	A088658	001	Mar 26, 1984
	15MG	A088659	001	Mar 26, 1984
	25MG	A088660	001	Mar 26, 1984
	50MG	A088661	001	Mar 26, 1984

THIOTEPA

INJECTABLE; INJECTION

THIOPLEX

+ IMMUNEX

15MG/VIAL **

N020058 001 Dec 22, 1994

THIOTEPA

FRESENIUS KABI USA

15MG/VIAL

A075698 001 Sep 20, 2001

IMMUNEX

15MG/VIAL

N011683 001

TEVA PARENTERAL

15MG/VIAL **

A075730 001 Apr 20, 2001

30MG/VIAL **

A075730 002 Apr 20, 2001

THIOTHIXENE

CAPSULE; ORAL

NAVANE

+ PFIZER

1MG **

N016584 001

+

2MG **

N016584 002

+

5MG **

N016584 003

+

10MG **

N016584 004

+

20MG **

N016584 005

THIOTHIXENE

AM THERAP

1MG

A071884 001 Aug 12, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

	2MG	A071885 001	Aug 12, 1987
	5MG	A071886 001	Aug 12, 1987
	10MG	A071887 001	Aug 12, 1987
	20MG	A072200 001	Dec 17, 1987
CHARTWELL RX	1MG	A070600 001	Jun 05, 1987
	2MG	A070601 001	Jun 05, 1987
	5MG	A070602 001	Jun 05, 1987
	10MG	A070603 001	Jun 05, 1987
EPIC PHARMA LLC	1MG	A071529 002	Jun 24, 1987
	2MG	A071529 003	Jun 24, 1987
	5MG	A071529 001	Jun 24, 1987
	10MG	A071529 004	Jun 24, 1987
WATSON LABS	2MG	A071626 001	Jun 25, 1987
	5MG	A071627 001	Jun 25, 1987
	10MG	A071628 001	Jun 25, 1987

THIOTHIXENE HYDROCHLORIDE

CONCENTRATE; ORAL

NAVANE

+ PFIZER

EQ 5MG BASE/ML

N016758 001

THIOTHIXENE HYDROCHLORIDE

ALPHARMA US PHARMS

EQ 5MG BASE/ML

A070969 001 Oct 16, 1987

PACO

EQ 1MG BASE/ML

A071917 001 Sep 20, 1989

EQ 5MG BASE/ML

A071939 001 Dec 16, 1988

TEVA

EQ 5MG BASE/ML

A071184 001 Jun 22, 1987

TEVA PHARMS

EQ 5MG BASE/ML

A071554 001 Oct 16, 1987

THIOTHIXENE HYDROCHLORIDE INTENSOL

HIKMA

EQ 5MG BASE/ML

A073494 001 Jun 30, 1992

INJECTABLE; INJECTION

NAVANE

PFIZER

EQ 2MG BASE/ML

N016904 001

EQ 10MG BASE/VIAL

N016904 002

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

CEPHALON

6MG

N020646 006 Nov 29, 2005

8MG

N020646 007 Nov 29, 2005

10MG

N020646 008 Nov 29, 2005

20MG

N020646 004 Sep 30, 1997

TIAGABINE HYDROCHLORIDE

AMNEAL PHARMS CO

2MG

A208181 001 Dec 08, 2017

4MG

A208181 002 Dec 08, 2017

12MG

A208181 003 Dec 08, 2017

16MG

A208181 004 Dec 08, 2017

WILSHIRE PHARMS INC

2MG

A206857 001 Oct 13, 2017

4MG

A206857 002 Oct 13, 2017

12MG

A206857 003 Oct 13, 2017

16MG

A206857 004 Oct 13, 2017

TICAGRELOR

TABLET; ORAL

TICAGRELOR

AMNEAL

90MG

A208531 001 Jan 23, 2019

MYLAN

60MG

A208597 001 Jul 09, 2021

90MG

A208597 002 Jul 09, 2021

SIGMAPHARM LABS LLC

90MG

A208596 001 Apr 07, 2020

SUNSHINE

90MG

A208508 001 Apr 06, 2020

WATSON LABS INC

60MG

A208390 001 Sep 04, 2018

90MG

A208390 002 Sep 04, 2018

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

GLAXOSMITHKLINE

EQ 1GM BASE/VIAL

N050497 001

EQ 3GM BASE/VIAL

A062690 001 Dec 19, 1986

EQ 3GM BASE/VIAL

N050497 002

EQ 6GM BASE/VIAL

N050497 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

EQ 20GM BASE/VIAL	N050497 004	
EQ 30GM BASE/VIAL	N050497 005	Apr 04, 1984

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLID

ROCHE PALO	125MG	N019979 001	Mar 24, 1993
	250MG	N019979 002	Oct 31, 1991

TICLOPIDINE HYDROCHLORIDE

ACTAVIS ELIZABETH	250MG	A075253 001	Aug 20, 1999
CHARTWELL RX	250MG	A075318 001	Aug 20, 1999
	250MG	A075326 001	Aug 20, 1999
MYLAN	250MG	A075161 001	Sep 13, 1999
	250MG	A075316 001	Nov 02, 1999
SUN PHARM INDS INC	250MG	A075526 001	Sep 26, 2002
TEVA	250MG	A075149 001	Aug 20, 1999
WATSON LABS	250MG	A075309 001	Apr 26, 2000

TIGECYCLINE

POWDER; INTRAVENOUS

TIGECYCLINE

XELLIA PHARMS APS	50MG/VIAL	A205722 001	Oct 18, 2019
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TILUDRONATE DISODIUM

TABLET; ORAL

SKELID

+ SANOFI AVENTIS US	EQ 200MG BASE **	N020707 001	Mar 07, 1997
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TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL

EPIC PHARMA LLC	EQ 0.25% BASE	A205309 001	Sep 30, 2016
	EQ 0.5% BASE	A205309 002	Sep 30, 2016

TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

AKORN	EQ 0.25% BASE	A074465 001	Mar 25, 1997
	EQ 0.25% BASE	A074515 001	Mar 25, 1997
	EQ 0.5% BASE	A074466 001	Mar 25, 1997
	EQ 0.5% BASE	A074516 001	Mar 25, 1997
APOTEX INC	EQ 0.25% BASE	A075411 001	Sep 08, 2000
	EQ 0.5% BASE	A075412 001	Sep 08, 2000
FOUGERA	EQ 0.25% BASE	A074667 001	Mar 25, 1997
	EQ 0.5% BASE	A074668 001	Mar 25, 1997
HIKMA	EQ 0.5% BASE	A075163 001	Sep 10, 2002

TABLET; ORAL

BLOCADREN

+ MERCK	5MG **	N018017 001	
+	10MG **	N018017 002	
+	20MG **	N018017 004	

TIMOLOL MALEATE

ANI PHARMS	5MG	A072917 001	Jul 31, 1991
	10MG	A072918 001	Jul 31, 1991
	20MG	A072919 001	Jul 31, 1991
CHARTWELL RX	5MG	A072550 001	Apr 13, 1989
	10MG	A072551 001	Apr 13, 1989
	20MG	A072552 001	Apr 13, 1989
QUANTUM PHARMICS	5MG	A072466 001	May 19, 1989
	10MG	A072467 001	May 19, 1989
	20MG	A072468 001	May 19, 1989
TEVA	5MG	A072648 001	Jun 16, 1993
	10MG	A072649 001	Jun 16, 1993
	20MG	A072650 001	Jun 16, 1993
USL PHARMA	5MG	A072001 001	Apr 11, 1989
	10MG	A072002 001	Apr 11, 1989
	20MG	A072003 001	Apr 11, 1989
WATSON LABS	5MG	A072269 001	Apr 11, 1989
	10MG	A072270 001	Apr 11, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TIMOLOL MALEATETABLET; ORAL
TIMOLOL MALEATE

20MG A072271 001 Apr 11, 1989

TINIDAZOLETABLET; ORAL
TINIDAZOLEHIKMA 250MG A201172 001 Apr 30, 2012
500MG A201172 002 Apr 30, 2012TINZAPARIN SODIUMINJECTABLE; INJECTION
INNOHEP

LEO PHARMA AS 20,000 IU/ML N020484 001 Jul 14, 2000

TIOCONAZOLECREAM; TOPICAL
TZ-3

PFIZER 1% N018682 001 Feb 18, 1983

TIOPRONINTABLET, DELAYED RELEASE; ORAL
TIOPRONINPAR PHARM 100MG A217219 001 Feb 24, 2023
300MG A217219 002 Feb 24, 2023TIPRANAVIRSOLUTION; ORAL
APTIVUS

+ BOEHRINGER INGELHEIM 100MG/ML N022292 001 Jun 23, 2008

TIROFIBAN HYDROCHLORIDEINJECTABLE; INJECTION
AGGRASTAT

MEDICURE EQ 12.5MG BASE/50ML (EQ 0.25MG BASE/ML) N020912 001 May 14, 1998

SOLUTION; INTRAVENOUS
AGGRASTAT

MEDICURE EQ 25MG BASE/500ML (EQ 0.05MG BASE/ML) N020913 001 May 14, 1998

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

TIZANIDINE HYDROCHLORIDE
MYLAN PHARMS INCEQ 2MG BASE A091502 001 Nov 09, 2012
EQ 4MG BASE A091502 002 Nov 09, 2012
EQ 6MG BASE A091502 003 Nov 09, 2012
PAR PHARM INC EQ 2MG BASE A207199 001 Mar 14, 2017
EQ 4MG BASE A207199 002 Mar 14, 2017
EQ 6MG BASE A207199 003 Mar 14, 2017

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE
ANI PHARMSEQ 2MG BASE A076283 001 Jul 12, 2002
EQ 2MG BASE A076284 001 Jul 03, 2002
EQ 2MG BASE A076321 001 Sep 30, 2004
EQ 2MG BASE A076371 001 Apr 09, 2003
EQ 4MG BASE A076283 002 Jul 12, 2002
EQ 4MG BASE A076284 002 Jul 03, 2002
EQ 4MG BASE A076321 002 Sep 30, 2004
EQ 4MG BASE A076371 002 Apr 09, 2003
AUROBINDO PHARMA USA EQ 2MG BASE A076282 001 Dec 16, 2003
EQ 4MG BASE A076282 002 Dec 16, 2003
PAR PHARM INC EQ 2MG BASE A207170 001 Jan 26, 2017
EQ 4MG BASE A207170 002 Jan 26, 2017
RISING EQ 2MG BASE A076354 001 Mar 28, 2003
EQ 4MG BASE A076354 002 Mar 28, 2003

ZANAFLEX

+ LEGACY PHARMA USA EQ 2MG BASE ** N020397 002 Feb 04, 2000

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOBRAMYCIN

SOLUTION; INHALATION

TOBRAMYCIN

LUOXIN AUROVITAS	300MG/5ML	A210871 001	Jan 22, 2021
MYLAN	300MG/5ML	A209554 001	Oct 13, 2017

SOLUTION/DROPS; OPHTHALMIC

AKTOB

EPIC PHARMA LLC	0.3%	A064096 001	Jan 31, 1996
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TOBRAMYCIN

ALCON PHARMS LTD	0.3%	A063176 001	May 25, 1994
APOTEX INC	0.3%	A065087 001	Feb 25, 2002

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

NEBCIN

LILLY	EQ 10MG BASE/ML	A062008 004	
	EQ 10MG BASE/ML	A062707 001	Apr 29, 1987
+	EQ 10MG BASE/ML **	N050477 005	
	EQ 40MG BASE/ML	A062008 001	
+	EQ 1.2GM BASE/VIAL **	N050519 001	

TOBRAMYCIN SULFATE

APOTHECON

EQ 10MG BASE/ML	A064021 001	May 31, 1994
EQ 40MG BASE/ML	A064021 002	May 31, 1994
EQ 40MG BASE/ML	A064026 001	May 31, 1994

EPIC PHARMA LLC

EQ 40MG BASE/ML	A205179 001	Sep 16, 2014
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HIKMA

EQ 10MG BASE/ML	A063113 001	Apr 26, 1991
EQ 10MG BASE/ML	A063128 001	Nov 27, 1991
EQ 40MG BASE/ML	A063118 001	Jul 29, 1991
EQ 40MG BASE/ML	A063127 001	Nov 27, 1991

HOSPIRA

EQ 10MG BASE/ML	A063080 001	Apr 30, 1991
EQ 10MG BASE/ML	A063112 001	Apr 30, 1991
EQ 40MG BASE/ML	A063161 001	May 29, 1991

IGI LABS INC

EQ 10MG BASE/ML	A063119 001	Oct 31, 1994
EQ 40MG BASE/ML	A063120 001	Oct 31, 1994
EQ 40MG BASE/ML	A063121 001	Oct 31, 1994
EQ 40MG BASE/ML	A063122 001	Oct 31, 1994

TEVA PHARMS USA

EQ 40MG BASE/ML	A063100 001	Jan 30, 1992
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WATSON LABS INC

EQ 10MG BASE/ML	A062945 001	Aug 09, 1989
EQ 40MG BASE/ML	A062945 002	Aug 09, 1989

TOBRAMYCIN SULFATE (PHARMACY BULK)

HOSPIRA

EQ 40MG BASE/ML **	A063116 001	May 18, 1992
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TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA

EQ 1.2MG BASE/ML	A063081 003	Jul 31, 1990
EQ 1.6MG BASE/ML	A063081 006	Jun 02, 1993
EQ 80MG BASE/100ML	A063081 001	Jul 31, 1990

TOCAINIDE HYDROCHLORIDE

TABLET; ORAL

TONOCARD

ASTRAZENECA

400MG	N018257 001	Nov 09, 1984
600MG	N018257 002	Nov 09, 1984

TOFACITINIB CITRATE

TABLET; ORAL

TOFACITINIB CITRATE

AJANTA PHARMA LTD

EQ 10MG BASE	A212943 001	Jun 01, 2021
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MICRO LABS

EQ 5MG BASE	A209738 001	Mar 13, 2023
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ZYDUS PHARMS

EQ 5MG BASE	A209829 001	Mar 13, 2023
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TABLET, EXTENDED RELEASE; ORAL

TOFACITINIB

ZYDUS PHARMS

EQ 11MG BASE	A214264 001	Aug 19, 2021
EQ 22MG BASE	A214264 002	Aug 19, 2021

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

BARR

100MG	A070162 001	Jan 14, 1986
250MG	A070163 001	Jan 14, 1986
500MG	A070164 001	Jan 14, 1986

CHARTWELL RX

100MG	A071633 001	Dec 09, 1987
250MG	A070289 001	Mar 13, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLAZAMIDETABLET; ORAL
TOLAZAMIDE

	500MG		A070290	001	Mar 13, 1986
COSETTE	100MG		N018894	001	Nov 02, 1984
	250MG		N018894	002	Nov 02, 1984
	500MG		N018894	003	Nov 02, 1984
DURAMED PHARMS BARR	100MG		A070165	001	Jan 10, 1986
	250MG		A070166	001	Jan 10, 1986
	500MG		A070167	001	Jan 10, 1986
INTERPHARM	250MG		A071270	001	Sep 23, 1986
	500MG		A071271	001	Sep 23, 1986
MYLAN PHARMS INC	250MG		A070259	001	Jan 02, 1986
	500MG		A070259	003	Mar 17, 1986
PAR PHARM	100MG		A070159	001	Jan 06, 1986
	250MG		A070160	001	Jan 06, 1986
	500MG		A070161	001	Jan 06, 1986
SUN PHARM INDUSTRIES	100MG		A071357	001	Jul 16, 1987
	250MG		A071358	001	Jul 16, 1987
	500MG		A071359	001	Jul 16, 1987
SUPERPHARM	250MG		A070763	001	Jun 16, 1986
	500MG		A070764	001	Jun 16, 1986
USL PHARMA	100MG		A071355	001	Jan 11, 1988
	250MG		A070168	001	Apr 02, 1986
	500MG		A070169	001	Apr 02, 1986
WATSON LABS	100MG		A070242	001	Aug 01, 1986
	100MG		A070513	001	Jan 09, 1986
	250MG		A070243	001	Aug 01, 1986
	250MG		A070514	001	Jan 09, 1986
	500MG		A070244	001	Aug 01, 1986
	500MG		A070515	001	Jan 09, 1986
TOLINASE					
+ PHARMACIA AND UPJOHN	100MG **		N015500	002	
+	250MG **		N015500	004	
+	500MG **		N015500	005	

TOLAZOLINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRISCOLINE

NOVARTIS 25MG/ML N006403 005 Feb 22, 1985

TOLBUTAMIDE

TABLET; ORAL

ORINASE

PHARMACIA AND UPJOHN	250MG **		N010670	002	
	500MG **		N010670	001	
TOLBUTAMIDE					
ALRA	500MG		A086141	001	
ANI PHARMS	500MG		A087093	001	
ASCOT	500MG		A087541	001	Mar 01, 1983
BARR	500MG		A087121	001	
CHARTWELL RX	500MG		A086574	001	
DAVA PHARMS INC	500MG		A086926	001	
MYLAN PHARMS INC	500MG		A086445	001	
PARKE DAVIS	500MG		A086047	001	
PUREPAC PHARM	500MG		A088950	001	Jun 17, 1985
SANDOZ	500MG		N012678	001	
SUPERPHARM	500MG		A088893	001	Nov 19, 1984
VANGARD	500MG		A087876	001	Apr 20, 1982
WATSON LABS	250MG		A089110	001	May 29, 1987
	500MG		A086109	001	
	500MG		A087318	001	
	500MG		A089111	001	May 29, 1987

TOLBUTAMIDE SODIUM

INJECTABLE; INJECTION

ORINASE DIAGNOSTIC

PHARMACIA AND UPJOHN EQ 1GM BASE/VIAL N012095 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLCAPONE

TABLET; ORAL

TASMAR

BAUSCH 200MG N020697 002 Jan 29, 1998

TOLCAPONE

ALVOGEN 100MG A207729 001 Jul 29, 2020

DR REDDYS LABS SA 100MG A210095 001 Aug 01, 2019

TOLMETIN SODIUM

CAPSULE; ORAL

TOLECTIN DS

+ ORTHO MCNEIL JANSSEN EQ 400MG BASE N018084 001

TOLMETIN SODIUM

ANI PHARMS EQ 400MG BASE A073308 001 Jan 24, 1992

EQ 400MG BASE A073392 001 Jan 24, 1992

EQ 400MG BASE A073519 001 May 29, 1992

FOSUN PHARMA EQ 400MG BASE A073462 001 Apr 30, 1992

SUN PHARM INDUSTRIES EQ 400MG BASE A073311 001 Nov 27, 1991

TEVA EQ 400MG BASE A073290 001 Nov 27, 1991

TABLET; ORAL

TOLECTIN

+ ORTHO MCNEIL JANSSEN EQ 200MG BASE N017628 001

TOLECTIN 600

+ ORTHO MCNEIL JANSSEN EQ 600MG BASE N017628 002 Mar 08, 1989

TOLMETIN SODIUM

ANI PHARMS EQ 600MG BASE A073527 001 Jun 30, 1992

COSETTE EQ 600MG BASE A074399 001 Mar 28, 1996

EQ 600MG BASE A074729 001 Feb 27, 1997

FOSUN PHARMA EQ 200MG BASE A073588 001 Jul 31, 1992

EQ 600MG BASE A074002 001 Sep 27, 1993

SUN PHARM INDUSTRIES EQ 200MG BASE A073310 001 Nov 27, 1991

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

TOLTERODINE TARTRATE

AUROBINDO PHARMA USA 2MG A201486 001 Oct 31, 2013

4MG A201486 002 Oct 31, 2013

TABLET; ORAL

TOLTERODINE TARTRATE

APOTEX CORP 1MG A200164 001 Sep 25, 2012

2MG A200164 002 Sep 25, 2012

MYLAN PHARMS INC 1MG A202641 001 Nov 27, 2012

2MG A202641 002 Nov 27, 2012

TOLVAPTAN

TABLET; ORAL

SAMSCA

+ OTSUKA 60MG ** N022275 003 May 19, 2009

TOPIRAMATE

CAPSULE; ORAL

TOPAMAX SPRINKLE

+ JANSSEN PHARMS 50MG ** N020844 003 Oct 26, 1998

TOPIRAMATE

BARR 15MG A076448 001 Apr 15, 2009

25MG A076448 002 Apr 15, 2009

FOSUN PHARMA 15MG A079206 001 Oct 14, 2009

25MG A079206 002 Oct 14, 2009

STRIDES PHARMA 15MG A078418 001 Oct 14, 2009

25MG A078418 002 Oct 14, 2009

WATSON LABS 15MG A077868 001 Apr 15, 2009

25MG A077868 002 Apr 15, 2009

CAPSULE, EXTENDED RELEASE; ORAL

TOPIRAMATE

ZYDUS PHARMS 200MG A216167 001 Feb 09, 2023

TABLET; ORAL

TOPAMAX

JANSSEN PHARMS 300MG N020505 003 Dec 24, 1996

400MG N020505 006 Dec 24, 1996

TOPIRAMATE

ACTAVIS TOTOWA 25MG A078637 001 Feb 27, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOPIRAMATETABLET; ORAL
TOPIRAMATE

	50MG	A078637 002	Feb 27, 2013
	100MG	A078637 003	Feb 27, 2013
	200MG	A078637 004	Feb 27, 2013
BARR	25MG	A076315 001	Mar 27, 2009
	100MG	A076315 002	Mar 27, 2009
	200MG	A076315 003	Mar 27, 2009
CHARTWELL	25MG	A078410 001	Sep 11, 2013
	50MG	A078410 002	Sep 11, 2013
	100MG	A078410 003	Sep 11, 2013
	200MG	A078410 004	Sep 11, 2013
HIKMA PHARMS	25MG	A091185 001	Nov 25, 2013
	50MG	A091185 002	Nov 25, 2013
	100MG	A091185 003	Nov 25, 2013
	200MG	A091185 004	Nov 25, 2013
MYLAN	25MG	A076314 001	Mar 27, 2009
	50MG	A076314 002	Mar 27, 2009
	100MG	A076314 003	Mar 27, 2009
	200MG	A076314 004	Mar 27, 2009
PLIVA HRVATSKA DOO	25MG	A077905 001	Mar 30, 2009
	50MG	A077905 002	Mar 30, 2009
	100MG	A077905 003	Mar 30, 2009
	200MG	A077905 004	Mar 30, 2009
ROXANE	25MG	A076306 001	Mar 27, 2009
	50MG	A076306 002	Mar 27, 2009
	100MG	A076306 003	Mar 27, 2009
	200MG	A076306 004	Mar 27, 2009
TEVA	25MG	A076317 001	Mar 27, 2009
	50MG	A076317 002	Mar 27, 2009
	100MG	A076317 003	Mar 27, 2009
	200MG	A076317 004	Mar 27, 2009
TORRENT PHARMS	25MG	A079153 001	Mar 27, 2009
	50MG	A079153 002	Mar 27, 2009
	100MG	A079153 003	Mar 27, 2009
	200MG	A079153 004	Mar 27, 2009
UPSHER SMITH LABS	25MG	A078499 001	Jan 07, 2010
	50MG	A078499 002	Jan 07, 2010
	100MG	A078499 003	Jan 07, 2010
	200MG	A078499 004	Jan 07, 2010
WATSON LABS	25MG	A077643 001	Mar 27, 2009
	50MG	A077643 002	Mar 27, 2009
	100MG	A077643 003	Mar 27, 2009
	200MG	A077643 004	Mar 27, 2009
WOCKHARDT	25MG	A090353 001	Sep 01, 2010
	50MG	A090353 002	Sep 01, 2010
	100MG	A090353 003	Sep 01, 2010
	200MG	A090353 004	Sep 01, 2010

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

TOPOTECAN HYDROCHLORIDE

FRESENIUS KABI USA	EQ 4MG BASE/VIAL	A091376 001	Nov 29, 2010
MEITHEAL	EQ 4MG BASE/VIAL	A201166 001	Aug 08, 2012
MYLAN LABS LTD	EQ 4MG BASE/VIAL	A091542 001	Aug 28, 2012
SUN PHARM INDS LTD	EQ 4MG BASE/VIAL	A202203 001	Aug 29, 2013

SOLUTION; INTRAVENOUS

TOPOTECAN

+ SANDOZ INC	EQ 1MG BASE/ML (EQ 1MG BASE/ML) **	N200199 001	Feb 25, 2011
+	EQ 3MG BASE/3ML (EQ 1MG BASE/ML) **	N200199 002	Feb 25, 2011
+	EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **	N200199 003	Feb 25, 2011

TOPOTECAN HYDROCHLORIDE

DASH PHARMS	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	A206074 001	Nov 24, 2017
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TORSEMIDE

INJECTABLE; INJECTION

DEMADEX

+	ROCHE	50MG/5ML (10MG/ML) **	N020137	002	Aug 23, 1993
+		20MG/2ML (10MG/ML) **	N020137	001	Aug 23, 1993

TORSEMIDE

	AM REGENT	20MG/2ML (10MG/ML)	A090656	001	Apr 21, 2010
		50MG/5ML (10MG/ML)	A090656	002	Apr 21, 2010
	WEST-WARD PHARMS INT	20MG/2ML (10MG/ML)	A078007	001	Jun 11, 2008
		50MG/5ML (10MG/ML)	A078007	002	Jun 11, 2008

TABLET; ORAL

DEMADEX

+	MYLAN SPECIALITY LP	5MG **	N020136	001	Aug 23, 1993
+		10MG **	N020136	002	Aug 23, 1993
+		20MG **	N020136	003	Aug 23, 1993
+		100MG **	N020136	004	Aug 23, 1993

SOAANZ

+	SARFE PHARMS	20MG	N213218	001	Jun 14, 2021
+		60MG	N213218	002	Jun 14, 2021

TORSEMIDE

	SUN PHARM INDS	5MG	A078478	001	Feb 26, 2008
		10MG	A078478	002	Feb 26, 2008
		20MG	A078478	003	Feb 26, 2008
		100MG	A078478	004	Feb 26, 2008
	TEVA	5MG	A076110	001	May 14, 2002
		10MG	A076110	002	May 14, 2002
		20MG	A076110	003	May 14, 2002
		100MG	A076110	004	May 14, 2002

TRAMADOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CONZIP

+	CIPHER PHARMS INC	150MG	N022370	004	Aug 01, 2011
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TABLET; ORAL

TRAMADOL HYDROCHLORIDE

	ACCORD HLTHCARE	50MG	A202390	001	May 16, 2013
	ACTAVIS ELIZABETH	50MG	A075960	001	Jun 19, 2002
	ASTA	50MG	A075974	001	Jul 12, 2002
	GRAVITI PHARMS	50MG	A075968	001	Jun 25, 2002
	IVAX SUB TEVA PHARMS	50MG	A075963	001	Jul 03, 2002
	MACLEODS PHARMS LTD	50MG	A205702	001	Sep 25, 2015
	MYLAN	50MG	A075986	001	Jun 21, 2002
	MYLAN PHARMS INC	50MG	A075980	001	Nov 21, 2002
	NORTHSTAR HLTHCARE	50MG	A078935	001	May 26, 2010
	PLIVA	50MG	A075982	001	Jul 01, 2002
	SPECGX LLC	50MG	A075983	001	Jun 25, 2002
	SUN PHARM INDUSTRIES	50MG	A076100	001	Jun 20, 2002
	WATSON LABS	50MG	A075962	001	Jun 24, 2002

ULTRAM

+	JANSSEN PHARMS	50MG **	N020281	002	Mar 03, 1995
+		100MG **	N020281	001	Mar 03, 1995

TABLET, EXTENDED RELEASE; ORAL

RYZOLT

+	PURDUE PHARMA	100MG **	N021745	001	Dec 30, 2008
+		200MG **	N021745	002	Dec 30, 2008
+		300MG **	N021745	003	Dec 30, 2008

TRAMADOL HYDROCHLORIDE

	ACTAVIS ELIZABETH	100MG	A091609	001	Jun 27, 2012
		200MG	A091609	002	Jun 27, 2012
		300MG	A091609	003	Jun 27, 2012
	ANCHEN PHARMS	100MG	A200491	001	Jun 27, 2012
		200MG	A200491	002	Jun 27, 2012
		300MG	A200491	003	Jun 27, 2012
	AUROBINDO PHARMA LTD	100MG	A204421	001	Oct 20, 2015
		200MG	A204421	002	Oct 20, 2015
		300MG	A204421	003	Oct 20, 2015
	MYLAN	100MG	A205257	001	Dec 22, 2015
		200MG	A205257	002	Dec 22, 2015
		300MG	A205257	003	Dec 22, 2015
	STRIDES PHARMA	100MG	A078783	001	Nov 13, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRAMADOL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

TRAMADOL HYDROCHLORIDE

	200MG	A078783 002	Nov 13, 2009
	300MG	A078783 003	Sep 20, 2011
SUN PHARM	200MG	A091607 002	Dec 30, 2011
ULTRAM ER			
+ VALEANT PHARMS	100MG **	N021692 001	Sep 08, 2005
+	200MG **	N021692 002	Sep 08, 2005
+	300MG **	N021692 003	Sep 08, 2005

TABLET, ORALLY DISINTEGRATING;ORAL

RYBIX ODT

SHIONOGI INC	50MG	N021693 001	May 05, 2005
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TRAMETINIB DIMETHYL SULFOXIDE

TABLET;ORAL

MEKINIST

+ NOVARTIS	EQ 1MG	N204114 002	May 29, 2013
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TRANDOLAPRIL

TABLET;ORAL

MAVIK

+ ABBVIE	1MG **	N020528 001	Apr 26, 1996
+	2MG **	N020528 002	Apr 26, 1996
+	4MG **	N020528 003	Apr 26, 1996

TRANDOLAPRIL

CHARTWELL MOLECULAR	1MG	A077307 002	Jun 12, 2007
	2MG	A077307 001	Jun 12, 2007
	4MG	A077307 003	Jun 12, 2007
DR REDDYS LABS LTD	1MG	A078493 001	Aug 25, 2008
	2MG	A078493 002	Aug 25, 2008
	4MG	A078493 003	Aug 25, 2008
EPIC PHARMA LLC	1MG	A077256 001	Jun 12, 2007
	2MG	A077256 002	Jun 12, 2007
	4MG	A077256 003	Jun 12, 2007
INVAGEN PHARMS	1MG	A078320 001	Jun 12, 2007
	2MG	A078320 002	Jun 12, 2007
	4MG	A078320 003	Jun 12, 2007
MYLAN	1MG	A078346 001	Apr 28, 2008
	2MG	A078346 002	Apr 28, 2008
	4MG	A078346 003	Apr 28, 2008
TEVA PHARMS	1MG	A077489 001	Dec 12, 2006
	2MG	A077489 002	Dec 12, 2006
	4MG	A077489 003	Dec 12, 2006
WATSON LABS	1MG	A077805 001	Jun 12, 2007
	2MG	A077805 002	Jun 12, 2007
	4MG	A077805 003	Jun 12, 2007

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

TARKA

+ ABBVIE	1MG;240MG **	N020591 003	Oct 22, 1996
+	2MG;180MG	N020591 001	Oct 22, 1996
+	2MG;240MG	N020591 004	Oct 22, 1996
+	4MG;240MG	N020591 002	Oct 22, 1996

TRANEXAMIC ACID

INJECTABLE;INJECTION

TRANEXAMIC ACID

AKORN	100MG/ML	A206594 001	Sep 28, 2017
CHARTWELL RX	100MG/ML	A202755 001	Feb 25, 2016
RISING	100MG/ML	A206634 001	Jun 09, 2016
ZYDUS PHARMS	100MG/ML	A205228 001	Jul 17, 2017

TABLET;ORAL

CYKLOKAPRON

PHARMACIA AND UPJOHN	500MG	N019280 001	Dec 30, 1986
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TRANEXAMIC ACID

APOTEX	650MG	A202286 001	Jan 27, 2014
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRAVOPROST

SOLUTION/DROPS;OPHTHALMIC

IZBA

+ NOVARTIS 0.003% ** N204822 001 May 15, 2014

TRAVATAN

+ ALCON PHARMS LTD 0.004% ** N021257 001 Mar 16, 2001

TRAZODONE HYDROCHLORIDE

TABLET;ORAL

DESYREL

+ PRAGMA 50MG ** N018207 001

+ 100MG ** N018207 002

+ 150MG ** N018207 003 Mar 25, 1985

+ 300MG ** N018207 004 Nov 07, 1988

TRAZODONE HYDROCHLORIDE

ALVOGEN

50MG A071636 001 Apr 18, 1988

100MG A071514 001 Apr 18, 1988

AM THERAP

50MG A071139 001 Oct 29, 1986

100MG A071140 001 Oct 29, 1986

FOSUN PHARMA

100MG A072483 001 Apr 30, 1990

MYLAN

50MG A071405 001 Feb 27, 1991

100MG A071406 001 Feb 27, 1991

MYLAN PHARMS INC

50MG A090514 001 Jun 02, 2009

100MG A090514 002 Jun 02, 2009

150MG A090514 003 Jun 02, 2009

300MG A090514 004 Jun 02, 2009

QUANTUM PHARMICS

100MG A070921 001 Dec 01, 1986

RISING

50MG A072484 001 Apr 30, 1990

TEVA

150MG A074357 001 Apr 30, 1997

USL PHARMA

50MG A070491 001 Apr 29, 1987

100MG A070492 001 Apr 29, 1987

WATSON LABS

50MG A070857 001 Oct 10, 1986

50MG A071112 001 Nov 17, 1986

100MG A070858 001 Oct 10, 1986

100MG A071113 001 Nov 17, 1986

TRIALODINE

QUANTUM PHARMICS

50MG A070942 001 Dec 01, 1986

TABLET, EXTENDED RELEASE;ORAL

OLEPTRO

+ ANGELINI PHARMA 150MG ** N022411 001 Feb 02, 2010

+ 300MG ** N022411 002 Feb 02, 2010

TREPROSTINIL

SOLUTION;INTRAVENOUS, SUBCUTANEOUS

REMODULIN

UNITED THERAP 20MG/20ML (1MG/ML) N208276 001 Jul 30, 2018

50MG/20ML (2.5MG/ML) N208276 002 Jul 30, 2018

100MG/20ML (5MG/ML) N208276 003 Jul 30, 2018

200MG/20ML (10MG/ML) N208276 004 Jul 30, 2018

TRETINOIN

CAPSULE;ORAL

VESANOID

+ CHEPLAPHARM 10MG ** N020438 001 Nov 22, 1995

CREAM;TOPICAL

RENOVA

+ VALEANT PHARMS NORTH 0.05% ** N019963 001 Dec 29, 1995

TRETINOIN

ALLERGAN 0.0375% A090098 001 Mar 22, 2010

0.075% A202209 001 Oct 11, 2012

ZO SKIN HEALTH

0.05% A076498 001 Sep 15, 2005

GEL;TOPICAL

AVITA

MYLAN 0.025% N020400 001 Jan 29, 1998

TRETINOIN

MYLAN 0.04% A202567 001 Jul 17, 2013

0.1% A202026 001 Jul 17, 2013

SOLUTION;TOPICAL

RETIN-A

+ VALEANT INTL 0.05% N016921 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRETINOIN

SOLUTION; TOPICAL

TRETINOIN

TEVA PHARMS	0.05%	A074873 001	Jun 19, 1998
WOCKHARDT	0.05%	A075260 001	Jan 25, 1999

SWAB; TOPICAL

RETIN-A

VALEANT INTL	0.05%	N016921 002	
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TRIAMCINOLONE

TABLET; ORAL

ARISTOCORT

ASTELLAS

1MG	N011161 009
2MG	N011161 004
4MG	N011161 007
8MG	N011161 011
16MG	N011161 010

KENACORT

DELCOR ASSET CORP

1MG	N011283 003
2MG	N011283 008
4MG	N011283 006
8MG	N011283 010

TRIAMCINOLONE

BARR

2MG	A084286 001
2MG	A084318 001
4MG	A084267 001
4MG	A084319 001
8MG	A084268 001
8MG	A084320 001

IMPAX LABS

4MG	A084340 001
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IVAX SUB TEVA PHARMS

4MG	A083750 001
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MYLAN

2MG	A084406 001
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PUREPAC PHARM

2MG	A084020 002
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4MG	A084020 003
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ROXANE

2MG	A084708 001
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4MG	A084709 001
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8MG	A084707 001
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SANDOZ

4MG	A085601 001
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TEVA

4MG	A084775 001
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WATSON LABS

4MG	A084270 001
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4MG	A085834 001
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TRIAMCINOLONE ACETONIDE

AEROSOL, METERED; INHALATION

AZMACORT

ABBVIE

0.1MG/INH	N018117 001	Apr 23, 1982
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AEROSOL, METERED; NASAL

NASACORT

CHATTEM SANOFI

0.055MG/INH	N019798 001	Jul 11, 1991
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CREAM; TOPICAL

ARISTOCORT

ASTELLAS

0.025%	A083017 003
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0.1%	A083016 004
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+

0.5%	A083015 002
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ARISTOCORT A

ASTELLAS

0.025%	A083017 004
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0.025%	A088818 001	Oct 16, 1984
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0.1%	A083016 005
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0.1%	A088819 001	Oct 16, 1984
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0.5%	A083015 003
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0.5%	A088820 001	Oct 16, 1984
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FLUTEX

IVAX PHARMS

0.025%	A085539 001
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0.1%	A085539 002
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0.5%	A085539 003
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KENALOG

+ DELCOR ASSET CORP

0.5%	A083943 001
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KENALOG-H

DELCOR ASSET CORP

0.1%	A086240 001
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

TRIA CET

TEVA	0.025%	A084908 001
	0.1%	A084908 002
	0.5%	A084908 003

TRIA CORT

SOLVAY	0.1%	A087113 001
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TRIAMCINOLONE ACETONIDE

ACTAVIS MID ATLANTIC	0.1%	A087798 001	Jun 04, 1982
ALPHARMA US PHARMS	0.025%	A087797 001	Jun 07, 1982
AMBIX	0.025%	A087932 001	May 09, 1983
MORTON GROVE	0.025%	A088094 001	Sep 01, 1983
	0.1%	A088095 001	Sep 01, 1983
	0.5%	A088096 001	Sep 01, 1983
+ MYLAN	0.025% **	N011601 003	
+	0.1% **	N011601 006	
PHARMADERM	0.025%	A087990 001	Jul 07, 1983
	0.1%	A087991 001	Jul 07, 1983
	0.5%	A087992 001	Jul 07, 1983
PHARMAFAIR	0.025%	A087921 001	Aug 10, 1982
	0.1%	A087912 001	Aug 10, 1982
	0.5%	A087922 001	Aug 10, 1982
TARO	0.025%	A040038 001	Oct 26, 1994
	0.025%	A086277 001	
	0.1%	A086276 001	
	0.5%	A086275 001	
TOPIDERM	0.025%	A089274 001	Feb 21, 1989
	0.1%	A089275 001	Feb 21, 1989
	0.5%	A089276 001	Feb 21, 1989

TRIA TEX

IVAX PHARMS	0.025%	A087430 001	Nov 01, 1988
	0.1%	A087429 001	Nov 01, 1988
	0.5%	A087428 001	Nov 01, 1988

TRYMEX

SAVAGE LABS	0.025%	A088196 001	Mar 25, 1983
	0.1%	A088197 001	Mar 25, 1983
	0.5%	A088198 001	Mar 25, 1983

GEL;TOPICAL

ARISTOGEL

ASTELLAS	0.1%	A083380 001
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INJECTABLE;INJECTION

TRIAMCINOLONE ACETONIDE

PARNELL	3MG/ML	N019503 001	Oct 16, 1987
SANDOZ	10MG/ML	A090166 001	May 27, 2009
	40MG/ML	A090164 001	Jun 01, 2009
WATSON LABS	40MG/ML	A085825 001	

INJECTABLE;INTRA-ARTICULAR, INTRAMUSCULAR, INTRAVITREAL

TRIVARIS

+ ALLERGAN	8MG/0.1ML (8MG/0.1ML) **	N022220 001	Jun 16, 2008
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LOTION;TOPICAL

KENALOG

DEL COR ASSET CORP	0.025% **	A084343 001
+	0.025% **	N011602 003
	0.1% **	A084343 002
+	0.1% **	N011602 001

TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS	0.025%	A087191 001	Sep 08, 1982
	0.1%	A087192 001	Sep 08, 1982
PAI HOLDINGS PHARM	0.025%	A204608 001	Jul 07, 2016
	0.1%	A204606 001	Jul 07, 2016

OINTMENT;TOPICAL

ARISTOCORT

ASTELLAS	0.1%	A080750 004
+	0.5% **	A080745 002

ARISTOCORT A

ASTELLAS	0.1%	A080750 003	
	0.1%	A088780 001	Oct 01, 1984
+	0.5% **	A080745 003	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

ARISTOCORT A					
	0.5%		A088781	001	Oct 05, 1984
FLUTEX					
IVAX PHARMS	0.025%		A087375	001	Nov 01, 1988
	0.1%		A087377	001	Nov 01, 1988
	0.5%		A087376	001	Nov 01, 1988
KENALOG					
+ DELCOR ASSET CORP	0.5% **		A083944	001	
TRIAMCINOLONE ACETONIDE					
ACTAVIS MID ATLANTIC	0.1%		A087799	001	Jun 07, 1982
ALPHARMA US PHARMS	0.5%		A089913	001	Dec 23, 1988
MORTON GROVE	0.025%		A088090	001	Sep 01, 1983
	0.1%		A088091	001	Sep 01, 1983
	0.5%		A088092	001	Sep 01, 1983
+ MYLAN	0.025% **		N011600	003	
+ PAI HOLDINGS PHARM	0.5%		A208590	001	Mar 03, 2017
PHARMADERM	0.025%		A088692	001	Aug 02, 1984
	0.1%		A088690	001	Aug 02, 1984
STRIDES PHARMA	0.1%		A211315	001	Mar 18, 2020
	0.5%		A211315	002	Mar 18, 2020
TARO	0.025%		A040040	001	Sep 30, 1994
	0.025%		A040374	001	Jun 05, 2001
	0.1%		A087902	001	Dec 27, 1982
	0.5%		A040386	001	Jun 05, 2001
TRIANEX					
CMP PHARMA INC	0.05%		A089595	001	Mar 23, 1995
TRYMEX					
SAVAGE LABS	0.025%		A088693	001	Aug 02, 1984
	0.1%		A088691	001	Aug 02, 1984
PASTE; DENTAL					
KENALOG IN ORABASE					
+ DELCOR ASSET CORP	0.1% **		N012097	001	
ORALONE					
TARO	0.1%		A071383	001	Jul 06, 1987
SPRAY; TOPICAL					
KENALOG					
+ SUN PHARM INDS INC	0.147MG/GM		N012104	001	
SPRAY, METERED; NASAL					
ALLERNAZE					
LUPIN ATLANTIS	0.05MG/SPRAY		N020120	001	Feb 04, 2000
NASACORT HFA					
SANOFI AVENTIS US	0.055MG/SPRAY		N020784	001	Apr 07, 2004
TRIAMCINOLONE ACETONIDE					
PERRIGO PHARMA INTL	0.055MG/SPRAY		A078104	001	Jul 30, 2009

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

ARISTOCORT					
FOSUN PHARMA	25MG/ML		N011685	003	
+ WATSON LABS	40MG/ML **		N012802	001	
TRIAMCINOLONE DIACETATE					
AKORN	25MG/ML		A085122	001	
	40MG/ML		A086394	001	
WATSON LABS	40MG/ML		A084072	001	
	40MG/ML		A085529	001	
SYRUP; ORAL					
ARISTOCORT					
ASTELLAS	2MG/5ML		N011960	004	
KENACORT					
DELCOR ASSET CORP	EQ 4MG BASE/5ML		N012515	001	

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE HEXACETONIDE

INJECTABLE; INJECTION

ARISTOSPAN

+ SANDOZ

5MG/ML **

N016466 001

+

20MG/ML **

N016466 002

TRIAMTERENE

CAPSULE; ORAL

TRIAMTERENE

EYWA

50MG

A214768 001 Jul 06, 2022

100MG

A214768 002 Jul 06, 2022

TRIAZOLAM

TABLET; ORAL

HALCION

PFIZER

0.5MG

N017892 002 Nov 15, 1982

TRIAZOLAM

MYLAN PHARMS INC

0.125MG

A074031 001 Mar 25, 1994

WATSON LABS

0.25MG

A074031 002 Mar 25, 1994

WATSON LABS

0.125MG

A074445 001 Oct 20, 1995

WATSON LABS

0.25MG

A074445 002 Oct 20, 1995

TRICHLORMETHIAZIDE

TABLET; ORAL

METAHYDRIN

SANOFI AVENTIS US

2MG

N012594 001 Jun 16, 1988

SANOFI AVENTIS US

4MG

N012594 002 Jun 16, 1988

NAQUA

SCHERING

2MG

N012265 001

SCHERING

4MG

N012265 002

TRICHLOREX

LANNETT

4MG

A083436 001

LANNETT

4MG

A085630 001

TRICHLORMAS

MAST MM

4MG

A086259 001

TRICHLORMETHIAZIDE

CHARTWELL RX

4MG

A085568 001

IMPAX LABS

4MG

A083967 001

PAR PHARM

2MG

A087007 001

PAR PHARM

4MG

A087005 001

SANDOZ

4MG

A086171 001

WATSON LABS

2MG

A083847 001

WATSON LABS

2MG

A086458 001

WATSON LABS

4MG

A083462 001

WATSON LABS

4MG

A083855 001

WATSON LABS

4MG

A085962 001

TRICLOFOS SODIUM

SOLUTION; ORAL

TRICLOS

SANOFI AVENTIS US

1.5GM/15ML

N016830 001

TABLET; ORAL

TRICLOS

SANOFI AVENTIS US

750MG

N016809 002

TRIDIHETHYL CHLORIDE

INJECTABLE; INJECTION

PATHILON

LEDERLE

10MG/ML

N009729 001

TABLET; ORAL

PATHILON

LEDERLE

25MG

N009489 005

TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

CLOVIQUE

CHARTWELL RX

250MG

A209731 001 Oct 21, 2019

TRIENTINE HYDROCHLORIDE

ACCORD HLTHCARE

250MG

A212929 001 Aug 30, 2021

AMNEAL

250MG

A210619 001 Feb 08, 2019

CHARTWELL RX

250MG

A209415 001 Sep 16, 2019

LUPIN LTD

250MG

A211637 001 May 21, 2020

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

STELAZINE

+ GLAXOSMITHKLINE EQ 10MG BASE/ML ** N011552 006

TRIFLUOPERAZINE HYDROCHLORIDE

FOSUN PHARMA EQ 10MG BASE/ML A085787 001 Apr 15, 1982

WOCKHARDT EQ 10MG BASE/ML A088143 001 Jul 26, 1983

INJECTABLE; INJECTION

STELAZINE

+ GLAXOSMITHKLINE EQ 2MG BASE/ML ** N011552 005

TABLET; ORAL

STELAZINE

+ GLAXOSMITHKLINE EQ 1MG BASE ** N011552 001

+ EQ 2MG BASE ** N011552 002

+ EQ 5MG BASE ** N011552 003

+ EQ 10MG BASE ** N011552 004

TRIFLUOPERAZINE HYDROCHLORIDE

ATHEM EQ 1MG BASE A040153 001 Oct 25, 1996

EQ 2MG BASE A040153 002 Oct 25, 1996

EQ 5MG BASE A040153 003 Oct 25, 1996

EQ 10MG BASE A040153 004 Oct 25, 1996

DURAMED PHARMS BARR EQ 1MG BASE A088967 001 Apr 23, 1985

EQ 2MG BASE A088968 001 Apr 23, 1985

EQ 5MG BASE A088969 001 Apr 23, 1985

EQ 10MG BASE A088970 001 Apr 23, 1985

IVAX PHARMS EQ 1MG BASE A087612 001 Nov 19, 1982

EQ 2MG BASE A087613 001 Nov 19, 1982

EQ 5MG BASE A087328 001 Nov 19, 1982

EQ 10MG BASE A087614 001 Nov 19, 1982

WATSON LABS EQ 1MG BASE A085975 001 Jun 23, 1988

EQ 2MG BASE A085976 001 Jun 23, 1988

EQ 5MG BASE A085973 001 Jun 23, 1988

EQ 10MG BASE A088710 001 Jun 23, 1988

TRIFLUPROMAZINE

SUSPENSION; ORAL

VESPRIN

APOTHECON EQ 50MG HYDROCHLORIDE/5ML N011491 004

TRIFLUPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

VESPRIN

APOTHECON 3MG/ML N011325 005

10MG/ML N011325 004

20MG/ML N011325 001

TABLET; ORAL

VESPRIN

BRISTOL MYERS SQUIBB 10MG N011123 001

25MG N011123 002

50MG N011123 003

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

HIKMA 1% A205438 001 Jul 28, 2017

TRIHEXYPHENIDYL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

ARTANE

LEDERLE 5MG N006773 010

5MG N012947 001

ELIXIR; ORAL

ARTANE

LEDERLE 2MG/5ML N006773 009

TRIHEXYPHENIDYL HYDROCHLORIDE

PHARM VENTURES 2MG/5ML A089514 001 Apr 07, 1989

TABLET; ORAL

ARTANE

+ LEDERLE 2MG ** N006773 005

+ 5MG ** N006773 003

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIHEXYPHENIDYL HYDROCHLORIDE

TABLET; ORAL

TREMIM

SCHERING	2MG	A080381	001
	5MG	A080381	003

TRIHEXYPHENIDYL HYDROCHLORIDE

HIKMA	2MG	A040337	002	Feb 16, 2000
	5MG	A040337	001	Feb 16, 2000
NYLOS	5MG	A085622	001	
VANGARD	2MG	A088035	001	Jul 30, 1982
WATSON LABS	2MG	A040184	001	Feb 06, 1998
	2MG	A085117	001	
	5MG	A040184	002	Feb 06, 1998
	5MG	A085105	001	

TRILOSTANE

CAPSULE; ORAL

MODRASTANE

BIOENVISION	30MG	N018719	002	Dec 31, 1984
	60MG	N018719	001	Dec 31, 1984

TRIMEPRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 5MG BASE	N011316	004	
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SYRUP; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE/5ML	N011316	003	
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TRIMEPRAZINE TARTRATE

ALPHARMA US PHARMS	EQ 2.5MG BASE/5ML	A085015	001	Feb 18, 1982
MORTON GROVE	EQ 2.5MG BASE/5ML	A088285	001	Apr 11, 1985

TABLET; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE	N011316	001	
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TRIMETHADIONE

CAPSULE; ORAL

TRIDIONE

ABBVIE	300MG	N005856	005	
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SOLUTION; ORAL

TRIDIONE

ABBVIE	200MG/5ML	N005856	002	
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TABLET; ORAL

TRIDIONE

+ ABBVIE	150MG	N005856	009	
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TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION

ARFONAD

ROCHE	50MG/ML	N008983	001	
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TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TIGAN

+ KING PHARMS LLC	300MG **	N017531	006	Dec 13, 2001
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INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

AM REGENT	100MG/ML	A091330	001	Mar 08, 2011
HOSPIRA	100MG/ML	A088804	001	Apr 03, 1987
SMITH AND NEPHEW	100MG/ML	A088960	001	Apr 04, 1986
	100MG/ML	A089043	001	Apr 04, 1986
SOLOPAK	100MG/ML	A089094	001	Apr 04, 1986
WATSON LABS	100MG/ML	A086577	001	Oct 19, 1982
	100MG/ML	A087939	001	Dec 28, 1982

TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE

AM REGENT	100MG/ML	A091329	001	Mar 08, 2011
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIMETHOPRIM

TABLET; ORAL

PROLOPRIM

MONARCH PHARMS	100MG	N017943 001	
	200MG	N017943 003	Jul 14, 1982

TRIMETHOPRIM

SUN PHARM INDUSTRIES	100MG	A070494 001	Jan 22, 1986
	200MG	A070495 001	Sep 24, 1986

TEVA	200MG **	A071259 001	Jun 18, 1987
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TRIMPEX

ROCHE	100MG	N017952 001	
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TRIMPEX 200

ROCHE	200MG	N017952 002	Nov 09, 1982
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TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

ALLEGIS	EQ 25MG BASE/5ML	N074374 001	Jun 23, 1995
+	EQ 50MG BASE/5ML	N074973 001	Jan 24, 2000

TRIMETREXATE GLUCURONATE

INJECTABLE; INJECTION

NEUTREXIN

MEDIMMUNE ONCOLOGY	EQ 25MG BASE/VIAL	N020326 001	Dec 17, 1993
	EQ 200MG BASE/VIAL	N020326 002	Jul 31, 1998

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

SURMONTIL

+	ODYSSEY PHARMS	EQ 25MG BASE **	N016792 001
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+		EQ 50MG BASE **	N016792 002
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+		EQ 100MG BASE **	N016792 003 Sep 15, 1982
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TRIMIPRAMINE MALEATE

USL PHARMA	EQ 25MG BASE	A071283 001	Dec 08, 1987
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	EQ 50MG BASE	A071284 001	Dec 08, 1987
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	EQ 100MG BASE	A071285 001	Dec 08, 1987
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TRIOXSALEN

TABLET; ORAL

TRISORALEN

VALEANT PHARM INTL	5MG	N012697 001	
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TRIPLENNAMINE CITRATE

ELIXIR; ORAL

PBZ

NOVARTIS	EQ 25MG HYDROCHLORIDE/5ML	N005914 004	
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TRIPLENNAMINE HYDROCHLORIDE

TABLET; ORAL

PBZ

NOVARTIS	25MG	A083149 001	
	50MG	N005914 002	

TRIPLENNAMINE HYDROCHLORIDE

ANABOLIC	50MG	A083037 001	
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BARR	50MG	A080744 001	
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HEATHER	50MG	A083989 001	
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IMPAX LABS	50MG	A080785 001	
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LANNETT	50MG	A083557 001	
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NYLOS	50MG	A085412 001	
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PARKE DAVIS	25MG	A083625 001	
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	50MG	A083626 001	
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WATSON LABS	50MG	A080713 001	
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	50MG	A080790 001	
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	50MG	A085188 001	
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TABLET, EXTENDED RELEASE; ORAL

PBZ-SR

NOVARTIS	50MG	N010533 002	
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	100MG	N010533 001	
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIPLE SULFA (SULFABENZAMIDE;SULFACETAMIDE;SULFATHIAZOLE)

CREAM;VAGINAL

GYNE-SULF

COSETTE 3.7%;2.86%;3.42% A088607 001 Jun 09, 1986

SULTRIN

ORTHO MCNEIL PHARM 3.7%;2.86%;3.42% N005794 001

TRIPLE SULFA

ALPHARMA US PHARMS 3.7%;2.86%;3.42% A087864 001 Sep 01, 1982

FOUGERA 3.7%;2.86%;3.42% A086424 001

PADAGIS US 3.7%;2.86%;3.42% A087285 001 Nov 15, 1982

TRYSUL

SAVAGE LABS 3.7%;2.86%;3.42% A087887 001 Jul 23, 1982

VAGILIA

COSETTE 3.7%;2.86%;3.42% A088821 001 Nov 09, 1987

TABLET;VAGINAL

SULTRIN

ORTHO MCNEIL PHARM 184MG;143.75MG;172.5MG N005794 002

TRIPLE SULFA

FOUGERA 184MG;143.75MG;172.5MG A088463 001 Jan 03, 1985

PHARMADERM 184MG;143.75MG;172.5MG A088462 001 Jan 03, 1985

TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

ACTIDIL

GLAXOSMITHKLINE 1.25MG/5ML N011496 002 Jul 01, 1983

MYIDYL

USL PHARMA 1.25MG/5ML A087963 001 Jan 18, 1983

TRIPROLIDINE HYDROCHLORIDE

ALPHARMA US PHARMS 1.25MG/5ML A085940 001

HALSEY 1.25MG/5ML A088735 001 Jan 17, 1985

PHARM ASSOC 1.25MG/5ML A087514 001 Feb 10, 1982

TABLET;ORAL

ACTIDIL

GLAXOSMITHKLINE 2.5MG N011110 002 Jul 01, 1983

TRIPROLIDINE HYDROCHLORIDE

VITARINE 2.5MG A085610 001

WATSON LABS 2.5MG A085094 001

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

SUSPENSION;ORAL

LANTRISUL

LANNETT 167MG/5ML;167MG/5ML;167MG/5ML A080123 002

NEOTRIZINE

LILLY 167MG/5ML;167MG/5ML;167MG/5ML N006317 012

SULFALOID

FOREST PHARMS 167MG/5ML;167MG/5ML;167MG/5ML A080100 001

SULFOSE

WYETH AYERST 167MG/5ML;167MG/5ML;167MG/5ML A080013 002

TERFONYL

BRISTOL MYERS SQUIBB 167MG/5ML;167MG/5ML;167MG/5ML N006904 002

TRIPLE SULFA

ALPHARMA US PHARMS 167MG/5ML;167MG/5ML;167MG/5ML A080280 001

TRIPLE SULFAS

LEDERLE 167MG/5ML;167MG/5ML;167MG/5ML N006920 003

TABLET;ORAL

NEOTRIZINE

LILLY 167MG;167MG;167MG N006317 011

SULFA-TRIPLE #2

IMPAX LABS 167MG;167MG;167MG A080079 001

SULFALOID

FOREST PHARMS 167MG;167MG;167MG A080099 001

SULFOSE

WYETH AYERST 167MG;167MG;167MG A080013 001

TERFONYL

BRISTOL MYERS SQUIBB 167MG;167MG;167MG N006904 001

TRIPLE SULFA

PUREPAC PHARM 167MG;167MG;167MG A080086 001

TRIPLE SULFAS

LEDERLE 167MG;167MG;167MG N006920 002

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

TABLET;ORAL

TRIPLE SULFOID

PAL PAK

167MG;167MG;167MG

A080094 001

TROGLITAZONE

TABLET;ORAL

PRELAY

SANKYO

200MG

N020719 001 Jan 29, 1997

300MG

N020719 003 Aug 04, 1997

400MG

N020719 002 Jan 29, 1997

REZULIN

PFIZER PHARMS

200MG

N020720 001 Jan 29, 1997

300MG

N020720 003 Aug 04, 1997

400MG

N020720 002 Jan 29, 1997

TROLAMINE POLYPEPTIDE OLEATE CONDENSATE

SOLUTION/DROPS;OTIC

CERUMENEX

PHARM RES ASSOC

10%

N011340 002

TROLEANDOMYCIN

CAPSULE;ORAL

TAO

PFIZER

EQ 250MG BASE

N050336 002

SUSPENSION;ORAL

TAO

PFIZER

EQ 125MG BASE/5ML

N050332 001

TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

MYDRIACYL

ALCON

0.5% **

N012111 002

1% **

N012111 004

MYDRIAFAIR

PHARMAFAIR

0.5%

A088274 001 Sep 16, 1983

1%

A088230 001 Sep 16, 1983

TROPICAMIDE

AKORN

1%

A088447 001 Aug 28, 1985

ALCON PHARMS LTD

1%

A089172 001 Dec 28, 1990

MIZA PHARMS USA

0.5%

A087636 001 Jul 30, 1982

1%

A087637 001 Aug 09, 1982

WATSON LABS

0.5%

A089171 001 Dec 28, 1990

TROSPIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

SANCTURA XR

+ ALLERGAN

60MG **

N022103 001 Aug 03, 2007

TROSPIUM CHLORIDE

UPSHER SMITH LABS

60MG

A091635 001 Apr 29, 2015

TABLET;ORAL

SANCTURA

+ ALLERGAN

20MG **

N021595 001 May 28, 2004

TROVAFLOXACIN MESYLATE

TABLET;ORAL

TROVAN

PFIZER

EQ 100MG BASE

N020759 001 Dec 18, 1997

EQ 200MG BASE

N020759 002 Dec 18, 1997

TRYPAN BLUE

SOLUTION;OPHTHALMIC

MEMBRANEBLUE

+ DORC

0.15%

N022278 001 Feb 20, 2009

TUBOCURARINE CHLORIDE

INJECTABLE;INJECTION

TUBOCURARINE CHLORIDE

BRISTOL MYERS SQUIBB

3MG/ML

N005657 001

HOSPIRA

3MG/ML

N006095 001

LILLY

3MG/ML

N006325 001

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TYROPANOATE SODIUM

CAPSULE;ORAL

BILOPAQUE

GE HEALTHCARE

750MG

N013731 001

UMBRALISIB TOSYLATE

TABLET;ORAL

UKONIQ

+ TG THERAPS

EQ 200MG BASE

N213176 001 Feb 05, 2021

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS;OPHTHALMIC

RESCULA

+ SUCAMPO PHARMA LLC

0.15% **

N021214 001 Aug 03, 2000

URACIL MUSTARD

CAPSULE;ORAL

URACIL MUSTARD

SHIRE

1MG

N012892 001

UREA

INJECTABLE;INJECTION

STERILE UREA

HOSPIRA

40GM/VIAL

N017698 001

UREAPHIL

HOSPIRA

40GM/VIAL

N012154 001

UREA C-13

FOR SOLUTION;ORAL

BREATHTEK UBT FOR H-PYLORI

OTSUKA AMERICA

EQ 75MG/POUCH

N020586 002 May 10, 2001

HELICOSOL

METABOLIC SOLUTIONS

125MG/VIAL

N021092 001 Dec 17, 1999

MERETEK UBT KIT (W/ PRANACTIN)

OTSUKA AMERICA

125MG/VIAL

N020586 001 Sep 17, 1996

PYLORI-CHEK BREATH TEST

DXS DEVICES

100MG/VIAL

N020900 001 Feb 04, 1999

UREA, C-14

CAPSULE;ORAL

PYTEST KIT

+ AVENT

1uCi

N020617 002 May 09, 1997

URSODIOL

CAPSULE;ORAL

ACTIGALL

TEVA BRANDED PHARM

150MG

N019594 001 Dec 31, 1987

URSODIOL

IMPAX LABS INC

300MG

A077895 001 Jul 27, 2006

MYLAN

300MG

A090530 001 Feb 17, 2010

TEVA PHARMS

300MG

A075592 001 May 25, 2000

TABLET;ORAL

URSODIOL

IMPAX LABS INC

250MG

A200826 001 Dec 23, 2011

500MG

A200826 002 Dec 23, 2011

TEVA PHARMS USA

250MG

A079184 001 May 13, 2009

500MG

A079184 002 May 13, 2009

VALACYCLOVIR HYDROCHLORIDE

TABLET;ORAL

VALACYCLOVIR HYDROCHLORIDE

HIKMA

EQ 500MG BASE

A078656 001 May 24, 2010

EQ 1GM BASE

A078656 002 May 24, 2010

MYLAN

EQ 500MG BASE

A078070 001 May 24, 2010

EQ 1GM BASE

A078070 002 May 24, 2010

TEVA PHARMS

EQ 500MG BASE

A077655 001 May 24, 2010

EQ 1GM BASE

A077655 002 May 24, 2010

WATSON LABS INC

EQ 500MG BASE

A090370 001 Mar 16, 2011

EQ 1GM BASE

A090370 002 Mar 16, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VALDECOXIB

TABLET; ORAL

BEXTRA

GD SEARLE	10MG	N021341 002	Nov 16, 2001
	20MG	N021341 003	Nov 16, 2001

VALGANCICLOVIR HYDROCHLORIDE

TABLET; ORAL

VALGANCICLOVIR HYDROCHLORIDE

CIPLA	EQ 450MG BASE	A209672 001	Nov 09, 2018
MYLAN	EQ 450MG BASE	A205151 001	Mar 03, 2021

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPAICON

+ ABBVIE

EQ 100MG BASE/ML ** N020593 001 Dec 30, 1996

VALPROATE SODIUM

MYLAN

EQ 100MG BASE/ML A208120 001 Dec 22, 2021

VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

+ ABBVIE

250MG ** N018081 001

VALPROIC ACID

PAR PHARM

250MG A070431 001 Feb 28, 1986

SCHERER RP

250MG A070195 001 Jul 02, 1987

UPSHER SMITH LABS

250MG A070631 001 Jun 11, 1987

CAPSULE, DELAYED RELEASE; ORAL

STAVZOR

+ BIONPHARMA

125MG ** N022152 001 Jul 29, 2008

+

250MG ** N022152 002 Jul 29, 2008

+

500MG ** N022152 003 Jul 29, 2008

SYRUP; ORAL

DEPAKENE

+ ABBVIE

250MG/5ML ** N018082 001

VALPROIC ACID

LANNETT CO INC

250MG/5ML A077960 001 Oct 13, 2006

NOSTRUM LABS INC

250MG/5ML A077105 001 Jul 29, 2005

PHARMOBEDIANT CNSLTG

250MG/5ML A070868 001 Jul 01, 1986

VALSARTAN

CAPSULE; ORAL

DIOVAN

NOVARTIS

80MG N020665 001 Dec 23, 1996

160MG N020665 002 Dec 23, 1996

SOLUTION; ORAL

PREXXARTAN

+ CARMEL BIOSCIENCES

20MG/5ML ** N209139 001 Dec 19, 2017

+

80MG/20ML ** N209139 002 Dec 19, 2017

TABLET; ORAL

VALSARTAN

IVAX PHARMS

40MG A077530 001 Jan 04, 2016

80MG A077530 002 Jan 04, 2016

160MG A077530 003 Jan 04, 2016

320MG A077530 004 Jan 04, 2016

TORRENT

40MG A202728 001 Jan 05, 2015

80MG A202728 002 Jan 05, 2015

160MG A202728 003 Jan 05, 2015

320MG A202728 004 Jan 05, 2015

UNICHEM

40MG A209261 001 May 04, 2018

80MG A209261 002 May 04, 2018

160MG A209261 003 May 04, 2018

320MG A209261 004 May 04, 2018

WATSON LABS INC

40MG A090642 001 Jan 05, 2015

80MG A090642 002 Jan 05, 2015

160MG A090642 003 Jan 05, 2015

320MG A090642 004 Jan 05, 2015

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VANCOMYCINSOLUTION; INTRAVENOUS, ORAL
VANCOMYCIN

+ XELLIA PHARMS APS 5GM/100ML (50MG/ML) N213895 001 Aug 26, 2021

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOMYCIN HYDROCHLORIDE

FRESENIUS KABI USA EQ 125MG BASE A065453 001 Jun 18, 2012

EQ 250MG BASE A065453 002 Jun 18, 2012

WATSON LABS EQ 125MG BASE A065510 001 Apr 09, 2012

EQ 250MG BASE A065510 002 Apr 09, 2012

FOR SOLUTION; ORAL

VANCOCIN HYDROCHLORIDE

ANI PHARMS EQ 500MG BASE/6ML A061667 001

VANCOLED

LEDERLE EQ 250MG BASE/5ML A063321 002 Oct 15, 1993

EQ 500MG BASE/6ML A063321 003 Oct 15, 1993

INJECTABLE; INJECTION

VANCOCIN HYDROCHLORIDE

STERISCIENCE EQ 500MG BASE/VIAL ** A060180 001

EQ 500MG BASE/VIAL A062476 001 Mar 15, 1984

EQ 500MG BASE/VIAL A062716 001 Mar 13, 1987

EQ 500MG BASE/VIAL ** A062812 001 Nov 17, 1987

EQ 1GM BASE/VIAL ** A060180 002 Mar 21, 1986

EQ 1GM BASE/VIAL A062476 002 Mar 21, 1986

EQ 1GM BASE/VIAL A062716 002 Mar 13, 1987

EQ 1GM BASE/VIAL ** A062812 002 Nov 17, 1987

EQ 10GM BASE/VIAL ** A062812 003 Nov 17, 1987

VANCOLED

HIKMA EQ 500MG BASE/VIAL ** A062682 001 Jul 22, 1986

EQ 1GM BASE/VIAL ** A062682 002 Mar 30, 1988

EQ 2GM BASE/VIAL ** A062682 003 May 11, 1988

EQ 5GM BASE/VIAL ** A062682 004 May 11, 1988

EQ 10GM BASE/VIAL ** A062682 005 May 11, 1988

VANCOMYCIN HYDROCHLORIDE

AVET LIFESCIENCES

EQ 500MG BASE/VIAL A202275 001 Oct 31, 2013

EQ 1GM BASE/VIAL A202275 002 Oct 31, 2013

EQ 10GM BASE/VIAL A202464 001 Oct 09, 2013

EQ 5GM BASE/VIAL A202274 001 Oct 31, 2013

HIKMA EQ 500MG BASE/VIAL A062879 001 Aug 02, 1988

EQ 500MG BASE/VIAL A203300 001 Aug 11, 2020

EQ 1GM BASE/VIAL A062879 002 Aug 02, 1988

KNACK EQ 500MG BASE/VIAL A213059 001 Feb 15, 2022

EQ 750MG BASE/VIAL A213059 002 Feb 15, 2022

EQ 1GM BASE/VIAL A213059 003 Feb 15, 2022

MYLAN LABS LTD EQ 10GM BASE/VIAL A091469 001 Jul 01, 2011

SANDOZ EQ 500MG BASE/VIAL A090250 001 Apr 27, 2010

EQ 1GM BASE/VIAL A090250 002 Apr 27, 2010

TEVA PHARMS USA EQ 500MG BASE/VIAL A201251 001 Dec 23, 2015

EQ 1GM BASE/VIAL A201251 002 Dec 23, 2015

EQ 5GM BASE/VIAL A201250 001 Dec 23, 2015

EQ 10GM BASE/VIAL A201250 002 Dec 23, 2015

XELLIA PHARMS APS EQ 500MG BASE/VIAL A091377 001 Sep 09, 2015

EQ 1GM BASE/VIAL A091377 002 Sep 09, 2015

EQ 5GM BASE/VIAL A206243 001 Dec 23, 2015

EQ 10GM BASE/VIAL A206243 002 Dec 23, 2015

VANCOR

PHARMACIA AND UPJOHN EQ 500MG BASE/VIAL A062956 001 Aug 01, 1988

EQ 1GM BASE/VIAL A062956 002 Aug 01, 1988

POWDER; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

+ MYLAN LABS LTD EQ 250MG BASE/VIAL ** N209481 001 Jul 10, 2018

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

+	BAYER HLTHCARE	EQ 2.5MG BASE **	N021400 003	Aug 19, 2003
+		EQ 5MG BASE **	N021400 001	Aug 19, 2003
+		EQ 10MG BASE **	N021400 002	Aug 19, 2003
+		EQ 20MG BASE **	N021400 004	Aug 19, 2003

VARDENAFIL HYDROCHLORIDE

STEVENS J

		EQ 5MG BASE	A210738 001	Oct 31, 2018
		EQ 10MG BASE	A210738 002	Oct 31, 2018
		EQ 20MG BASE	A210738 003	Oct 31, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

STAXYN

+	BAYER HLTHCARE	10MG	N200179 001	Jun 17, 2010
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VARENICLINE TARTRATE

TABLET; ORAL

CHANTIX

+	PF PRISM CV	EQ 0.5MG BASE **	N021928 001	May 10, 2006
+		EQ 1MG BASE **	N021928 002	May 10, 2006

VASOPRESSIN

SOLUTION; INTRAVENOUS

VASOSTRICT

+	PAR STERILE PRODUCTS	50UNITS/50ML (1UNITS/ML)	N204485 006	Apr 12, 2023
+		60UNITS/100ML (0.6UNITS/ML)	N204485 004	Apr 15, 2020

VASOPRESSIN TANNATE

INJECTABLE; INJECTION

PITRESSIN TANNATE

+	PARKE DAVIS	5PRESSOR UNITS/ML **	N003402 001	
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VECURONIUM BROMIDE

INJECTABLE; INJECTION

NORCURON

+	ORGANON USA INC	10MG/VIAL **	N018776 002	Apr 30, 1984
+		20MG/VIAL **	N018776 003	Jan 03, 1992

VECURONIUM BROMIDE

HIKMA

		10MG/VIAL	A075218 001	Aug 23, 1999
		20MG/VIAL	A075218 002	Aug 23, 1999

HOSPIRA

		4MG/VIAL	A075558 001	Sep 11, 2001
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WATSON LABS

		10MG/VIAL	A074334 001	Aug 31, 1995
		20MG/VIAL	A074334 002	Aug 31, 1995

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

UPJOHN

		EQ 100MG BASE	N020699 003	Oct 20, 1997
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VENLAFAXINE HYDROCHLORIDE

ANCHEN PHARMS

		EQ 37.5MG BASE	A078087 001	Mar 16, 2012
		EQ 75MG BASE	A078087 002	Mar 16, 2012
		EQ 150MG BASE	A078087 003	Mar 16, 2012

MYLAN

		EQ 37.5MG BASE	A078789 001	Jun 01, 2011
		EQ 75MG BASE	A078789 002	Jun 01, 2011
		EQ 150MG BASE	A078789 003	Jun 01, 2011

TORRENT

		EQ 37.5MG BASE	A090899 001	Jun 01, 2011
		EQ 75MG BASE	A090899 002	Jun 01, 2011
		EQ 150MG BASE	A090899 003	Jun 01, 2011

TABLET; ORAL

EFFEXOR

+	WYETH PHARMS INC	EQ 12.5MG BASE **	N020151 001	Dec 28, 1993
+		EQ 25MG BASE **	N020151 002	Dec 28, 1993
+		EQ 37.5MG BASE **	N020151 006	Dec 28, 1993
+		EQ 50MG BASE **	N020151 003	Dec 28, 1993
+		EQ 75MG BASE **	N020151 004	Dec 28, 1993
+		EQ 100MG BASE **	N020151 005	Dec 28, 1993

VENLAFAXINE HYDROCHLORIDE

CHARTWELL RX

		EQ 25MG BASE	A077515 001	Jun 13, 2008
		EQ 37.5MG BASE	A077515 002	Jun 13, 2008
		EQ 50MG BASE	A077515 003	Jun 13, 2008
		EQ 75MG BASE	A077515 004	Jun 13, 2008
		EQ 100MG BASE	A077515 005	Jun 13, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

VENLAFAXINE HYDROCHLORIDE

MYLAN	EQ 25MG BASE	A077166 001	Jun 13, 2008
	EQ 37.5MG BASE	A077166 002	Jun 13, 2008
	EQ 50MG BASE	A077166 003	Jun 13, 2008
	EQ 75MG BASE	A077166 004	Jun 13, 2008
	EQ 100MG BASE	A077166 005	Jun 13, 2008
PLIVA HRVATSKA DOO	EQ 25MG BASE	A078517 001	Jun 13, 2008
	EQ 37.5MG BASE	A078517 002	Jun 13, 2008
	EQ 50MG BASE	A078517 003	Jun 13, 2008
	EQ 75MG BASE	A078517 004	Jun 13, 2008
	EQ 100MG BASE	A078517 005	Jun 13, 2008
PRINSTON INC	EQ 25MG BASE	A090027 001	Aug 04, 2010
	EQ 37.5MG BASE	A090027 002	Aug 04, 2010
	EQ 50MG BASE	A090027 003	Aug 04, 2010
	EQ 75MG BASE	A090027 004	Aug 04, 2010
	EQ 100MG BASE	A090027 005	Aug 04, 2010

TABLET, EXTENDED RELEASE; ORAL

VENLAFAXINE HYDROCHLORIDE

SUNNY	EQ 75MG BASE	A214423 001	Jan 04, 2022
	EQ 150MG BASE	A214423 002	Jan 04, 2022

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

RISING	100MG	A078306 001	Aug 09, 2007
	200MG	A078306 002	Aug 09, 2007
	300MG	A078306 003	Aug 09, 2007

INJECTABLE; INJECTION

CALAN

GD SEARLE LLC	2.5MG/ML	N019038 001	Mar 30, 1984
ISOPTIN			
+ MT ADAMS	2.5MG/ML **	N018485 001	
VERAPAMIL HYDROCHLORIDE			
ABRAXIS PHARM	2.5MG/ML	A070348 001	May 01, 1986
BEDFORD	2.5MG/ML	A072888 001	Jul 28, 1995
HOSPIRA	2.5MG/ML	A070577 001	Feb 02, 1987
	2.5MG/ML	A070739 001	May 06, 1987
	2.5MG/ML	A070740 001	May 06, 1987
INTL MEDICATION	2.5MG/ML	A070451 001	Dec 16, 1985
LUITPOLD	2.5MG/ML	A070225 001	Nov 12, 1985
	2.5MG/ML	A070617 001	Nov 12, 1985
MARSAM PHARMS LLC	2.5MG/ML	A072233 001	Feb 26, 1993
	2.5MG/ML	A073485 001	Sep 27, 1993
SMITH AND NEPHEW	2.5MG/ML	A070696 001	Jul 31, 1987
	2.5MG/ML	A070697 001	Jul 31, 1987
SOLOPAK	2.5MG/ML	A070695 001	Jul 31, 1987

SOLUTION; INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

EXELA PHARMA	10MG/4ML (2.5MG/ML)	N018925 002	Apr 05, 2018
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TABLET; ORAL

CALAN

PFIZER	40MG **	N018817 003	Feb 23, 1988
+	80MG **	N018817 001	Sep 10, 1984
+	120MG **	N018817 002	Sep 10, 1984
	160MG **	N018817 004	Feb 23, 1988

ISOPTIN

+ MT ADAMS	40MG **	N018593 003	Nov 23, 1987
+	80MG **	N018593 001	Mar 08, 1982
+	120MG **	N018593 002	Mar 08, 1982

VERAPAMIL HYDROCHLORIDE

ACTAVIS ELIZABETH	80MG	A071019 001	Sep 24, 1986
	120MG	A070468 001	Sep 24, 1986
CHARTWELL RX	40MG	A073168 001	Jul 31, 1992
	80MG	A071423 001	May 24, 1988
	120MG	A071424 001	May 25, 1988
MUTUAL PHARM	80MG	A070482 001	Sep 24, 1986
	120MG	A070483 001	Sep 24, 1986
PLIVA	40MG	A072751 001	Feb 23, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VERAPAMIL HYDROCHLORIDE

TABLET; ORAL

VERAPAMIL HYDROCHLORIDE

	80MG	A072124	001	Jan 26, 1989
	120MG	A072125	001	Jan 26, 1989
RISING	80MG	A071483	002	Feb 15, 1989
	120MG	A071483	001	Feb 15, 1989
SUN PHARM INDUSTRIES	80MG	A071489	002	Jan 13, 1988
	120MG	A071489	001	Jan 13, 1988
WARNER CHILCOTT	80MG	A070340	001	Sep 24, 1986
	120MG	A070341	001	Sep 24, 1986
WATSON LABS	40MG	A072799	001	Apr 28, 1989
	40MG	A072923	001	Jun 29, 1993
	80MG	A070855	001	Sep 24, 1986
	80MG	A071366	001	Oct 01, 1986
	120MG	A070856	001	Sep 24, 1986
	120MG	A071367	001	Oct 01, 1986
TABLET, EXTENDED RELEASE; ORAL				
CALAN SR				
+ PFIZER	120MG	N019152	003	Mar 06, 1991
+	180MG **	N019152	002	Dec 15, 1989
+	240MG	N019152	001	Dec 16, 1986
COVERA-HS				
PFIZER	180MG	N020552	001	Feb 26, 1996
	240MG	N020552	002	Feb 26, 1996
VERAPAMIL HYDROCHLORIDE				
APOTEX CORP	120MG	A200878	001	Apr 20, 2012
	180MG	A200878	002	Apr 20, 2012
	240MG	A200878	003	Apr 20, 2012
IVAX SUB TEVA PHARMS	120MG	A073568	002	Oct 10, 1997
	180MG	A074330	001	Jan 31, 1994
	240MG	A073568	001	Jul 31, 1992
PLIVA	240MG	A072922	001	Mar 01, 1996
RISING	120MG	A074587	002	Feb 21, 1997
	180MG	A074587	003	Sep 09, 1997
	240MG	A074587	001	Mar 23, 1996
SUN PHARM INDS INC	120MG	A090529	001	Dec 30, 2011
	180MG	A090529	002	Dec 30, 2011
	240MG	A090529	003	Dec 30, 2011

VERATRUM VIRIDE ROOT

TABLET; ORAL

VERTAVIS

MEDPOINTE PHARM HLC 130CSR UNIT N005691 002

VIDARABINE

INJECTABLE; INJECTION

VIRA-A

PARKEDALE EQ 187.4MG BASE/ML N050523 001

OINTMENT; OPHTHALMIC

VIRA-A

PARKEDALE 3% N050486 001

VIGABATRIN

FOR SOLUTION; ORAL

VIGABATRIN

ACCORD HLTHCARE	500MG/PACKET	A214425	001	Nov 13, 2020
CHARTWELL RX	500MG/PACKET	A211790	001	Mar 10, 2022
GRANULES	500MG/PACKET	A213469	001	Apr 24, 2020
SPECGX LLC	500MG/PACKET	A212626	001	Jul 28, 2021
ZYDUS	500MG/PACKET	A214671	001	Mar 02, 2023

VINBLASTINE SULFATE

INJECTABLE; INJECTION

VELBAN

+ LILLY 10MG/VIAL ** N012665 001

VINBLASTINE SULFATE

ABRAXIS PHARM	10MG/VIAL	A089011	001	Nov 18, 1985
HOSPIRA	10MG/VIAL	A089565	001	Aug 18, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VINCRIStINE SULFATE

INJECTABLE; INJECTION

ONCOVIN

+	LILLY	1MG/VIAL **	N014103 001	
+		1MG/ML **	N014103 003	Mar 07, 1984
+		5MG/VIAL **	N014103 002	

VINCASAR PFS

	TEVA PARENTERAL	1MG/ML	A071426 001	Jul 17, 1987
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VINCREX

	BRISTOL MYERS SQUIBB	5MG/VIAL	A070867 001	Jul 12, 1988
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VINCRIStINE SULFATE

	ABIC	1MG/ML	A070873 001	Feb 19, 1987
	ABRAXIS PHARM	1MG/ML	A070411 001	Sep 10, 1986
	FRESENIUS KABI USA	1MG/ML	A076296 001	Dec 20, 2002
		1MG/ML	A076401 001	Oct 28, 2003
	HOSPIRA	1MG/VIAL	A071559 001	Apr 11, 1988
		2MG/VIAL	A071560 001	Apr 11, 1988
		5MG/VIAL	A071561 001	Apr 11, 1988

VINCRIStINE SULFATE PFS

	TEVA PHARMS USA	1MG/ML	A075493 001	Sep 01, 1999
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INJECTABLE, LIPOSOMAL; INTRAVENOUS

MARQIBO KIT

+	ACROTECH	5MG/5ML (1MG/ML)	N202497 001	Aug 09, 2012
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VINORELBINE TARTRATE

INJECTABLE; INJECTION

NAVELBINE

+	PIERRE FABRE	EQ 10MG BASE/ML **	N020388 001	Dec 23, 1994
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VINORELBINE TARTRATE

	EBEWE PHARMA	EQ 10MG BASE/ML	A078408 001	Feb 13, 2008
	FRESENIUS KABI USA	EQ 10MG BASE/ML	A076849 001	Apr 18, 2005
	HOSPIRA	EQ 10MG BASE/ML	A076827 001	Jun 02, 2005
	MYLAN LABS LTD	EQ 10MG BASE/ML	A200148 001	Aug 31, 2012
	NOVAST LABS	EQ 10MG BASE/ML	A208997 001	Aug 05, 2019
	TEVA PHARMS USA	EQ 10MG BASE/ML	A076028 001	Feb 03, 2003

VIOMYCIN SULFATE

INJECTABLE; INJECTION

VIOCIN SULFATE

	PFIZER	EQ 1GM BASE/VIAL	A061086 001	
		EQ 5GM BASE/VIAL	A061086 002	

VITAMIN A

CAPSULE; ORAL

AQUASOL A

	ASTRAZENECA	25,000USP UNITS	A083080 002	
		50,000USP UNITS	A083080 001	

VITAMIN A

	BANNER PHARMACAPS	50,000USP UNITS	A083973 001	
	CHASE CHEM	50,000 IU	A083351 001	
	EVERYLIFE	50,000 IU	A083134 001	
	IMPAX LABS	50,000USP UNITS	A080952 001	
	WEST WARD	50,000USP UNITS	A080985 001	

VITAMIN A PALMITATE

CAPSULE; ORAL

AFAXIN

	STERLING WINTHROP	EQ 50,000 UNITS BASE	A083187 001	
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ALPHALIN

	LILLY	EQ 50,000 UNITS BASE	A080883 001	
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DEL-VI-A

	DEL RAY LABS	EQ 50,000 UNITS BASE	A080830 001	
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VI-DOM-A

	BAYER PHARMS	EQ 50,000 UNITS BASE	A080972 001	
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VITAMIN A

	BANNER PHARMACAPS	EQ 50,000 UNITS BASE	A080702 001	
	BRISTOL MYERS SQUIBB	EQ 50,000 UNITS BASE	A080860 001	
	CHASE CHEM	EQ 50,000 UNITS BASE	A080746 001	
		EQ 50,000 UNITS BASE	A083207 001	
	ELKINS SINN	EQ 50,000 UNITS BASE	A085479 001	
	EVERYLIFE	EQ 50,000 UNITS BASE	A080943 001	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VITAMIN A PALMITATE

CAPSULE; ORAL

VITAMIN A

	EQ 50,000 UNITS BASE	A083114 001
IMPAX LABS	EQ 50,000 UNITS BASE	A080953 001
	EQ 50,000 UNITS BASE	A080955 001
IVAX SUB TEVA PHARMS	EQ 50,000 UNITS BASE	A083035 001
	EQ 50,000 UNITS BASE	A083190 001
MK LABS	EQ 25,000 UNITS BASE	A083457 002
	EQ 50,000 UNITS BASE	A083457 001
WEST WARD	EQ 50,000 UNITS BASE	A080967 001
WHARTON LABS	EQ 50,000 UNITS BASE	A083665 001

VITAMIN A PALMITATE

ARCUM	EQ 50,000 UNITS BASE	A083311 001
	EQ 50,000 UNITS BASE	A083321 001
BANNER PHARMACAPS	EQ 50,000 UNITS BASE	A083948 001
	EQ 50,000 UNITS BASE	A083981 001

VITAMIN A SOLUBILIZED

TEVA	EQ 50,000 UNITS BASE	A080921 001
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INJECTABLE; INJECTION

VITAMIN A PALMITATE

BEL MAR	EQ 50,000 UNITS BASE/ML	A080819 001
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VORICONAZOLE

FOR SUSPENSION; ORAL

VORICONAZOLE

RISING	200MG/5ML	A202361 001	May 28, 2013
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INJECTABLE; INTRAVENOUS

VORICONAZOLE

EUGIA PHARMA	200MG/VIAL	A212162 001	Feb 02, 2023
MYLAN LABS LTD	200MG/VIAL	A210849 001	Oct 11, 2022

TABLET; ORAL

VORICONAZOLE

TEVA PHARMS	50MG	A091658 001	Apr 06, 2012
	200MG	A091658 002	Apr 06, 2012

VORTIOXETINE HYDROBROMIDE

TABLET; ORAL

TRINTELLIX

+ TAKEDA PHARMS USA	EQ 15MG BASE **	N204447 003	Sep 30, 2013
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VORTIOXETINE HYDROBROMIDE

ZYDUS PHARMS	EQ 5MG BASE	A211146 001	Sep 17, 2021
	EQ 10MG BASE	A211146 002	Sep 17, 2021
	EQ 20MG BASE	A211146 003	Sep 17, 2021

WARFARIN POTASSIUM

TABLET; ORAL

ATHROMBIN-K

PHARM RES ASSOC	2MG	N011771 007
	5MG	N011771 004
	10MG	N011771 005
	25MG	N011771 006

WARFARIN SODIUM

INJECTABLE; INJECTION

COUMADIN

BRISTOL MYERS SQUIBB	5MG/VIAL	N009218 024	Feb 07, 1995
	50MG/VIAL	N009218 020	
	75MG/VIAL	N009218 012	

TABLET; ORAL

ATHROMBIN

PHARM RES ASSOC	5MG	N011771 003
	10MG	N011771 002
	25MG	N011771 001

COUMADIN

+ BRISTOL MYERS SQUIBB	1MG **	N009218 022	Mar 01, 1990
	2MG	N009218 013	
	2.5MG	N009218 018	
	3MG	N009218 025	Nov 18, 1996
	4MG	N009218 023	Aug 24, 1993
	5MG	N009218 007	

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

WARFARIN SODIUM

TABLET; ORAL

COUMADIN

+	6MG	N009218 026	Nov 18, 1996
+	7.5MG	N009218 016	
+	10MG	N009218 005	

PANWARFIN

ABBOTT

2MG	N017020 001
2.5MG	N017020 002
5MG	N017020 003
7.5MG	N017020 004
10MG	N017020 005

WARFARIN SODIUM

AUROBINDO PHARMA USA

1MG	A040415 001	Sep 27, 2004
2MG	A040415 002	Sep 27, 2004
2.5MG	A040415 003	Sep 29, 2004
3MG	A040415 004	Sep 27, 2004
4MG	A040415 005	Sep 27, 2004
5MG	A040415 006	Sep 27, 2004
6MG	A040415 007	Sep 27, 2004
7.5MG	A040415 008	Sep 27, 2004
10MG	A040415 009	Sep 27, 2004

CHARTWELL RX

1MG	A040196 001	Sep 30, 1997
2MG	A040196 002	Sep 30, 1997
2.5MG	A040196 003	Sep 30, 1997
3MG	A040196 008	Jul 26, 2000
4MG	A040196 004	Sep 30, 1997
5MG	A040196 005	Sep 30, 1997
6MG	A040196 009	Jul 26, 2000
7.5MG	A040196 006	Sep 30, 1997
10MG	A040196 007	Sep 30, 1997

USL PHARMA

2MG	A088719 001	Jun 27, 1985
2.5MG	A088720 001	Aug 06, 1985
5MG	A088721 001	Jul 02, 1985

WATSON LABS

2MG	A086123 001	Aug 17, 1982
2.5MG	A086120 001	Aug 17, 1982
5MG	A086119 001	Aug 17, 1982
7.5MG	A086118 001	Aug 17, 1982
10MG	A086122 001	Aug 17, 1982

XENON XE-127

GAS; INHALATION

XENON XE 127

MALLINCKRODT

5mCi/VIAL	N018536 001	Oct 01, 1982
10mCi/VIAL	N018536 002	Oct 01, 1982

XENON XE-133

GAS; INHALATION

XENON XE 133

GE HEALTHCARE

1 CI/AMP	N017256 002
10mCi/VIAL	N017687 002
20mCi/VIAL	N017687 003
5-100 CI/CYLINDER	N017550 001
0.25-5 CI/AMP	N017550 003

XENON XE 133-V.S.S.

GE HEALTHCARE

10mCi/VIAL	N017687 001
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INJECTABLE; INJECTION

XENON XE 133

GE HEALTHCARE

1.3-1.7 CI/AMP	N017256 001
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LANTHEUS MEDCL

6.3mCi/ML	N017283 001
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SOLUTION; INHALATION, INJECTION

XENEISOL

MALLINCKRODT

18-25mCi/AMP	N017262 002
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XYLOSE

POWDER; ORAL

XYLO-PFAN

SAVAGE LABS

25GM/BOT	N017605 001
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XYLOSE

LYNE

25GM/BOT	N018856 001	Mar 26, 1987
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Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZALCITABINETABLET; ORAL
HIVID

ROCHE	0.375MG	N020199 001	Jun 19, 1992
	0.75MG	N020199 002	Jun 19, 1992

ZALEPLON

CAPSULE; ORAL

ZALEPLON

HIKMA PHARMS	5MG	A078147 001	Nov 25, 2008
	10MG	A078147 002	Nov 25, 2008
MYLAN	5MG	A077238 001	Jun 06, 2008
	10MG	A077238 002	Jun 06, 2008
TEVA PHARMS	5MG	A077239 001	Jun 06, 2008
	10MG	A077239 002	Jun 06, 2008
UPSHER SMITH LABS	5MG	A078095 001	Jun 06, 2008
	5MG	A078706 001	Jun 06, 2008
	10MG	A078095 002	Jun 06, 2008
	10MG	A078706 002	Jun 06, 2008

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

TERSERA	200MCG/2ML (100MCG/ML)	N021060 003	Dec 28, 2004
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ZIDOVUDINE

INJECTABLE; INJECTION

ZIDOVUDINE

AM REGENT	10MG/ML	A091457 001	May 06, 2010
LIAONING CHENGDA	10MG/ML	A204538 001	Nov 26, 2013

TABLET; ORAL

RETROVIR

VIIV HLTHCARE	200MG	N020518 001	Dec 19, 1995
+	300MG **	N020518 002	Oct 04, 1996

ZIDOVUDINE

AUROBINDO PHARMA	60MG	N022294 001	Jul 23, 2009
HEC PHARM	300MG	A202058 001	Oct 07, 2011
HIKMA	300MG	A076844 001	Sep 19, 2005
MYLAN LABS LTD	100MG	N200732 001	Feb 23, 2011
RANBAXY LABS LTD	300MG	A077327 001	Sep 19, 2005

ZILEUTON

TABLET; ORAL

ZYFLO

CHIESI	300MG	N020471 001	Dec 09, 1996
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TABLET, EXTENDED RELEASE; ORAL

ZILEUTON

LUPIN LTD	600MG	A211972 001	Nov 05, 2019
TEVA PHARMS USA	600MG	A211043 001	May 03, 2022

ZYFLO CR

+	CHIESI	600MG **	N022052 001	May 30, 2007
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ZINC SULFATE

INJECTABLE; INJECTION

ZINC SULFATE

ABRAXIS PHARM	EQ 1MG ZINC/ML	N019229 002	May 05, 1987
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ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

ZIPRASIDONE HYDROCHLORIDE

MYLAN	EQ 20MG BASE	A202395 001	Oct 10, 2013
	EQ 40MG BASE	A202395 002	Oct 10, 2013
	EQ 60MG BASE	A202395 003	Oct 10, 2013
	EQ 80MG BASE	A202395 004	Oct 10, 2013

SUSPENSION; ORAL

GEODON

PFIZER INC	EQ 10MG BASE/ML	N021483 001	Mar 29, 2006
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DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZOLEDRONIC ACID

INJECTABLE; INTRAVENOUS

ZOLEDRONIC ACID

ACTAVIS INC	EQ 4MG BASE/5ML	A202472	001	Mar 04, 2013
AVET LIFESCIENCES	EQ 5MG BASE/100ML	A201801	001	Mar 29, 2013
BRECKENRIDGE	EQ 4MG BASE/5ML	A091170	001	Mar 04, 2013
DR REDDYS	EQ 4MG BASE/100ML	A204344	001	Nov 19, 2018
EUGIA PHARMA	EQ 4MG BASE/5ML	A207751	001	Sep 26, 2016
	EQ 5MG BASE/100ML	A209125	001	Dec 08, 2017
NOVAST LABS	EQ 4MG BASE/5ML	A208968	001	Feb 19, 2020
SHILPA	EQ 4MG BASE/5ML	A208513	001	May 15, 2019
SUN PHARMA GLOBAL	EQ 4MG BASE/VIAL	A090018	001	Mar 04, 2013
	EQ 4MG BASE/5ML	A202746	001	Mar 04, 2013

ZOMETA

+ NOVARTIS	EQ 4MG BASE/VIAL **	N021223	001	Aug 20, 2001
+	EQ 4MG BASE/5ML	N021223	002	Mar 07, 2003
+	EQ 4MG BASE/100ML	N021223	003	Jun 17, 2011

ZOLMITRIPTAN

TABLET; ORAL

ZOLMITRIPTAN

ANI PHARMS	2.5MG	A090861	001	Mar 04, 2014
	5MG	A090861	002	Mar 04, 2014
APOTEX INC	2.5MG	A202078	001	May 14, 2013
	5MG	A202078	002	May 14, 2013
MACLEODS PHARMS LTD	2.5MG	A203772	001	Sep 30, 2015
	5MG	A203772	002	Sep 30, 2015
NATCO PHARMA USA	2.5MG	A203186	001	May 14, 2013
	5MG	A203186	002	May 14, 2013
SUN PHARMA GLOBAL	2.5MG	A203476	001	Nov 13, 2014
	5MG	A203476	002	Nov 13, 2014

ZOMIG

+ IPR	2.5MG	N020768	001	Nov 25, 1997
+	5MG	N020768	002	Nov 25, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

ZOLMITRIPTAN

APOTEX INC	2.5MG	A202476	001	May 14, 2013
	5MG	A202476	002	May 14, 2013
MACLEODS PHARMS LTD	2.5MG	A204336	001	Oct 22, 2015
	5MG	A204336	002	Oct 22, 2015
RISING	2.5MG	A202855	001	Sep 20, 2019
	5MG	A202855	002	Sep 20, 2019

ZOMIG-ZMT

+ ASTRAZENECA	2.5MG	N021231	001	Feb 13, 2001
+	5MG	N021231	002	Sep 17, 2001

ZOLPIDEM TARTRATE

SPRAY, METERED; ORAL

ZOLPIMIST

+ AYTU	5MG/SPRAY	N022196	001	Dec 19, 2008
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TABLET; ORAL

ZOLPIDEM TARTRATE

DR REDDYS LABS LTD	5MG	A077985	001	Apr 23, 2007
	10MG	A077985	002	Apr 23, 2007
HIKMA	5MG	A078129	001	Apr 30, 2008
	10MG	A078129	002	Apr 30, 2008
MYLAN PHARMS INC	5MG	A078016	001	Apr 23, 2007
	10MG	A078016	002	Apr 23, 2007
RISING	5MG	A076578	001	Apr 23, 2007
	10MG	A076578	002	Apr 23, 2007
STRIDES PHARMA	5MG	A076062	001	Apr 23, 2007
	5MG	A078616	001	Nov 21, 2008
	10MG	A076062	002	Apr 23, 2007
	10MG	A078616	002	Nov 21, 2008
SUN PHARM INDS INC	5MG	A077359	001	Apr 23, 2007
	10MG	A077359	002	Apr 23, 2007
SUN PHARM INDS LTD	5MG	A078055	001	Apr 23, 2007
	10MG	A078055	002	Apr 23, 2007
SUN PHARM INDUSTRIES	5MG	A077288	001	Apr 23, 2007
	10MG	A077288	002	Apr 23, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or effectiveness reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZOLPIDEM TARTRATE

TABLET;ORAL

ZOLPIDEM TARTRATE

SYNTHON PHARMS	5MG	A077540 001	Apr 23, 2007
	10MG	A077540 002	Apr 23, 2007
WATSON LABS	5MG	A077773 001	Apr 23, 2007
	10MG	A077773 002	Apr 23, 2007
WOCKHARDT	5MG	A078426 001	May 15, 2007
	10MG	A078426 002	May 15, 2007
YUNG SHIN PHARM	5MG	A077990 001	Apr 23, 2007
	10MG	A077990 002	Apr 23, 2007

TABLET;SUBLINGUAL

INTERMEZZO

+ PURDUE PHARMA	1.75MG	N022328 001	Nov 23, 2011
+	3.5MG	N022328 002	Nov 23, 2011

ZOLPIDEM TARTRATE

MYLAN	5MG	A202657 001	Aug 08, 2016
	10MG	A202657 002	Aug 08, 2016

TABLET, EXTENDED RELEASE;ORAL

ZOLPIDEM TARTRATE

ACTAVIS ELIZABETH	6.25MG	A078179 002	Oct 13, 2010
	12.5MG	A078179 001	Jun 06, 2011
ACTAVIS LABS FL INC	6.25MG	A090153 001	Mar 25, 2013
	12.5MG	A090153 002	Mar 25, 2013
ANCHEN PHARMS	6.25MG	A078148 002	Apr 14, 2011
	12.5MG	A078148 001	Dec 03, 2010
BRECKENRIDGE	6.25MG	A213592 001	Jun 04, 2020
	12.5MG	A213592 002	Jun 04, 2020
SYNTHON PHARMS	6.25MG	A078483 001	Apr 12, 2011
	12.5MG	A078483 002	Jun 06, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

TOVALT ODT

+ BIOVAIL LABS INTL	5MG	N021412 001	Apr 25, 2007
+	10MG	N021412 002	Apr 25, 2007

ZONISAMIDE

CAPSULE;ORAL

ZONEGRAN

+ CONCORDIA	50MG **	N020789 002	Aug 22, 2003
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ZONISAMIDE

ANI PHARMS	25MG	A077639 001	Dec 22, 2005
	25MG	A077641 003	Dec 22, 2005
	50MG	A077639 002	Dec 22, 2005
	50MG	A077641 002	Dec 22, 2005
	100MG	A077639 003	Dec 22, 2005
	100MG	A077641 001	Dec 22, 2005
EPIC PHARMA LLC	25MG	A077876 001	Feb 21, 2007
	50MG	A077876 002	Feb 21, 2007
	100MG	A077876 003	Feb 21, 2007
HERITAGE PHARMA AVET	25MG	A077650 001	Apr 20, 2006
	50MG	A077650 002	Apr 20, 2006
	100MG	A077650 003	Apr 20, 2006
MYLAN PHARMS INC	25MG	A077647 001	Dec 22, 2005
	50MG	A077647 002	Dec 22, 2005
	100MG	A077647 003	Dec 22, 2005
RISING	25MG	A077637 001	Dec 22, 2005
	50MG	A077637 002	Dec 22, 2005
	100MG	A077637 003	Dec 22, 2005
ROXANE	25MG	A077648 001	Dec 22, 2005
	50MG	A077648 002	Dec 22, 2005
	100MG	A077648 003	Dec 22, 2005
SUN PHARM INDUSTRIES	25MG	A077635 001	Dec 22, 2005
	50MG	A077635 002	Dec 22, 2005
	100MG	A077635 003	Dec 22, 2005
UPSHER SMITH LABS	25MG	A077644 001	Dec 22, 2005
	50MG	A077644 002	Dec 22, 2005
	100MG	A077644 003	Dec 22, 2005

ORPHAN PRODUCTS DESIGNATIONS AND APPROVALS LIST

The list of Orphan Designations and Approvals is available at:

<http://www.fda.gov/ForIndustry/DevelopingProductsforRareDiseasesConditions/default.htm>

**DRUG PRODUCTS WHICH MUST DEMONSTRATE *IN VIVO* BIOAVAILABILITY
ONLY IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION**

ACETAMINOPHEN;ASPIRIN;BUTALBITAL
CAPSULE OR TABLET; ORAL
160-165MG;160-165MG;50MG
325MG;325MG;50MG

ASPIRIN;CAFFEINE;CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
160MG;32MG;200MG;16MG

ACETAMINOPHEN;ASPIRIN;BUTALBITAL;
CAFFEINE
CAPSULE OR TABLET; ORAL
160-165MG;160-165MG;50MG;40MG
325MG;325MG;50MG;40MG

ASPIRIN;CARISOPRODOL
TABLET; ORAL
325MG;200MG

ACETAMINOPHEN;BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG;50MG

ASPIRIN;CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
325MG;200MG;16MG

ACETAMINOPHEN;BUTALBITAL;CAFFEINE
CAPSULE OR TABLET; ORAL
325MG;50MG;40MG

ASPIRIN;MEPROBAMATE
TABLET; ORAL
325MG;200MG

AMINOPHYLLINE
TABLET; ORAL
100MG;200MG

ASPIRIN;METHOCARBAMOL
TABLET; ORAL
325MG;400MG

ASPIRIN;BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG;50MG
650MG;50MG

CHLOROTHIAZIDE
TABLET; ORAL
250MG

ASPIRIN;BUTALBITAL;CAFFEINE
CAPSULE OR TABLET; ORAL
325MG;50MG;40MG
650MG;50MG;40MG

HYDROXYZINE HYDROCHLORIDE
TABLET; ORAL
10MG;25MG;
50MG;100MG

ASPIRIN;CAFFEINE;CARISOPRODOL
TABLET; ORAL
160MG;32MG;200MG

PREDNISONE
TABLET; ORAL
1MG;2.5MG;5MG;10MG;
20MG;25MG;50MG

APPENDIX A - PRODUCT NAME INDEX

** A **

ABACAVIR SULFATE, ABACAVIR SULFATE
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE
ABELCET, AMPHOTERICIN B
ABILIFY, ARIPIPIRAZOLE
ABILIFY ASIMTUFII, ARIPIPIRAZOLE
ABILIFY MAINTENA KIT, ARIPIPIRAZOLE
ABILIFY MYCITE KIT, ARIPIPIRAZOLE
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ABLYSINOL, ALCOHOL
ABRAXANE, PACLITAXEL
ABREVA, DOCOSANOL (OTC)
ABSORICA, ISOTRETINOIN
ABSORICA LD, ISOTRETINOIN
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ACANYA, BENZOYL PEROXIDE
ACARBOSE, ACARBOSE
ACCOLATE, ZAFIRLUKAST
ACCRUFER, FERRIC MALTOL
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACEPHEN, ACETAMINOPHEN (OTC)
ACETADOTE, ACETYLCYSTEINE
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ACETAMINOPHEN
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
ACETAZOLAMIDE, ACETAZOLAMIDE
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ACETIC ACID, ACETIC ACID, GLACIAL
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ACETYLCYSTEINE, ACETYLCYSTEINE
ACIPHEX, RABEPRAZOLE SODIUM
ACITRETIN, ACITRETIN
ACTHAR GEL, CORTICOTROPIN
ACTICLATE, DOXYCYCLINE HYCLATE
ACTIGALL, URSODIOL
ACTIQ, FENTANYL CITRATE
ACTIVELLA, ESTRADIOL
ACTONEL, RISEDRONATE SODIUM
ACTOPLUS MET, METFORMIN HYDROCHLORIDE
ACTOS, PIOGLITAZONE HYDROCHLORIDE
ACULAR, KETOROLAC TROMETHAMINE
ACULAR LS, KETOROLAC TROMETHAMINE
ACUVAIL, KETOROLAC TROMETHAMINE
ACYCLOVIR, ACYCLOVIR
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ACZONE, DAPSONE
ADAPALENE, ADAPALENE (OTC)
ADAPALENE, ADAPALENE
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
ADASUVE, LOXAPINE
ADCIRCA, TADALAFIL
ADDERALL XR 10, AMPHETAMINE ASPARTATE
ADDERALL XR 15, AMPHETAMINE ASPARTATE
ADDERALL XR 20, AMPHETAMINE ASPARTATE
ADDERALL XR 25, AMPHETAMINE ASPARTATE
ADDERALL XR 30, AMPHETAMINE ASPARTATE
ADDERALL XR 5, AMPHETAMINE ASPARTATE
ADDYI, FLIBANSERIN
ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
ADEMPAS, RIOCIQUAT
ADENOSINE, ADENOSINE
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
ADLARITY, DONEPEZIL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** A **

ADRENACLICK, EPINEPHRINE
ADRENALIN, EPINEPHRINE
ADREVIEW, IOBENGUANE SULFATE I-123
ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
ADVAIR HFA, FLUTICASONE PROPIONATE
ADVIL, IBUPROFEN (OTC)
ADVIL, IBUPROFEN SODIUM (OTC)
ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL COLD AND SINUS, IBUPROFEN (OTC)
ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)
ADVIL DUAL ACTION WITH ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ADVIL LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MULTI-SYMPTOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ADZENYS XR-ODT, AMPHETAMINE
AEMCOLO, RIFAMYCIN SODIUM
AFINITOR, EVEROLIMUS
AFINITOR DISPERZ, EVEROLIMUS
AFIRMELLE, ETHINYL ESTRADIOL
AGAMREE, VAMOROLONE
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
AIRDUO DIGIHALER, FLUTICASONE PROPIONATE
AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
AIRSUPRA, ALBUTEROL SULFATE
AK-FLUOR 10%, FLUORESCEIN SODIUM
AK-FLUOR 25%, FLUORESCEIN SODIUM
AKEEGA, ABIRATERONE ACETATE
AKLIEF, TRIFAROTENE
AKOVAZ, EPHEDRINE SULFATE
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKTEN, LIDOCAINE HYDROCHLORIDE
AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
AKYNZEO, NETUPITANT
ALA-CORT, HYDROCORTISONE
ALA-SCALP, HYDROCORTISONE
ALAVERT, LORATADINE (OTC)
ALAWAY, KETOTIFEN FUMARATE (OTC)
ALBENDAZOLE, ALBENDAZOLE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALCAINE, PROPARACAINE HYDROCHLORIDE
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
ALDACTAZIDE, HYDROCHLOROTHIAZIDE
ALDACTONE, SPIRONOLACTONE
ALECENSA, ALECTINIB HYDROCHLORIDE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALEVE, NAPROXEN SODIUM (OTC)
ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ALFENTA, ALFENTANIL HYDROCHLORIDE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALIMTA, PEMETREXED DISODIUM
ALINIA, NITAZOXANIDE
ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
ALKINDI SPRINKLE, HYDROCORTISONE
ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)

APPENDIX A - PRODUCT NAME INDEX

** A **

ALLI, ORLISTAT (OTC)
ALLOPURINOL, ALLOPURINOL
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
ALLZITAL, ACETAMINOPHEN
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
ALOMIDE, LODOXAMIDE TROMETHAMINE
ALOPRIM, ALLOPURINOL SODIUM
ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
ALPHAGAN P, BRIMONIDINE TARTRATE
ALPRAZOLAM, ALPRAZOLAM
ALPROSTADIL, ALPROSTADIL
ALREX, LOTEPIREDNOL ETABONATE
ALTABAX, RETAPAMULIN
ALTACE, RAMIPRIL
ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
ALTAVERA, ETHINYL ESTRADIOL
ALTOPREV, LOVASTATIN
ALTRENO, TRETINOIN
ALUNBRIG, BRIGATINIB
ALVAIZ, ELTROMBOPAG CHOLINE
ALVESCO, CICLESONIDE
ALVIMOPAN, ALVIMOPAN
ALYACEN 1/35, ETHINYL ESTRADIOL
ALYACEN 7/7/7, ETHINYL ESTRADIOL
ALYQ, TADALAFIL
AMABELZ, ESTRADIOL
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMARYL, GLIMEPIRIDE
AMBIEN, ZOLPIDEM TARTRATE
AMBIEN CR, ZOLPIDEM TARTRATE
AMBISOME, AMPHOTERICIN B
AMBRISENTAN, AMBRISENTAN
AMCINONIDE, AMCINONIDE
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE
AMICAR, AMINOCAPROIC ACID
AMIDATE, ETOMIDATE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMINO ACIDS, AMINO ACIDS
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOPHYLLINE, AMINOPHYLLINE
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN-PF 10%, AMINO ACIDS
AMINOSYN-PF 7%, AMINO ACIDS
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMITIZA, LUBIPROSTONE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
AMMONIA N 13, AMMONIA N-13
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
AMMONIUM LACTATE, AMMONIUM LACTATE
AMMONUL, SODIUM BENZOATE
AMNESTEEM, ISOTRETINOIN
AMONDYS 45, CASIMERSEN
AMOXAPINE, AMOXAPINE
AMOXICILLIN, AMOXICILLIN

APPENDIX A - PRODUCT NAME INDEX**** A ****

AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXIL, AMOXICILLIN
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
AMPHOTERICIN B, AMPHOTERICIN B
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
AMPYRA, DALFAMPRIDINE
AMRINONE LACTATE, INAMRINONE LACTATE
AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE
AMVUTTRA, VUTRISIRAN SODIUM
AMYVID, FLORBETAPIR F-18
AMZEEQ, MINOCYCLINE HYDROCHLORIDE
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ANAPROX DS, NAPROXEN SODIUM
ANASTROZOLE, ANASTROZOLE
ANCOBON, FLUCYTOSINE
ANDROGEL, TESTOSTERONE
ANDROID 25, METHYLTESTOSTERONE
ANECTINE, SUCCINYLCHOLINE CHLORIDE
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
ANGELIQ, DROSPIRENONE
ANGIOMAX, BIVALIRUDIN
ANGIOMAX RTU, BIVALIRUDIN
ANNOVERA, ETHINYL ESTRADIOL
ANORO ELLIPTA, UMECLIDINIUM BROMIDE
ANTARA (MICRONIZED), FENOFIBRATE
ANTHELIOS 20, AVOBENZONE (OTC)
ANTHELIOS 40, AVOBENZONE (OTC)
ANTHELIOS SX, AVOBENZONE (OTC)
ANTIVERT, MECLIZINE HYDROCHLORIDE
ANUSOL HC, HYDROCORTISONE
ANZEMET, DOLASETRON MESYLATE
APHEXDA, MOTIXAFORTIDE ACETATE
APIXABAN, APIXABAN
APLENZIN, BUPROPION HYDROBROMIDE
APOKYN, APOMORPHINE HYDROCHLORIDE
APOMORPHINE HYDROCHLORIDE, APOMORPHINE HYDROCHLORIDE
APONVIE, APREPITANT
APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
APREMILAST, APREMILAST
APREPITANT, APREPITANT
APRETUDE, CABOTEGRAVIR
APRISO, MESALAMINE
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
APTIOM, ESLICARBAZEPINE ACETATE
APTIVUS, TIPRANAVIR
AQUASOL A, VITAMIN A PALMITATE
ARAKODA, TAFENOQUINE SUCCINATE
ARANELLE, ETHINYL ESTRADIOL
ARAVA, LEFLUNOMIDE
ARAZLO, TAZAROTENE
ARESTIN, MINOCYCLINE HYDROCHLORIDE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
ARGATROBAN, ARGATROBAN
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
ARICEPT, DONEPEZIL HYDROCHLORIDE
ARIDOL KIT, MANNITOL
ARIKAYCE KIT, AMIKACIN SULFATE
ARIMIDEX, ANASTROZOLE
ARIPIPRAZOLE, ARIPIPRAZOLE

APPENDIX A - PRODUCT NAME INDEX

** A **

ARISTADA, ARIPIPRAZOLE LAUROXIL
ARISTADA INITIO KIT, ARIPIPRAZOLE LAUROXIL
ARIXTRA, FONDAPARINUX SODIUM
ARMODAFINIL, ARMODAFINIL
ARMONAIR DIGIHALER, FLUTICASONE PROPIONATE
ARNUITY ELLIPTA, FLUTICASONE FUROATE
AROMASIN, EXEMESTANE
ARRANON, NELARABINE
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ARTESUNATE, ARTESUNATE
ARTHROTEC, DICLOFENAC SODIUM
ASCLERA, POLIDOCANOL
ASCOR, ASCORBIC ACID
ASENAPINE MALEATE, ASENAPINE MALEATE
ASHLYNA, ETHINYL ESTRADIOL
ASMANEX HFA, MOMETASONE FUROATE
ASMANEX TWISTHALER, MOMETASONE FUROATE
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
ASPRUZYO SPRINKLE, RANOLAZINE
ASTAGRAF XL, TACROLIMUS
ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
ATACAND, CANDESARTAN CILEXETIL
ATACAND HCT, CANDESARTAN CILEXETIL
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
ATELVIA, RISEDRONATE SODIUM
ATENOLOL, ATENOLOL
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATHENTIA NEXT, LEVONORGESTREL (OTC)
ATIVAN, LORAZEPAM
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
ATORVALIQ, ATORVASTATIN CALCIUM
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
ATOVAQUONE, ATOVAQUONE
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRALIN, TRETINOIN
ATRIPLA, EFAVIRENZ
ATROPINE SULFATE, ATROPINE SULFATE
ATROVENT HFA, IPRATROPIUM BROMIDE
AUBAGIO, TERIFLUNOMIDE
AUGMENTIN '125', AMOXICILLIN
AUGMENTIN '250', AMOXICILLIN
AUGMENTIN '875', AMOXICILLIN
AUGMENTIN ES-600, AMOXICILLIN
AUGTYRO, REPOTRECTINIB
AUROVELA 1.5/30, ETHINYL ESTRADIOL
AUROVELA 1/20, ETHINYL ESTRADIOL
AUROVELA 24 FE, ETHINYL ESTRADIOL
AUROVELA FE 1.5/30, ETHINYL ESTRADIOL
AUROVELA FE 1/20, ETHINYL ESTRADIOL
AURYXIA, FERRIC CITRATE
AUSTEDO, DEUTETRABENAZINE
AUSTEDO XR, DEUTETRABENAZINE
AUVELITY, BUPROPION HYDROCHLORIDE
AUVI-Q, EPINEPHRINE
AVAGARD, ALCOHOL (OTC)
AVAGE, TAZAROTENE
AVALIDE, HYDROCHLOROTHIAZIDE
AVAPRO, IRBESARTAN
AVEED, TESTOSTERONE UNDECANOATE
AVIANE-28, ETHINYL ESTRADIOL
AVITA, TRETINOIN
AVODART, DUTASTERIDE
AVYCAZ, AVIBACTAM SODIUM

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** A **

AXID AR, NIZATIDINE (OTC)
 AXUMIN, FLUCICLOVINE F-18
 AYUNA, ETHINYL ESTRADIOL
 AYVAKIT, AVAPRITINIB
 AZACITIDINE, AZACITIDINE
 AZACTAM, AZTREONAM
 AZASAN, AZATHIOPRINE
 AZASITE, AZITHROMYCIN
 AZATHIOPRINE, AZATHIOPRINE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZEDRA, IOBENGUANE I-131
 AZELAIC ACID, AZELAIC ACID
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 AZELEX, AZELAIC ACID
 AZILECT, RASAGILINE MESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 AZOPT, BRINZOLAMIDE
 AZOR, AMLODIPINE BESYLATE
 AZSTARYS, DEXMETHYLPHENIDATE HYDROCHLORIDE
 AZTREONAM, AZTREONAM
 AZULFIDINE, SULFASALAZINE
 AZULFIDINE EN-TABS, SULFASALAZINE

** B **

BACITRACIN, BACITRACIN
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 BACLOFEN, BACLOFEN
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
 BACTRIM, SULFAMETHOXAZOLE
 BACTRIM DS, SULFAMETHOXAZOLE
 BAFIERTAM, MONOMETHYL FUMARATE
 BAL, DIMERCAPROL
 BALANCED SALT, CALCIUM CHLORIDE
 BALCOLTRA, ETHINYL ESTRADIOL
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BALVERSA, ERDAFITINIB
 BALZIVA-28, ETHINYL ESTRADIOL
 BANZEL, RUFINAMIDE
 BAQSIMI, GLUCAGON
 BARACLUDE, ENTECAVIR
 BARHEMSYS, AMISULPRIDE
 BAXDELA, DELAFLOXACIN MEGLUMINE
 BEKYREE, DESOGESTREL
 BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 BELEODAQ, BELINOSTAT
 BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BELSOMRA, SUVOREXANT
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENTYL, DICYCLOMINE HYDROCHLORIDE
 BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
 BENZAMYCIN, BENZOYL PEROXIDE
 BENZNIDAZOLE, BENZNIDAZOLE
 BENZONATATE, BENZONATATE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE

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BEPREVE, BEPOTASTINE BESILATE
BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
BETA-VAL, BETAMETHASONE VALERATE
BETADINE, POVIDONE-IODINE
BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
BETAINE, BETAINE
BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
BETAPACE, SOTALOL HYDROCHLORIDE
BETAPACE AF, SOTALOL HYDROCHLORIDE
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BETHKIS, TOBRAMYCIN
BETIMOL, TIMOLOL
BETOPTIC, BETAXOLOL HYDROCHLORIDE
BETOPTIC S, BETAXOLOL HYDROCHLORIDE
BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
BEXAROTENE, BEXAROTENE
BEYAZ, DROSPIRENONE
BICALUTAMIDE, BICALUTAMIDE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
BICNU, CARMUSTINE
BIDIL, HYDRALAZINE HYDROCHLORIDE
BIJUVA, ESTRADIOL
BIKTARVY, BICTEGRAVIR SODIUM
BILTRICIDE, PRAZIQUANTEL
BIMATOPROST, BIMATOPROST
BINOSTO, ALENDRONATE SODIUM
BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE
BIOCRUB, CHLORHEXIDINE GLUCONATE (OTC)
BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH SUBSALICYLATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BIVALIRUDIN, BIVALIRUDIN
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
BLISOVI 24 FE, ETHINYL ESTRADIOL
BLISOVI FE 1.5/30, ETHINYL ESTRADIOL
BLISOVI FE 1/20, ETHINYL ESTRADIOL
BLOXIVERZ, NEOSTIGMINE METHYLSULFATE
BLUDIGO, INDIGOTINDISULFONATE SODIUM
BONJESTA, DOXYLAMINE SUCCINATE
BONSITY, TERIPARATIDE
BONTRIL PDM, PHENDIMETRAZINE TARTRATE
BORTEZOMIB, BORTEZOMIB
BOSENTAN, BOSENTAN
BOSULIF, BOSUTINIB MONOHYDRATE
BRAFTOVI, ENCORAFENIB
BRENZAVVY, BEXAGLIFLOZIN
BREO ELLIPTA, FLUTICASONE FUROATE
BRETHINE, TERBUTALINE SULFATE
BREVIBLOC, ESMOLOL HYDROCHLORIDE
BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVITAL SODIUM, METHOHEXITAL SODIUM
BREXAFEMME, IBREXAFUNGERP CITRATE
BREXPIPAZOLE, BREXPIPAZOLE
BREYNA, BUDESONIDE
BREZTRI AEROSPHERE, BUDESONIDE
BRIDION, SUGAMMADEX SODIUM
BRIELLYN, ETHINYL ESTRADIOL
BRILINTA, TICAGRELOR

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BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BRINZOLAMIDE, BRINZOLAMIDE
 BRISDELLE, PAROXETINE MESYLATE
 BRIVIACT, BRIVARACETAM
 BRIXADI, BUPRENORPHINE
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE,
 BROMSITE, BROMFENAC SODIUM
 BRONCHITOL, MANNITOL
 BROVANA, ARFORMOTEROL TARTRATE
 BRUKINSA, ZANUBRUTINIB
 BRYHALI, HALOBETASOL PROPIONATE
 BSS, CALCIUM CHLORIDE
 BSS PLUS, CALCIUM CHLORIDE
 BUDESONIDE, BUDESONIDE (OTC)
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUMEX, BUMETANIDE
 BUPHENYL, SODIUM PHENYLBUTYRATE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPRENORPHINE, BUPRENORPHINE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 BUSULFEX, BUSULFAN
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 BUTAPAP, ACETAMINOPHEN
 BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTRANS, BUPRENORPHINE
 BYDUREON BCISE, EXENATIDE SYNTHETIC
 BYETTA, EXENATIDE SYNTHETIC
 BYFAVO, REMIMAZOLAM BESYLATE
 BYLVAY, ODEVIXIBAT
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE

**** C ****

CABAZITAXEL, CABAZITAXEL
 CABENUVA KIT, CABOTEGRAVIR
 CABERGOLINE, CABERGOLINE
 CABOMETYX, CABOZANTINIB S-MALATE
 CABTREQ, ADAPALENE
 CADUET, AMLODIPINE BESYLATE
 CAFKIT, CAFFEINE CITRATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

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CALCIUM GLUCONATE, CALCIUM GLUCONATE
CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
CALDOLOR, IBUPROFEN
CALQUENCE, ACALABRUTINIB
CALQUENCE, ACALABRUTINIB MALEATE
CAMBIA, DICLOFENAC POTASSIUM
CAMCEVI KIT, LEUPROLIDE MESYLATE
CAMILA, NORETHINDRONE
CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
CAMZYOS, MAVACAMTEN
CANASA, MESALAMINE
CANCIDAS, CASPOFUNGIN ACETATE
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
CAPECITABINE, CAPECITABINE
CAPEX, FLUOCINOLONE ACETONIDE
CAPITAL SOLEIL 15, AVOBENZONE (OTC)
CAPLYTA, LUMATEPERONE TOSYLATE
CAPRELSA, VANDETANIB
CAPTOPRIL, CAPTOPRIL
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CARAC, FLUOROURACIL
CARAFATE, SUCRALFATE
CARBAGLU, CARGLUMIC ACID
CARBAMAZEPINE, CARBAMAZEPINE
CARBATROL, CARBAMAZEPINE
CARBIDOPA, CARBIDOPA
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
CARBOPLATIN, CARBOPLATIN
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
CARDIZEM, DILTIAZEM HYDROCHLORIDE
CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
CARDURA, DOXAZOSIN MESYLATE
CARDURA XL, DOXAZOSIN MESYLATE
CARFILZOMIB, CARFILZOMIB
CARGLUMIC ACID, CARGLUMIC ACID
CARISOPRODOL, CARISOPRODOL
CARMUSTINE, CARMUSTINE
CARNITOR, LEVOCARNITINE
CARNITOR SF, LEVOCARNITINE
CAROSPIR, SPIRONOLACTONE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CARTIA XT, DILTIAZEM HYDROCHLORIDE
CARVEDILOL, CARVEDILOL
CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
CASODEX, BICALUTAMIDE
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
CASPORYN HC, HYDROCORTISONE
CATAFLAM, DICLOFENAC POTASSIUM
CATAPRES-TTS-1, CLONIDINE
CATAPRES-TTS-2, CLONIDINE
CATAPRES-TTS-3, CLONIDINE
CAVERJECT, ALPROSTADIL
CAVERJECT IMPULSE, ALPROSTADIL
CAYSTON, AZTREONAM

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** C **

CEFACLOR, CEFACLOR
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
CEFAZOLIN IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFDINIR, CEFDINIR
CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
CEFIXIME, CEFIXIME
CEFOTETAN, CEFOTETAN DISODIUM
CEFOXITIN, CEFOXITIN SODIUM
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
CEFOXITIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFTAZIDIME, CEFTAZIDIME
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE SODIUM, CEFTRIAXONE SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEFUROXIME SODIUM, CEFUROXIME SODIUM
CELEBREX, CELECOXIB
CELECOXIB, CELECOXIB
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CELEXA, CITALOPRAM HYDROBROMIDE
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CELONTIN, METHSUXIMIDE
CENTANY, MUPIROCIN
CEPHALEXIN, CEPHALEXIN
CEQUA, CYCLOSPORINE
CERDELGA, ELIGLUSTAT TARTRATE
CEREBYX, FOSPHENYTOIN SODIUM
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CERIANNA, FLUROESTRADIOL F-18
CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
CERVIDIL, DINOPROSTONE
CESAMET, NABILONE
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE
CETRORELIX ACETATE, CETRORELIX ACETATE
CETROTIDE, CETRORELIX ACETATE
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
CHABELINA FE, ETHINYL ESTRADIOL
CHEMET, SUCCIMER
CHENODIOL, CHENODIOL
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
CHILDREN'S ADVIL, IBUPROFEN (OTC)
CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
CHILDREN'S ALAWAY, KETOTIFEN FUMARATE (OTC)
CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
CHILDREN'S ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CLARITIN, LORATADINE (OTC)

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** C **

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
 CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN
 CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
 CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHLORZOAZONE, CHLORZOAZONE
 CHOLBAM, CHOLIC ACID
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CHOLINE C-11, CHOLINE C-11
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIALIS, TADALAFIL
 CIBINQO, ABROCITINIB
 CICLOPIROX, CICLOPIROX
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)
 CIDOFOVIR, CIDOFOVIR
 CILOSTAZOL, CILOSTAZOL
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 CIMDUO, LAMIVUDINE
 CIMETIDINE, CIMETIDINE (OTC)
 CIMETIDINE, CIMETIDINE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CINVANTI, APREPITANT
 CIPRO, CIPROFLOXACIN
 CIPRO, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
 CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
 CLADRIBINE, CLADRIBINE
 CLARAVIS, ISOTRETINOIN
 CLARINEX, DESLORATADINE

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** C **

CLARINEX-D 12 HOUR, DESLORATADINE
CLARISCAN, GADOTERATE MEGLUMINE
CLARITHROMYCIN, CLARITHROMYCIN
CLARITIN, LORATADINE (OTC)
CLARITIN HIVES RELIEF, LORATADINE (OTC)
CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
CLARITIN REDITABS, LORATADINE (OTC)
CLARITIN-D, LORATADINE (OTC)
CLARITIN-D 24 HOUR, LORATADINE (OTC)
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
CLENPIQ, CITRIC ACID
CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
CLEOCIN, CLINDAMYCIN PHOSPHATE
CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLEOCIN T, CLINDAMYCIN PHOSPHATE
CLEVIPREX, CLEVIDIPINE
CLIMARA, ESTRADIOL
CLIMARA PRO, ESTRADIOL
CLINDA-DERM, CLINDAMYCIN PHOSPHATE
CLINDAGEL, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
CLINDESSE, CLINDAMYCIN PHOSPHATE
CLINDETS, CLINDAMYCIN PHOSPHATE
CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,
CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
CLINOLIPID 20%, OLIVE OIL
CLOBAZAM, CLOBAZAM
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBEX, CLOBETASOL PROPIONATE
CLOCORTOLONE PIVALATE, CLOCORTOLONE PIVALATE
CLODERM, CLOCORTOLONE PIVALATE
CLOFARABINE, CLOFARABINE
CLOLAR, CLOFARABINE

APPENDIX A - PRODUCT NAME INDEX

** C **

CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE, CLONIDINE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
CLOTRIMAZOLE, CLOTRIMAZOLE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOZAPINE, CLOZAPINE
CLOZARIL, CLOZAPINE
COARTEM, ARTEMETHER
CODEINE SULFATE, CODEINE SULFATE
COL-PROBENECID, COLCHICINE
COLAZAL, BALSALAZIDE DISODIUM
COLCHICINE, COLCHICINE
COLCRYS, COLCHICINE
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
COLESTID, COLESTIPOL HYDROCHLORIDE
COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
COLOCORT, HYDROCORTISONE
COLY-MYCIN M, COLISTIMETHATE SODIUM
COLY-MYCIN S, COLISTIN SULFATE
COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
COMBIGAN, BRIMONIDINE TARTRATE
COMBIPATCH, ESTRADIOL
COMBIVENT RESPIMAT, ALBUTEROL SULFATE
COMBIVIR, LAMIVUDINE
COMBOGESIC IV, ACETAMINOPHEN
COMETRIQ, CABOZANTINIB S-MALATE
COMPLERA, EMTRICITABINE
COMPRO, PROCHLORPERAZINE
COMTAN, ENTACAPONE
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
CONDYLOX, PODOFILOX
CONRAY, IOTHALAMATE MEGLUMINE
CONTRAVE, BUPROPION HYDROCHLORIDE
CONZIP, TRAMADOL HYDROCHLORIDE
COPAXONE, GLATIRAMER ACETATE
COPIKTRA, DUVELISIB
CORDRAN, FLURANDRENOLIDE
COREG, CARVEDILOL
COREG CR, CARVEDILOL PHOSPHATE
CORGARD, NADOLOL
CORLANOR, IVABRADINE
CORLANOR, IVABRADINE HYDROCHLORIDE
CORLOPAM, FENOLDOPAM MESYLATE
CORMAX, CLOBETASOL PROPIONATE
CORPHEDRA, EPHEDRINE SULFATE
CORTEF, HYDROCORTISONE
CORTENEMA, HYDROCORTISONE
CORTIFOAM, HYDROCORTISONE ACETATE
CORTROSYN, COSYNTROPIN
CORVERT, IBUTILIDE FUMARATE
COSELA, TRILACICLIB DIHYDROCHLORIDE
COSMEGEN, DACTINOMYCIN
COSOPT, DORZOLAMIDE HYDROCHLORIDE
COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
COSYNTROPIN, COSYNTROPIN
COTELLIC, COBIMETINIB FUMARATE
COTEMPLA XR-ODT, METHYLPHENIDATE
COXANTO, OXAPROZIN
COZAAR, LOSARTAN POTASSIUM

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** C **

CRESEMBA, ISAVUCONAZONIUM SULFATE
 CRESTOR, ROSUVASTATIN CALCIUM
 CRINONE, PROGESTERONE
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CROTAN, CROTAMITON
 CRYSELLE, ETHINYL ESTRADIOL
 CUPRIC CHLORIDE, CUPRIC CHLORIDE
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CUPRIC SULFATE, CUPRIC SULFATE
 CUPRIMINE, PENICILLAMINE
 CUVPOSA, GLYCOPYRROLATE
 CUVRIOR, TRIENTINE TETRAHYDROCHLORIDE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYANOKIT, HYDROXOCOBALAMIN
 CYCLESSA, DESOGESTREL
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYCLOSET, BROMOCRIPTINE MESYLATE
 CYCLOSPORINE, CYCLOSPORINE
 CYKLOKAPRON, TRANEXAMIC ACID
 CYMBALTA, DULOXETINE HYDROCHLORIDE
 CYONANZ, ETHINYL ESTRADIOL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 CYSTADANE, BETAINE
 CYSTADROPS, CYSTEAMINE HYDROCHLORIDE
 CYSTAGON, CYSTEAMINE BITARTRATE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE
 CYTALUX, PAFOLACIANINE SODIUM
 CYTARABINE, CYTARABINE
 CYTOMEL, LIOTHYRONINE SODIUM
 CYTOTEC, MISOPROSTOL
 CYTOXAN, CYCLOPHOSPHAMIDE

** D **

DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DACARBAZINE, DACARBAZINE
 DACTINOMYCIN, DACTINOMYCIN
 DALFAMPRIDINE, DALFAMPRIDINE
 DALIRESP, ROFLUMILAST
 DALVANCE, DALBAVANCIN HYDROCHLORIDE
 DANAZOL, DANAZOL
 DANTRIUM, DANTROLENE SODIUM
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DAPSONE, DAPSONE
 DAPTOMYCIN, DAPTOMYCIN
 DAPTOMYCIN IN 0.9% SODIUM CHLORIDE, DAPTOMYCIN
 DARAPRIM, PYRIMETHAMINE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DARUNAVIR, DARUNAVIR
 DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL
 DATSCAN, IOFLUPANE I-123
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DAURISMO, GLASDEGIB MALEATE
 DAYBUE, TROFINETIDE
 DAYPRO, OXAPROZIN
 DAYSEE, ETHINYL ESTRADIOL

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DAYTRANA, METHYLPHENIDATE
 DAYVIGO, LEMBOREXANT
 DDAVP, DESMOPRESSIN ACETATE
 DECITABINE, DECITABINE
 DEFENCATH, HEPARIN SODIUM
 DEFERASIROX, DEFERASIROX
 DEFERIPRONE, DEFERIPRONE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEFINITY, PERFLUTREN
 DEFINITY RT, PERFLUTREN
 DEFITELIO, DEFIBROTIDE SODIUM
 DELESTROGEN, ESTRADIOL VALERATE
 DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELSTRIGO, DORAVIRINE
 DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
 DELZICOL, MESALAMINE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DEMSER, METYROSINE
 DENAVIR, PENCICLOVIR
 DEOXYCHOLIC ACID, DEOXYCHOLIC ACID
 DEPAKOTE, DIVALPROEX SODIUM
 DEPAKOTE ER, DIVALPROEX SODIUM
 DEPEN, PENICILLAMINE
 DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
 DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
 DERMABET, BETAMETHASONE VALERATE
 DERMOTIC, FLUOCINOLONE ACETONIDE
 DESCOVY, EMTRICITABINE
 DESFERAL, DEFEROXAMINE MESYLATE
 DESFLURANE, DESFLURANE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESLORATADINE, DESLORATADINE
 DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DESONIDE, DESONIDE
 DESOWEN, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
 DESVENLAFAXINE, DESVENLAFAXINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DETECTNET, COPPER CU-64 DOTATATE
 DETROL, TOLTERODINE TARTRATE
 DETROL LA, TOLTERODINE TARTRATE
 DEXAMETHASONE, DEXAMETHASONE
 DEXAMETHASONE INTENSOL, DEXAMETHASONE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXASPORIN, DEXAMETHASONE
 DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE

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** D **

DEXILANT, DEXLANSOPRAZOLE
DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
DEXTENZA, DEXAMETHASONE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE
DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 25%, DEXTROSE
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075%, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15%, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.225%, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,

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** D **

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER,
 DEXTROSE 50%, DEXTROSE
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 DEXYCU KIT, DEXAMETHASONE
 DHIVY, CARBIDOPA
 DIABETA, GLYBURIDE
 DIACOMIT, STIRIPENTOL
 DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIASTAT, DIAZEPAM
 DIASTAT ACUDIAL, DIAZEPAM
 DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM, DIATRIZOATE MEGLUMINE
 DIAZEPAM, DIAZEPAM
 DIAZEPAM INTENSOL, DIAZEPAM
 DIAZOXIDE, DIAZOXIDE
 DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
 DICHLORPHENAMIDE, DICHLORPHENAMIDE
 DICLEGIS, DOXYLAMINE SUCCINATE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 DIFFERIN, ADAPALENE (OTC)
 DIFFERIN, ADAPALENE
 DIFICID, FIDAXOMICIN
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DIFLUCAN, FLUCONAZOLE
 DIFLUNISAL, DIFLUNISAL
 DIFLUPREDNATE, DIFLUPREDNATE
 DIGOXIN, DIGOXIN
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DILANTIN, PHENYTOIN
 DILANTIN, PHENYTOIN SODIUM
 DILANTIN-125, PHENYTOIN
 DILAUDID, HYDROMORPHONE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMENHYDRINATE, DIMENHYDRINATE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DIOVAN, VALSARTAN
 DIOVAN HCT, HYDROCHLOROTHIAZIDE
 DIPENTUM, OLSALAZINE SODIUM
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIPRIVAN, PROPOFOL
 DIPROLENE, BETAMETHASONE DIPROPIONATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DISULFIRAM, DISULFIRAM
 DIURIL, CHLOROTHIAZIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DIVIGEL, ESTRADIOL

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** D **

DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
DOCOSANOL, DOCOSANOL (OTC)
DODEX, CYANOCOBALAMIN
DOFETILDE, DOFETILIDE
DOFETILIDE, DOFETILIDE
DOJOLVI, TRIHEPTANOIN
DOLISHALE, ETHINYL ESTRADIOL
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPRAM, DOXAPRAM HYDROCHLORIDE
DOPTelet, AVATROMBOPAG MALEATE
DORAL, QUAZEPAM
DORYX, DOXYCYCLINE HYCLATE
DORYX MPC, DOXYCYCLINE HYCLATE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DOTAREM, GADOTERATE MEGLUMINE
DOVATO, DOLUTEGRAVIR SODIUM
DOVONEX, CALCIPOTRIENE
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXERCALCIFEROL, DOXERCALCIFEROL
DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DOXY 100, DOXYCYCLINE HYCLATE
DOXY 200, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
DOXYCYCLINE, DOXYCYCLINE HYCLATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT
DRAXIMAGE DTPA, TECHNETIUM TC-99M PENTETATE KIT
DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
DRISDOL, ERGOCALCIFEROL
DRONABINOL, DRONABINOL
DROPERIDOL, DROPERIDOL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
DROXIA, HYDROXYUREA
DROXIDOPA, DROXIDOPA
DSUVIA, SUFENTANIL CITRATE
DUAC, BENZOYL PEROXIDE
DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
DUAVEE, BAZEDOXIFENE ACETATE
DUETACT, GLIMEPIRIDE
DUEXIS, FAMOTIDINE
DULERA, FORMOTEROL FUMARATE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
DUOBRII, HALOBETASOL PROPIONATE
DUODOTE, ATROPINE
DUOPA, CARBIDOPA
DURACLON, CLONIDINE HYDROCHLORIDE
DURAMORPH PF, MORPHINE SULFATE
DURAPREP, IODINE POVACRYLEX (OTC)
DUREZOL, DIFLUPREDNATE
DURYSTA, BIMATOPROST
DUTASTERIDE, DUTASTERIDE
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
DUVOID, BETHANECHOL CHLORIDE

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** D **

DYANAVEL XR, AMPHETAMINE
 DYANAVEL XR 10, AMPHETAMINE
 DYANAVEL XR 15, AMPHETAMINE
 DYANAVEL XR 20, AMPHETAMINE
 DYANAVEL XR 5, AMPHETAMINE
 DYCLOPRO, DYCLONINE HYDROCHLORIDE
 DYMISTA, AZELASTINE HYDROCHLORIDE
 DYRENIUM, TRIAMTERENE

** E **

E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)
 E-Z-HD, BARIUM SULFATE
 E-Z-PAQUE, BARIUM SULFATE
 E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
 E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
 EC-NAPROSYN, NAPROXEN
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ECOZA, ECONAZOLE NITRATE
 EDARBI, AZILSARTAN KAMEDOXOMIL
 EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
 EDECRIN, ETHACRYNATE SODIUM
 EDECRIN, ETHACRYNIC ACID
 EDETATE CALCIUM DISODIUM, EDETATE CALCIUM DISODIUM
 EDEX, ALPROSTADIL
 EDLUAR, ZOLPIDEM TARTRATE
 EDURANT, RILPIVIRINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFAVIRENZ; EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 EFINACONAZOLE, EFINACONAZOLE
 EFUDEX, FLUOROURACIL
 EGATEN, TRICLABENDAZOLE
 ELCYS, CYSTEINE HYDROCHLORIDE
 ELESTRIN, ESTRADIOL
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ELIDEL, PIMECROLIMUS
 ELIGARD KIT, LEUPROLIDE ACETATE
 ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
 ELIMITE, PERMETHRIN
 ELINEST, ETHINYL ESTRADIOL
 ELIQUIS, APIXABAN
 ELIXOPHYLLIN, THEOPHYLLINE
 ELLA, ULIPRISTAL ACETATE
 ELLENCE, EPIRUBICIN HYDROCHLORIDE
 ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 ELOXATIN, OXALIPLATIN
 ELUCIREM, GADOPICLENOL
 ELURYNG, ETHINYL ESTRADIOL
 ELYXYB, CELECOXIB
 EMBELINE, CLOBETASOL PROPIONATE
 EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
 EMEND, APREPITANT
 EMEND, FOSAPREPITANT DIMEGLUMINE
 EMERPHED, EPHEDRINE SULFATE
 EMFLAZA, DEFLAZACORT
 EMPAVELI, PEGCETACOPLAN
 EMSAM, SELEGILINE
 EMTRICITABINE, EMTRICITABINE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE

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** E **

EMTRIVA, EMTRICITABINE
EMVERM, MEBENDAZOLE
EMZAHH, NORETHINDRONE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ENALAPRILAT, ENALAPRILAT
ENDARI, L-GLUTAMINE
ENDOMETRIN, PROGESTERONE
ENILLORING, ETHINYL ESTRADIOL
ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
ENPRESSE-28, ETHINYL ESTRADIOL
ENSKYCE, DESOGESTREL
ENSTILAR, BETAMETHASONE DIPROPIONATE
ENTACAPONE, ENTACAPONE
ENTADFI, FINASTERIDE
ENTECAVIR, ENTECAVIR
ENTERO VU 24%, BARIUM SULFATE
ENTOCORT EC, BUDESONIDE
ENTRESTO, SACUBITRIL
ENVARUS XR, TACROLIMUS
EOVIST, GADOXETATE DISODIUM
EPANED, ENALAPRIL MALEATE
EPCLUSA, SOFOSBUVIR
EPHEDRINE SULFATE, EPHEDRINE SULFATE
EPIDIOLEX, CANNABIDIOL
EPIDUO, ADAPALENE
EPIDUO FORTE, ADAPALENE
EPIFOAM, HYDROCORTISONE ACETATE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
EPINEPHRINE, EPINEPHRINE
EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
EPIPEN, EPINEPHRINE
EPIPEN JR., EPINEPHRINE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
EPITOL, CARBAMAZEPINE
EPIVIR, LAMIVUDINE
EPIVIR-HBV, LAMIVUDINE
EPLERENONE, EPLERENONE
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
EPRONTIA, TOPIRAMATE
EPSOLAY, BENZOYL PEROXIDE
EPTIFIBATIDE, EPTIFIBATIDE
EPZICOM, ABACAVIR SULFATE
EQUETRO, CARBAMAZEPINE
ERAXIS, ANIDULAFUNGIN
ERGOALCIFEROL, ERGOALCIFEROL
ERGOLOID MESYLATES, ERGOLOID MESYLATES
ERGOMAR, ERGOTAMINE TARTRATE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ERIVEDGE, VISMODEGIB
ERLEADA, APALUTAMIDE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
ERMEZA, LEVOTHYROXINE SODIUM
ERRIN, NORETHINDRONE
ERTACZO, SERTACONAZOLE NITRATE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ERY-TAB, ERYTHROMYCIN
ERYC, ERYTHROMYCIN
ERYGEL, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHRA-DERM, ERYTHROMYCIN
ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN, ERYTHROMYCIN

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** E **

ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
 ESBRIET, PIRFENIDONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESTARYLLA, ETHINYL ESTRADIOL
 ESTAZOLAM, ESTAZOLAM
 ESTRACE, ESTRADIOL
 ESTRADIOL, ESTRADIOL
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ESTRADIOL AND NORGESTIMATE, ESTRADIOL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 ESTRING, ESTRADIOL
 ESTROGEL, ESTRADIOL
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHAMOLIN, ETHANOLAMINE OLEATE
 ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL
 ETHINYL ESTRADIOL; ETNOGESTREL, ETHINYL ESTRADIOL
 ETHOSUXIMIDE, ETHOSUXIMIDE
 ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ETHYOL, AMIFOSTINE
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 ETOPOSIDE, ETOPOSIDE
 ETRAVIRINE, ETRAVIRINE
 EUCRISA, CRISABOROLE
 EURAX, CROTAMITON
 EUTHYROX, LEVOTHYROXINE SODIUM **
 EVAMIST, ESTRADIOL
 EVEKEO, AMPHETAMINE SULFATE
 EVEKEO ODT, AMPHETAMINE SULFATE
 EVEROLIMUS, EVEROLIMUS
 EVISTA, RALOXIFENE HYDROCHLORIDE
 EVOMELA, MELPHALAN HYDROCHLORIDE
 EVOTAZ, ATAZANAVIR SULFATE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 EVRYSDI, RISDIPLAM
 EXCEDRIN (MIGRAINE RELIEF), ACETAMINOPHEN (OTC)
 EXELDERM, SULCONAZOLE NITRATE
 EXELON, RIVASTIGMINE
 EXEM FOAM KIT, AIR POLYMER-TYPE A
 EXEMESTANE, EXEMESTANE
 EXFORGE, AMLODIPINE BESYLATE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)
 EXJADE, DEFERASIROX
 EXKIVITY, MOBOCERTINIB SUCCINATE
 EXONDYS 51, ETEPLIRSEN
 EXPAREL, BUPIVACAINE
 EXSERVAN, RILUZOLE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 EXTINA, KETOCONAZOLE
 EXTRANEAL, ICODextrin
 EXXUA, GEPIRONE HYDROCHLORIDE

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** E **

EYSUVIS, LOTE PredNOL ETABONATE
 EZETIMIBE, EZETIMIBE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE

** F **

FABHALTA, IPTACOPAN HYDROCHLORIDE
 FABIOR, TAZAROTENE
 FALLBACK SOLO, LEVONORGESTREL (OTC)
 FALMINA, ETHINYL ESTRADIOL
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE (OTC)
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
 FANAPT, ILOPERIDONE
 FARESTON, TOREMIFENE CITRATE
 FARXIGA, DAPAGLIFLOZIN
 FASLODEX, FULVESTRANT
 FEBUXOSTAT, FEBUXOSTAT
 FELBAMATE, FELBAMATE
 FELBATOL, FELBAMATE
 FELDENE, PIROXICAM
 FELODIPINE, FELODIPINE
 FEMARA, LETROZOLE
 FEMRING, ESTRADIOL ACETATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FENOGLIDE, FENOFIBRATE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FENSOLVI KIT, LEUPROLIDE ACETATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-37, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL
 FENTANYL-87, FENTANYL
 FENTORA, FENTANYL CITRATE
 FERAHEME, FERUMOXYTOL
 FERRIPROX, DEFERIPRONE
 FERRLECIT, FERRIC OXYHYDROXIDE
 FERUMOXYTOL, FERUMOXYTOL
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FETROJA, CEFIDEROCOL SULFATE TOSYLATE
 FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
 FEXINIDAZOLE, FEXINIDAZOLE
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FILSPARI, SPARSENTAN
 FILSUVEZ, BIRCH TRITERPENES
 FINACEA, AZELAIC ACID
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FINTEPLA, FENFLURAMINE HYDROCHLORIDE
 FINZALA, ETHINYL ESTRADIOL
 FIORICET W/ CODEINE, ACETAMINOPHEN

APPENDIX A - PRODUCT NAME INDEX

** F **

FIRAZYR, ICATIBANT ACETATE
 FIRDAPSE, AMIFAMPRIDINE PHOSPHATE
 FIRMAGON, DEGARELIX ACETATE
 FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE
 FLAC, FLUOCINOLONE ACETONIDE
 FLAGYL, METRONIDAZOLE
 FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
 FLAREX, FLUOROMETHOLONE ACETATE
 FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLECTOR, DICLOFENAC EPOLAMINE
 FLEQSUVY, BACLOFEN
 FLOLAN, EPOPROSTENOL SODIUM
 FLOLIPID, SIMVASTATIN
 FLOMAX, TAMSULOSIN HYDROCHLORIDE
 FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)
 FLONASE SENSIMIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)
 FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 50, FLUTICASONE PROPIONATE
 FLOVENT HFA, FLUTICASONE PROPIONATE
 FLOXURIDINE, FLOXURIDINE
 FLUCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCYTOSINE, FLUCYTOSINE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 FLUMAZENIL, FLUMAZENIL
 FLUNISOLIDE, FLUNISOLIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUOCINONIDE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUORESCHEIN SODIUM, FLUORESCHEIN SODIUM
 FLUORESCHEIN SODIUM AND BENOXINATE HYDROCHLORIDE, BENOXINATE HYDROCHLORIDE
 FLUORESCITE, FLUORESCHEIN SODIUM
 FLUORODOPA F18, FLUORODOPA F-18
 FLUOROMETHOLONE, FLUOROMETHOLONE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURANDRENOLIDE, FLURANDRENOLIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 FLUTAMIDE, FLUTAMIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FML, FLUOROMETHOLONE
 FML FORTE, FLUOROMETHOLONE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCINVEZ, FOSAPREPITANT DIMEGLUMINE
 FOLIC ACID, FOLIC ACID
 FOLOTYN, PRALATREXATE
 FOMEPIZOLE, FOMEPIZOLE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM

APPENDIX A - PRODUCT NAME INDEX

** F **

FORANE, ISOFLURANE
 FORFIVO XL, BUPROPION HYDROCHLORIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 FORTAMET, METFORMIN HYDROCHLORIDE
 FORTEO, TERIPARATIDE
 FORTESTA, TESTOSTERONE
 FOSAMAX, ALENDRONATE SODIUM
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSCAVIR, FOSCARNET SODIUM
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FOSRENOL, LANTHANUM CARBONATE
 FOTIVDA, TIVOZANIB HYDROCHLORIDE
 FRAGMIN, DALTEPARIN SODIUM
 FROVA, FROVATRIPTAN SUCCINATE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FRUZAQLA, FRUQUINTINIB
 FULVESTRANT, FULVESTRANT
 FULVICIN P/G, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G 165, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G 330, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN-U/F, GRISEOFULVIN, MICROCRYSTALLINE
 FURADANTIN, NITROFURANTOIN
 FUROSCIX, FUROSEMIDE
 FUROSEMIDE, FUROSEMIDE
 FUSILEV, LEVOLEUCOVORIN CALCIUM
 FUZEON, ENFUVRTIDE
 FYARRO, SIROLIMUS
 FYAVOLV, ETHINYL ESTRADIOL
 FYCOMPA, PERAMPANEL
 FYREMADEL, GANIRELIX ACETATE

** G **

GABAPENTIN, GABAPENTIN
 GABITRIL, TIAGABINE HYDROCHLORIDE
 GABLOFEN, BACLOFEN
 GADAVIST, GADOBUTROL
 GADOBUTROL, GADOBUTROL
 GADOTERATE MEGLUMINE, GADOTERATE MEGLUMINE
 GALAFOLD, MIGALASTAT HYDROCHLORIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 GALLIUM GA 68 EDOTREOTIDE, GALLIUM GA-68 EDOTREOTIDE
 GALLIUM GA 68 GOZETOTIDE, GALLIUM GA-68 GOZETOTIDE
 GALZIN, ZINC ACETATE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GANZYK-RTU, GANCICLOVIR
 GASTROCROM, CROMOLYN SODIUM
 GASTROGRAFIN, DIATRIZOATE MEGLUMINE
 GATIFLOXACIN, GATIFLOXACIN
 GATTEX KIT, TEDUGLUTIDE RECOMBINANT
 GAVRETO, PRALSETINIB
 GEFITINIB, GEFITINIB
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GEMFIBROZIL, GEMFIBROZIL
 GEMMILY, ETHINYL ESTRADIOL
 GEMTESA, VIBEGRON
 GENERLAC, LACTULOSE
 GENGRAF, CYCLOSPORINE

APPENDIX A - PRODUCT NAME INDEX

** G **

GENOSYL, NITRIC OXIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 GENVOYA, COBICISTAT
 GEODON, ZIPRASIDONE HYDROCHLORIDE
 GEODON, ZIPRASIDONE MESYLATE
 GIAPREZA, ANGIOTENSIN II ACETATE
 GILDAGIA, ETHINYL ESTRADIOL
 GILDESS 24 FE, ETHINYL ESTRADIOL
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GILOTRIF, AFATINIB DIMALEATE
 GIMOTI, METOCLOPRAMIDE HYDROCHLORIDE
 GIVLAARI, GIVOSIRAN SODIUM
 GLATIRAMER ACETATE, GLATIRAMER ACETATE
 GLATOPA, GLATIRAMER ACETATE
 GLEEVEC, IMATINIB MESYLATE
 GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE
 GLEOSTINE, LOMUSTINE
 GLIADEL, CARMUSTINE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLOFIL-125, IOTHALAMATE SODIUM I-125
 GLOPERBA, COLCHICINE
 GLUCAGEN, GLUCAGON HYDROCHLORIDE
 GLUCAGON, GLUCAGON
 GLUCAGON, GLUCAGON HYDROCHLORIDE
 GLUCOTROL XL, GLIPIZIDE
 GLUMETZA, METFORMIN HYDROCHLORIDE
 GLYBURIDE, GLYBURIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GLYDO, LIDOCAINE HYDROCHLORIDE
 GLYNASE, GLYBURIDE
 GLYRX-PF, GLYCOPYRROLATE
 GLYSET, MIGLITOL
 GLYXAMBI, EMPAGLIFLOZIN
 GOCOVRI, AMANTADINE HYDROCHLORIDE
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 GOPRELTO, COCAINE HYDROCHLORIDE
 GRALISE, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GUAIFENESIN, GUAIFENESIN (OTC)
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 GVOKE HYOPEN, GLUCAGON
 GVOKE KIT, GLUCAGON
 GVOKE PFS, GLUCAGON
 GYNAZOLE-1, BUTOCONAZOLE NITRATE

** H **

HABITROL, NICOTINE (OTC)
 HAILEY 1.5/30, ETHINYL ESTRADIOL
 HAILEY 24 FE, ETHINYL ESTRADIOL
 HAILEY FE 1.5/30, ETHINYL ESTRADIOL
 HAILEY FE 1/20, ETHINYL ESTRADIOL
 HALAVEN, ERIBULIN MESYLATE

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** H **

HALCINONIDE, HALCINONIDE
 HALCION, TRIAZOLAM
 HALDOL, HALOPERIDOL DECANOATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HALOETTE, ETHINYL ESTRADIOL
 HALOG, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HARVONI, LEDIPASVIR
 HEATHER, NORETHINDRONE
 HECTOROL, DOXERCALCIFEROL
 HEMABATE, CARBOPROST TROMETHAMINE
 HEMADY, DEXAMETHASONE
 HEMANGEOL, PROPRANOLOL HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPSERA, ADEFOVIR DIPIVOXIL
 HEPZATO, MELPHALAN HYDROCHLORIDE
 HER STYLE, LEVONORGESTREL (OTC)
 HETLIOZ, TASIMELTEON
 HETLIOZ LQ, TASIMELTEON
 HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
 HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)
 HICON, SODIUM IODIDE I-131
 HIPREX, METHENAMINE HIPPURATE
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 HORIZANT, GABAPENTIN ENACARBIL
 HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 HYCODAN, HOMATROPINE METHYLBROMIDE
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDREA, HYDROXYUREA
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND ASPIRIN, ASPIRIN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 HYDROCORTISONE, HYDROCORTISONE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE IN ABSORBABLE, HYDROCORTISONE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYUREA, HYDROXYUREA
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 HYFTOR, SIROLIMUS

APPENDIX A - PRODUCT NAME INDEX

** H **

HYSINGLA ER, HYDROCODONE BITARTRATE
 HYZAAR, HYDROCHLOROTHIAZIDE

** I **

IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBRANCE, PALBOCICLIB
 IBSRELA, TENAPANOR HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN LYSINE, IBUPROFEN LYSINE
 IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 ICLEVIA, ETHINYL ESTRADIOL
 ICLUSIG, PONATINIB HYDROCHLORIDE
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IDHIFA, ENASIDENIB MESYLATE
 IDKIT:HP, CITRIC ACID
 IDOSE TR, TRAVOPROST
 IFEX, IFOSFAMIDE
 IFOSFAMIDE, IFOSFAMIDE
 IGALMI, DEXMEDETOMIDINE HYDROCHLORIDE
 IHEEZO, CHLOROPROCAINE HYDROCHLORIDE
 ILEVRO, NEPAFENAC
 ILLUCCIX, GALLIUM GA-68 GOZETOTIDE
 ILUVIEN, FLUOCINOLONE ACETONIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IMBRUVICA, IBRUTINIB
 IMCIVREE, SETMELANOTIDE ACETATE
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IMIQUIMOD, IMIQUIMOD
 IMITREX, SUMATRIPTAN
 IMITREX, SUMATRIPTAN SUCCINATE
 IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
 IMMPHENTIV, PHENYLEPHRINE HYDROCHLORIDE
 IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMODIUM MULTI-SYMPOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMPAVIDO, MILTEFOSINE
 IMPOYZ, CLOBETASOL PROPIONATE
 IMURAN, AZATHIOPRINE
 IMVEXXY, ESTRADIOL
 INBRIJA, LEVODOPA
 INCASSIA, NORETHINDRONE
 INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE
 INDAPAMIDE, INDAPAMIDE
 INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
 INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
 INDOCIN, INDOMETHACIN
 INDOCYANINE GREEN, INDOCYANINE GREEN
 INDOMETHACIN, INDOMETHACIN
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 INFANT'S ADVIL, IBUPROFEN (OTC)
 INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 INFED, FERRIC OXYHYDROXIDE
 INFUMORPH, MORPHINE SULFATE

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** I **

INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC, ASCORBIC ACID
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INGREZZA, VALBENAZINE TOSYLATE
INJECTAFER, FERRIC CARBOXYMALTOSE
INLYTA, AXITINIB
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
INOMAX, NITRIC OXIDE
INPEFA, SOTAGLIFLOZIN
INQOVI, CEDAZURIDINE
INREBIC, FEDRATINIB HYDROCHLORIDE
INSPRA, EPLERENONE
INTELENCE, ETRAVIRINE
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
INTRAROSA, PRASTERONE
INTROVALE, ETHINYL ESTRADIOL
INTUNIV, GUANFACINE HYDROCHLORIDE
INVANZ, ERTAPENEM SODIUM
INVEGA, PALIPERIDONE
INVEGA HAFYERA, PALIPERIDONE PALMITATE
INVEGA SUSTENNA, PALIPERIDONE PALMITATE
INVEGA TRINZA, PALIPERIDONE PALMITATE
INVELTYS, LOTEHPREDNOL ETABONATE
INVOKAMET, CANAGLIFLOZIN
INVOKAMET XR, CANAGLIFLOZIN
INVOKANA, CANAGLIFLOZIN
IODIXANOL, IODIXANOL
IOFLUPANE I-123, IOFLUPANE I-123
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IOPAMIDOL, IOPAMIDOL
IOPIDINE, APRACLONIDINE HYDROCHLORIDE
IOSAT, POTASSIUM IODIDE (OTC)
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
IRBESARTAN, IRBESARTAN
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRESSA, GEFITINIB
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
ISENTRESS, RALTEGRAVIR POTASSIUM
ISENTRESS HD, RALTEGRAVIR POTASSIUM
ISIBLOOM, DESOGESTREL
ISOFLURANE, ISOFLURANE
ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
ISONIAZID, ISONIAZID
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
ISOPTO ATROPINE, ATROPINE SULFATE
ISOPTO CARPINE, Pilocarpine Hydrochloride
ISORDIL, ISOSORBIDE DINITRATE
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
ISOSORBIDE DINITRATE AND HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
ISOSULFAN BLUE, ISOSULFAN BLUE
ISOTRETINOIN, ISOTRETINOIN
ISOVUE-200, IOPAMIDOL
ISOVUE-250, IOPAMIDOL
ISOVUE-300, IOPAMIDOL
ISOVUE-370, IOPAMIDOL
ISOVUE-M 200, IOPAMIDOL
ISOVUE-M 300, IOPAMIDOL
ISRADIPINE, ISRADIPINE
ISTALOL, TIMOLOL MALEATE
ISTODAX, ROMIDEPSIN
ISTURISA, OSILODROSTAT PHOSPHATE

APPENDIX A - PRODUCT NAME INDEX

** I **

ITRACONAZOLE, ITRACONAZOLE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 IVERMECTIN, IVERMECTIN (OTC)
 IVERMECTIN, IVERMECTIN
 IVY BLOCK, BENTOQUATAM (OTC)
 IWILFIN, EFLORNITHINE HYDROCHLORIDE
 IXEMPRA KIT, IXABEPILONE
 IYUZEH, LATANOPROST
 IZERVAY, AVACINCAPTAD PEGOL SODIUM

** J **

JADENU, DEFERASIROX
 JADENU SPRINKLE, DEFERASIROX
 JAIMIESS, ETHINYL ESTRADIOL
 JAKAFI, RUXOLITINIB PHOSPHATE
 JALYN, DUTASTERIDE
 JANTOVEN, WARFARIN SODIUM
 JANUMET, METFORMIN HYDROCHLORIDE
 JANUMET XR, METFORMIN HYDROCHLORIDE
 JANUVIA, SITAGLIPTIN PHOSPHATE
 JARDIANCE, EMPAGLIFLOZIN
 JATENZO, TESTOSTERONE UNDECANOATE
 JAYPIRCA, PIRTOBRUTINIB
 JELMYTO, MITOMYCIN
 JENCYCLA, NORETHINDRONE
 JENTADUETO, LINAGLIPTIN
 JENTADUETO XR, LINAGLIPTIN
 JESDUVROQ, DAPRODUSTAT
 JEVTANA KIT, CABAZITAXEL
 JOENJA, LENIOLISIB PHOSPHATE
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE
 JUBLIA, EFINACONAZOLE
 JULUCA, DOLUTEGRAVIR SODIUM
 JUNEL 1.5/30, ETHINYL ESTRADIOL
 JUNEL 1/20, ETHINYL ESTRADIOL
 JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 JUNEL FE 1/20, ETHINYL ESTRADIOL
 JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 JUXTAPID, LOMITAPIDE MESYLATE
 JYLAMVO, METHOTREXATE
 JYNARQUE, TOLVAPTAN

** K **

KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KAITLIB FE, ETHINYL ESTRADIOL
 KALETRA, LOPINAVIR
 KALEXATE, SODIUM POLYSTYRENE SULFONATE
 KALLIGA, DESOGESTREL
 KALYDECO, IVACAFTOR
 KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
 KARBINAL ER, CARBINOXAMINE MALEATE
 KARIVA, DESOGESTREL
 KATERZIA, AMLODIPINE BENZOATE
 KAZANO, ALOGLIPTIN BENZOATE
 KELNOR, ETHINYL ESTRADIOL
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENALOG-80, TRIAMCINOLONE ACETONIDE
 KENGREAL, CANGRELOR
 KEPPRA, LEVETIRACETAM
 KEPPRA XR, LEVETIRACETAM
 KERENDIA, FINERENONE
 KERYDIN, TAVABOROLE
 KETALAR, KETAMINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** K **

KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 KETOZOLE, KETOCONAZOLE
 KEVEYIS, DICHLORPHENAMIDE
 KHAPZORY, LEVOLEUCOVORIN
 KIMIDESS, DESOGESTREL
 KIMYRSA, ORITAVANCIN DIPHOSPHATE
 KINEVAC, SINCALIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 KISQALI, RIBOCICLIB SUCCINATE
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KITABIS PAK, TOBRAMYCIN
 KLARON, SULFACETAMIDE SODIUM
 KLISYRI, TIRBANIBULIN
 KLONOPIN, CLONAZEPAM
 KLOR-CON, POTASSIUM CHLORIDE
 KLOR-CON M10, POTASSIUM CHLORIDE
 KLOR-CON M15, POTASSIUM CHLORIDE
 KLOR-CON M20, POTASSIUM CHLORIDE
 KLOXXADO, NALOXONE HYDROCHLORIDE
 KONVOMEK, OMEPRAZOLE
 KORLYM, MIFEPRISTONE
 KORSUVA, DIFELIKEFALIN ACETATE
 KOSELUGO, SELUMETINIB SULFATE
 KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE
 KRAZATI, ADAGRASIB
 KRINTAFEL, TAFENOQUINE SUCCINATE
 KURVELO, ETHINYL ESTRADIOL
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE
 KYBELLA, DEOXYCHOLIC ACID
 KYLEENA, LEVONORGESTREL
 KYPROLIS, CARFILZOMIB
 KYZATREX, TESTOSTERONE UNDECANOATE

** L **

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN DEXTROSE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE, LABETALOL HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 LACRISERT, HYDROXYPROPYL CELLULOSE
 LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTULOSE, LACTULOSE
 LAMICTAL, LAMOTRIGINE
 LAMICTAL CD, LAMOTRIGINE
 LAMICTAL ODT, LAMOTRIGINE
 LAMICTAL XR, LAMOTRIGINE
 LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMISIL AT, TERBINAFINE (OTC)
 LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMIVUDINE, LAMIVUDINE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LAMPIT, NIFURTIMOX
 LANORINAL, ASPIRIN
 LANOXIN, DIGOXIN
 LANOXIN PEDIATRIC, DIGOXIN
 LANREOTIDE ACETATE, LANREOTIDE ACETATE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN

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** L **

LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LAPATINIB DITOSYLATE, LAPATINIB DITOSYLATE
 LARIN 1.5/30, ETHINYL ESTRADIOL
 LARIN 1/20, ETHINYL ESTRADIOL
 LARIN 24 FE, ETHINYL ESTRADIOL
 LARIN FE 1.5/30, ETHINYL ESTRADIOL
 LARIN FE 1/20, ETHINYL ESTRADIOL
 LAROTID, AMOXICILLIN
 LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LASIX, FUROSEMIDE
 LASTACAFT, ALCAFTADINE (OTC)
 LATANOPROST, LATANOPROST
 LATISSE, BIMATOPROST
 LATUDA, LURASIDONE HYDROCHLORIDE
 LEFLUNOMIDE, LEFLUNOMIDE
 LENALIDOMIDE, LENALIDOMIDE
 LENVIMA, LENVATINIB MESYLATE
 LEQVIO, INCLISIRAN SODIUM
 LERIBANE, ETHINYL ESTRADIOL
 LESCOL XL, FLUVASTATIN SODIUM
 LESSINA-28, ETHINYL ESTRADIOL
 LETAIRIS, AMBRISENTAN
 LETROZOLE, LETROZOLE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUKERAN, CHLORAMBUCIL
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEUPROLIDE ACETATE FOR DEPOT SUSPENSION, LEUPROLIDE ACETATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVO-T, LEVOTHYROXINE SODIUM **
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOCARNITINE SF, LEVOCARNITINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOCETIRIZINE HYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOLET, LEVOTHYROXINE SODIUM **
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LEVOMILNACIPRAN HYDROCHLORIDE, LEVOMILNACIPRAN HYDROCHLORIDE
 LEVONEST, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 LEVOPHED, NOREPINEPHRINE BITARTRATE
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LEVOXYL, LEVOTHYROXINE SODIUM **
 LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE
 LEXAPRO, ESCITALOPRAM OXALATE
 LEXETTE, HALOBETASOL PROPIONATE
 LEXISCAN, REGADENOSON
 LEXIVA, FOSAMPRENAVIR CALCIUM
 LIALDA, MESALAMINE
 LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LICART, DICLOFENAC EPOLAMINE
 LIDEX, FLUOCINONIDE
 LIDOCAINE, LIDOCAINE

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** L **

LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LIDODERM, LIDOCAINE
 LIFITEGRAST, LIFITEGRAST
 LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
 LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
 LIKMEZ, METRONIDAZOLE
 LILETTA, LEVONORGESTREL
 LINAGLIPTIN, LINAGLIPTIN
 LINAGLIPTIN AND METFORMIN HYDROCHLORIDE, LINAGLIPTIN
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
 LINZESS, LINACLOTIDE
 LIORESAL, BACLOFEN
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LIPIODOL, ETHIODIZED OIL
 LIPITOR, ATORVASTATIN CALCIUM
 LIPOFEN, FENOFIBRATE
 LIQREV, SILDENAFIL CITRATE
 LIQUID E-Z-PAQUE, BARIUM SULFATE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LISINOPRIL, LISINOPRIL
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LITFULO, RITLECITINIB TOSYLATE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LITHIUM CITRATE, LITHIUM CITRATE
 LITHOBID, LITHIUM CARBONATE
 LITHOSTAT, ACETOHYDROXAMIC ACID
 LIVALO, PITAVASTATIN CALCIUM
 LIVMARLI, MARALIXIBAT CHLORIDE
 LIVTENCITY, MARIBAVIR
 LO LOESTRIN FE, ETHINYL ESTRADIOL
 LO SIMPESE, ETHINYL ESTRADIOL
 LO-MALMOREDE, ETHINYL ESTRADIOL
 LO-ZUMANDIMINE, DROSPIRENONE
 LOCAMETZ, GALLIUM GA-68 GOZETOTIDE
 LOCHOLEST, CHOLESTYRAMINE
 LOCHOLEST LIGHT, CHOLESTYRAMINE
 LOCOID, HYDROCORTISONE BUTYRATE
 LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
 LODOCO, COLCHICINE
 Lodosyn, CARBIDOPA
 LOGILIA, ULIPRISTAL ACETATE
 LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
 LOMAIRA, PHENTERMINE HYDROCHLORIDE
 LOMOTIL, ATROPINE SULFATE
 LONSURF, TIPIRACIL HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPID, GEMFIBROZIL
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LOPRESSOR, METOPROLOL TARTRATE

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** L **

LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
 LOPROX, CICLOPIROX
 LOPURIN, ALLOPURINOL
 LORATADINE, LORATADINE (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE REDIDOSE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LORAZEPAM INTENSOL, LORAZEPAM
 LORBRENA, LORLATINIB
 LOREEV XR, LORAZEPAM
 LORYNA, DROSPIRENONE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSEASONIQUE, ETHINYL ESTRADIOL
 LOTEMAX, LOTEPRDNOL ETABONATE
 LOTEMAX SM, LOTEPRDNOL ETABONATE
 LOTENSIN, BENAZEPRIL HYDROCHLORIDE
 LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 LOTEPRDNOL ETABONATE, LOTEPRDNOL ETABONATE
 LOTREL, AMLODIPINE BESYLATE
 LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
 LOTRONEX, ALOSETRON HYDROCHLORIDE
 LOVASTATIN, LOVASTATIN
 LOVAZA, OMEGA-3-ACID ETHYL ESTERS
 LOVENOX, ENOXAPARIN SODIUM
 LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 LUBIPROSTONE, LUBIPROSTONE
 LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 LUMAKRAS, SOTORASIB
 LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
 LUMI-SPORYN, BACITRACIN ZINC
 LUMIFY, BRIMONIDINE TARTRATE (OTC)
 LUMIGAN, BIMATOPROST
 LUMRYZ, SODIUM OXYBATE
 LUNESTA, ESZOPICLONE
 LUPKYNIS, VOCLOSPORIN
 LUPRON DEPOT, LEUPROLIDE ACETATE
 LUPRON DEPOT-PED KIT, LEUPROLIDE ACETATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 LUTATHERA, LUTETIUM LU 177 DOTATATE
 LUVOX, FLUVOXAMINE MALEATE
 LUZU, LULICONAZOLE
 LYBALVI, OLANZAPINE
 LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
 LYNPARZA, OLAPARIB
 LYPQOZET, ATORVASTATIN CALCIUM
 LYRICA, PREGABALIN
 LYRICA CR, PREGABALIN
 LYSODREN, MITOTANE
 LYSTEDA, TRANEXAMIC ACID
 LYTGobi, FUTIBATINIB
 LYVISPAH, BACLOFEN

** M **

M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MACITENTAN, MACITENTAN
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MALARONE, ATOVAQUONE
 MALARONE PEDIATRIC, ATOVAQUONE

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** M **

MALATHION, MALATHION
MALMOREDE, ETHINYL ESTRADIOL
MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 25%, MANNITOL
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
MARAVIROC, MARAVIROC
MARCAINE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARINOL, DRONABINOL
MARLISSA, ETHINYL ESTRADIOL
MARPLAN, ISOCARBOXAZID
MATULANE, PROCARBAZINE HYDROCHLORIDE
MAVENCLAD, CLADRIBINE
MAVYRET, GLECAPREVIR
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
MAXIDEX, DEXAMETHASONE
MAXITROL, DEXAMETHASONE
MAXZIDE, HYDROCHLOROTHIAZIDE
MAXZIDE-25, HYDROCHLOROTHIAZIDE
MAYZENT, SIPONIMOD
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
MEDROL, METHYLPREDNISOLONE
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MEFENAMIC ACID, MEFENAMIC ACID
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEGACE ES, MEGESTROL ACETATE
MEGESTROL ACETATE, MEGESTROL ACETATE
MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
MEKTOVI, BINIMETINIB
MELAMISA, DROSPIRENONE
MELOXICAM, MELOXICAM
MELPHALAN, MELPHALAN
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
MEN'S ROGAINE, MINOXIDIL (OTC)
MENEST, ESTROGENS, ESTERIFIED
MENOSTAR, ESTRADIOL
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
MEPRON, ATOVAQUONE
MERCAPTOPYRINE, MERCAPTOPYRINE
MEROPENEM, MEROPENEM
MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM
MERREM, MEROPENEM
MERZEE, ETHINYL ESTRADIOL
MESALAMINE, MESALAMINE
MESNA, MESNA
MESNEX, MESNA
MESTINON, PYRIDOSTIGMINE BROMIDE
METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
METASTRON, STRONTIUM CHLORIDE SR-89

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** M **

METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 METHADOSE, METHADONE HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHERGINE, METHYLERGONOVINE MALEATE
 METHIMAZOLE, METHIMAZOLE
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOXSALEN, METHOXSALEN
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METHSUXIMIDE, METHSUXIMIDE
 METHYLENE BLUE, METHYLENE BLUE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
 METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
 METHYLPHENIDATE, METHYLPHENIDATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOLAZONE, METOLAZONE
 METOPIRONE, METYRAPONE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
 METROCREAM, METRONIDAZOLE
 METROGEL, METRONIDAZOLE
 METROGEL-VAGINAL, METRONIDAZOLE
 METROLOTION, METRONIDAZOLE
 METRONIDAZOLE, METRONIDAZOLE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 METYROSINE, METYROSINE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIACALCIN, CALCITONIN SALMON
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 MICA FUNGIN, MICA FUNGIN SODIUM
 MICA FUNGIN IN SODIUM CHLORIDE 0.9%, MICA FUNGIN SODIUM
 MICA FUNGIN SODIUM, MICA FUNGIN SODIUM
 MICARDIS, TELMISARTAN
 MICARDIS HCT, HYDROCHLOROTHIAZIDE
 MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
 MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE
 MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICORT-HC, HYDROCORTISONE ACETATE
 MICROZIDE, HYDROCHLOROTHIAZIDE
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM IN 0.8% SODIUM CHLORIDE, MIDAZOLAM
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** M **

MIDOL LIQUID GELS, IBUPROFEN (OTC)
MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MIEBO, PERFLUOROHEXYLOCTANE
MIFEPREX, MIFEPRISTONE
MIFEPRISTONE, MIFEPRISTONE
MIGERGOT, CAFFEINE
MIGLITOL, MIGLITOL
MIGLUSTAT, MIGLUSTAT
MIGRANAL, DIHYDROERGOTAMINE MESYLATE
MILI, ETHINYL ESTRADIOL
MILRINONE LACTATE, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE
MINIPRESS, PRAZOSIN HYDROCHLORIDE
MINIVELLE, ESTRADIOL
MINOCIN, MINOCYCLINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOLIRA, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL, MINOXIDIL (OTC)
MINOXIDIL, MINOXIDIL
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
MIOCHOL-E, ACETYLCHOLINE CHLORIDE
MIOSTAT, CARBACHOL
MIRABEGRON, MIRABEGRON
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)
MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE
MIRENA, LEVONORGESTREL
MIRTAZAPINE, MIRTAZAPINE
MIRVASO, BRIMONIDINE TARTRATE
MISOPROSTOL, MISOPROSTOL
MITIGARE, COLCHICINE
MITIGO, MORPHINE SULFATE
MITOMYCIN, MITOMYCIN
MITOSOL, MITOMYCIN
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
MIVACURIUM CHLORIDE, MIVACURIUM CHLORIDE
MOBIC, MELOXICAM
MODAFINIL, MODAFINIL
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE
MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3, MICONAZOLE NITRATE (OTC)
MONISTAT 3, MICONAZOLE NITRATE
MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
MONISTAT 7, MICONAZOLE NITRATE (OTC)
MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONO-LINYAH, ETHINYL ESTRADIOL
MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC,
MONODOX, DOXYCYCLINE
MONOFERRIC, FERRIC DERISOMALTOSE
MONOKET, ISOSORBIDE MONONITRATE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
MONUROL, FOSFOMYCIN TROMETHAMINE
MORPHINE SULFATE, MORPHINE SULFATE
MOTTEGRITY, PRUCALOPRIDE SUCCINATE
MOTOFEN, ATROPINE SULFATE
MOTPOLY XR, LACOSAMIDE
MOTRIN IB, IBUPROFEN (OTC)
MOUNJARO, TIRZEPATIDE

APPENDIX A - PRODUCT NAME INDEX

** M **

MOVANTIK, NALOXEGOL OXALATE
 MOVIPREP, ASCORBIC ACID
 MOXIDECTIN, MOXIDECTIN
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN
 MOZOBIL, PLERIXAFOR
 MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
 MS CONTIN, MORPHINE SULFATE
 MUCINEX, GUAIFENESIN (OTC)
 MUCINEX D, GUAIFENESIN (OTC)
 MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 MULPLETA, LUSUTROMBOPAG
 MULTAQ, DRONEDARONE HYDROCHLORIDE
 MULTIHANCE, GADOBENATE DIMEGLUMINE
 MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 5.5, MAGNESIUM CHLORIDE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 7.4, MAGNESIUM CHLORIDE
 MULTRYS, CUPRIC SULFATE
 MUPIROCIN, MUPIROCIN
 MUPIROCIN, MUPIROCIN CALCIUM
 MUSE, ALPROSTADIL
 MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
 MYCAMINE, MICA FUNGIN SODIUM
 MYCAPSSA, OCTREOTIDE ACETATE
 MYCOBUTIN, RIFABUTIN
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 MYDAYIS, AMPHETAMINE ASPARTATE
 MYDCOMBI, PHENYLEPHRINE HYDROCHLORIDE
 MYDRIACYL, TROPICAMIDE
 MYFEMBREE, ESTRADIOL
 MYFORTIC, MYCOPHENOLIC SODIUM
 MYKACET, NYSTATIN
 MYLERAN, BUSULFAN
 MYORISAN, ISOTRETINOIN
 MYOVIEV, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYOVIEV 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYRBETRIQ, MIRABEGRON
 MYRBETRIQ GRANULES, MIRABEGRON
 MYSOLINE, PRIMIDONE
 MYTESI, CROFELEMER
 MYZILRA, ETHINYL ESTRADIOL

** N **

NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALFON, FENOPROFEN CALCIUM
 NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM
 NALMEFENE HYDROCHLORIDE, NALMEFENE HYDROCHLORIDE
 NALOXONE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE (OTC)
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALTREXONE, NALTREXONE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMENDA XR, MEMANTINE HYDROCHLORIDE
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

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** N **

NAPRELAN, NAPROXEN SODIUM
 NAPROSYN, NAPROXEN
 NAPROXEN, NAPROXEN
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NARCAN, NALOXONE HYDROCHLORIDE (OTC)
 NARDIL, PHENELZINE SULFATE
 NAROPIN, ROPIVACAINE HYDROCHLORIDE
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
 NASCOBAL, CYANOCOBALAMIN
 NASONEX 24HR ALLERGY, MOMETASONE FUROATE (OTC)
 NATACYN, NATAMYCIN
 NATAZIA, DIENOGEST
 NATEGLINIDE, NATEGLINIDE
 NATESTO, TESTOSTERONE
 NATROBA, SPINOSAD
 NAYZILAM, MIDAZOLAM
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NEBUPENT, PENTAMIDINE ISETHIONATE
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NELARABINE, NELARABINE
 NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
 NEO-SYNALAR, FLUOCINOLONE ACETONIDE
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NEOPROFEN, IBUPROFEN LYSINE
 NEORAL, CYCLOSPORINE
 NEOSPORIN, GRAMICIDIN
 NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NEPHROSCAN, TECHNETIUM TC-99M SUCCIMER
 NERLYNX, NERATINIB MALEATE
 NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
 NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
 NESINA, ALOGLIPTIN BENZOATE
 NETSPOT, GALLIUM DOTATATE GA-68
 NEUPRO, ROTIGOTINE
 NEURACEQ, FLORBETABEN F-18
 NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
 NEURONTIN, GABAPENTIN
 NEVANAC, NEPAFENAC
 NEVIRAPINE, NEVIRAPINE
 NEXAVAR, SORAFENIB TOSYLATE
 NEXESTA FE, ETHINYL ESTRADIOL
 NEXICLON XR, CLONIDINE
 NEXIUM, ESOMEPRAZOLE MAGNESIUM
 NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)
 NEXIUM IV, ESOMEPRAZOLE SODIUM
 NEXLETOL, BEMPEDOIC ACID
 NEXLIZET, BEMPEDOIC ACID
 NEXPLANON, ETNOGESTREL
 NEXTERONE, AMIODARONE HYDROCHLORIDE
 NEXTSTELLIS, DROSPIRENONE
 NIACIN, NIACIN
 NIACOR, NIACIN

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** N **

NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
 NICODERM CQ, NICOTINE (OTC)
 NICORETTE, NICOTINE POLACRILEX (OTC)
 NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
 NICOTINE, NICOTINE (OTC)
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NICOTROL, NICOTINE
 NIFEDIPINE, NIFEDIPINE
 NIKKI, DROSPIRENONE
 NILANDRON, NILUTAMIDE
 NILOTINIB HYDROCHLORIDE, NILOTINIB HYDROCHLORIDE
 NILUTAMIDE, NILUTAMIDE
 NIMBEX, CISATRACURIUM BESYLATE
 NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 NIMODIPINE, NIMODIPINE
 NINLARO, IXAZOMIB CITRATE
 NIPENT, PENTOSTATIN
 NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
 NISOLDIPINE, NISOLDIPINE
 NITAZOXANIDE, NITAZOXANIDE
 NITHIODOTE, SODIUM NITRITE
 NITISINONE, NITISINONE
 NITRO-DUR, NITROGLYCERIN
 NITROFURANTOIN, NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NITROGLYCERIN, NITROGLYCERIN
 NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
 NITROLINGUAL PUMPSPRAY, NITROGLYCERIN
 NITROMIST, NITROGLYCERIN
 NITROSTAT, NITROGLYCERIN
 NITYR, NITISINONE
 NIX, PERMETHRIN (OTC)
 NIZATIDINE, NIZATIDINE
 NIZORAL ANTI-DANDRUFF, KETOCONAZOLE (OTC)
 NOR-QD, NORETHINDRONE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE, NOREPINEPHRINE BITARTRATE
 NORETHINDRONE, NORETHINDRONE
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORITATE, METRONIDAZOLE
 NORLIQVA, AMLODIPINE BESYLATE
 NORMOCARB HF 25, MAGNESIUM CHLORIDE
 NORMOCARB HF 35, MAGNESIUM CHLORIDE
 NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
 NORTHERA, DROXIDOPA
 NORTREL 0.5/35-28, ETHINYL ESTRADIOL
 NORTREL 1/35-21, ETHINYL ESTRADIOL
 NORTREL 1/35-28, ETHINYL ESTRADIOL
 NORTREL 7/7/7, ETHINYL ESTRADIOL
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NORVASC, AMLODIPINE BESYLATE

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** N **

NORVIR, RITONAVIR
 NOURIANZ, ISTRADefYLLINE
 NOXAFIL, POSACONAZOLE
 NOXAFIL POWDERMIX KIT, POSACONAZOLE
 NOXIVENT, NITRIC OXIDE
 NUBEQA, DAROLUTAMIDE
 NUCYNТА, TAPENTADOL HYDROCHLORIDE
 NUCYNТА ER, TAPENTADOL HYDROCHLORIDE
 NUEDEXТА, DEXTROMETHORPHAN HYDROBROMIDE
 NULIBRY, FOSDENOPTERIN HYDROBROMIDE
 NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350
 NUMBRINO, COCAINE HYDROCHLORIDE
 NUPLAZID, PIMAVANSERIN TARTRATE
 NURTEC ODT, RIMEGEPANT SULFATE
 NUTRILIPID 10%, SOYBEAN OIL
 NUTRILIPID 20%, SOYBEAN OIL
 NUVARING, ETHINYL ESTRADIOL
 NUVESSA, METRONIDAZOLE
 NUVIGIL, ARMODAFINIL
 NUZYRA, OMADACYCLINE TOSYLATE
 NYLIA 1/35, ETHINYL ESTRADIOL
 NYLIA 7/7/7, ETHINYL ESTRADIOL
 NYMALIZE, NIMODIPINE
 NYSTATIN, NYSTATIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTOP, NYSTATIN

** O **

OCALIVA, OBETICHOLIC ACID
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCUFLOX, OFLOXACIN
 ODEFSEY, EMTRICITABINE
 ODOMZO, SONIDEGIB PHOSPHATE
 OFEV, NINTEDANIB ESYLATE
 OFLOXACIN, OFLOXACIN
 OGEN 5, ESTROPIPATE
 OGSIVEO, NIROGACESTAT HYDROBROMIDE
 OJJAARA, MOMELOTINIB DIHYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLINVYK, OLICERIDINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OLPRUVA, SODIUM PHENYLBUTYRATE
 OLUMIANT, BARICITINIB
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEGAVEN, FISH OIL TRIGLYCERIDES
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMIDRIA, KETOROLAC TROMETHAMINE
 OMNARIS, CICLESONIDE
 OMNIPAQUE 12, IOHEXOL
 OMNIPAQUE 140, IOHEXOL
 OMNIPAQUE 180, IOHEXOL
 OMNIPAQUE 240, IOHEXOL
 OMNIPAQUE 300, IOHEXOL

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** O **

OMNIPAQUE 350, IOHEXOL
OMNIPAQUE 9, IOHEXOL
OMNIPRED, PREDNISOLONE ACETATE
OMNISCAN, GADODIAMIDE
ONDANSETRON, ONDANSETRON
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONEXTON, BENZOYL PEROXIDE
ONFI, CLOBAZAM
ONGENTYS, OPICAPONE
ONIVYDE, IRINOTECAN HYDROCHLORIDE
ONPATTRO, PATISIRAN SODIUM
ONSURA, ETHINYL ESTRADIOL
ONUREG, AZACITIDINE
ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
OPCICON ONE-STEP, LEVONORGESTREL (OTC)
OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
OPFOLDA, MIGLUSTAT
OPSUMIT, MACITENTAN
OPTIRAY 300, IOVERSOL
OPTIRAY 320, IOVERSOL
OPTIRAY 350, IOVERSOL
OPTISON, ALBUMIN HUMAN
OPVEE, NALMEFENE HYDROCHLORIDE
OPZELURA, RUXOLITINIB PHOSPHATE
ORABLOC, ARTICAIN HYDROCHLORIDE
ORACEA, DOXYCYCLINE
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
ORAQIX, LIDOCAINE
ORAVERSE, PHENTOLAMINE MESYLATE
ORAVIG, MICONAZOLE
ORBACTIV, ORITAVANCIN DIPHOSPHATE
ORENITRAM, TREPROSTINIL DIOLAMINE
ORFADIN, NITISINONE
ORGOVYX, RELUGOLIX
ORIAHNN (COPACKAGED), ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE
ORILISSA, ELAGOLIX SODIUM
ORKAMBI, IVACAFTOR
ORLADEYO, BEROTRALSTAT HYDROCHLORIDE
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
ORPHENGESIC, ASPIRIN
ORPHENGESIC FORTE, ASPIRIN
ORSERDU, ELACESTRANT DIHYDROCHLORIDE
ORVATEN, MIDODRINE HYDROCHLORIDE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
OSENI, ALOGLIPTIN BENZOATE
OSHIH, ETHINYL ESTRADIOL
OSMITROL 10% IN WATER, MANNITOL
OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMITROL 15% IN WATER, MANNITOL
OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMITROL 20% IN WATER, MANNITOL
OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMITROL 5% IN WATER, MANNITOL
OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMOLEX ER, AMANTADINE HYDROCHLORIDE
OSPHENA, OSPEMIFENE
OTEZLA, APREMILAST
OTICAIR, HYDROCORTISONE
OTOVEL, CIPROFLOXACIN HYDROCHLORIDE
OTREXUP, METHOTREXATE
OXACILLIN SODIUM, OXACILLIN SODIUM
OXALIPLATIN, OXALIPLATIN
OXAPROZIN, OXAPROZIN
OXAZEPAM, OXAZEPAM

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** O **

OXBRYTA, VOXELOTOR
 OXCARBAZEPINE, OXCARBAZEPINE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 OXISTAT, OXICONAZOLE NITRATE
 OXLUMO, LUMASIRAN SODIUM
 OXSORALEN-ULTRA, METHOXSALLEN
 OXTELLAR XR, OXCARBAZEPINE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYCONTIN, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 OXYTROL, OXYBUTYNYN
 OXYTROL FOR WOMEN, OXYBUTYNYN (OTC)
 OZEMPIC, SEMAGLUTIDE
 OZOBAX DS, BACLOFEN
 OZURDEX, DEXAMETHASONE

** P **

PACERONE, AMIODARONE HYDROCHLORIDE
 PACLITAXEL, PACLITAXEL
 PALBOCICLIB, PALBOCICLIB
 PALIPERIDONE, PALIPERIDONE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PANDEL, HYDROCORTISONE PROBUTATE
 PANRETIN, ALITRETINOIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARAGARD T 380A, COPPER
 PARICALCITOL, PARICALCITOL
 PARLODEL, BROMOCRIPTINE MESYLATE
 PARNATE, TRANYLCPROMINE SULFATE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 PAROXETINE, PAROXETINE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 PARSABIV, ETELICALCETIDE
 PASER, AMINOSALICYLIC ACID
 PATADAY ONCE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PATADAY TWICE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PAXIL, PAROXETINE HYDROCHLORIDE
 PAXIL CR, PAROXETINE HYDROCHLORIDE
 PAXLOVID (COPACKAGED), NIRMATRELVIR
 PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
 PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
 PEDMARK, SODIUM THIOSULFATE
 PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND
 PEMAZYRE, PEMIGATINIB
 PEMETREXED, PEMETREXED
 PEMETREXED, PEMETREXED DISODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PEMETREXED DITROMETHAMINE, PEMETREXED DITROMETHAMINE
 PEMFEXY, PEMETREXED
 PENCICLOVIR, PENCICLOVIR
 PENICILLAMINE, PENICILLAMINE
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
 PENICILLIN G PROCAINE, PENICILLIN G PROCAINE

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** P **

PENICILLIN G SODIUM, PENICILLIN G SODIUM
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
PENICILLIN-VK, PENICILLIN V POTASSIUM
PENNSAID, DICLOFENAC SODIUM
PENTAM, PENTAMIDINE ISETHIONATE
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
PENTASA, MESALAMINE
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
PENTOSTATIN, PENTOSTATIN
PENTOXIFYLLINE, PENTOXIFYLLINE
PEPCID AC, FAMOTIDINE (OTC)
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
PERCOCET, ACETAMINOPHEN
PERCODAN, ASPIRIN
PERFOROMIST, FORMOTEROL FUMARATE
PERIDEX, CHLORHEXIDINE GLUCONATE
PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
PERIOGARD, CHLORHEXIDINE GLUCONATE
PERMETHRIN, PERMETHRIN (OTC)
PERMETHRIN, PERMETHRIN
PERPHENAZINE, PERPHENAZINE
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PERSANTINE, DIPYRIDAMOLE
PERSERIS KIT, RISPERIDONE
PFIZERPEN, PENICILLIN G POTASSIUM
PHEBURANE, SODIUM PHENYLBUTYRATE
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENELZINE SULFATE, PHENELZINE SULFATE
PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PHENYTEK, PHENYTOIN SODIUM
PHENYTOIN, PHENYTOIN
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PHEXXI, CITRIC ACID
PHILITH, ETHINYL ESTRADIOL
PHOSLO GELCAPS, CALCIUM ACETATE
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
PHOTOFRIN, PORFIMER SODIUM
PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE
PHOTREXA VISCOS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE
PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PHYRAGO, DASATINIB
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYTONADIONE, PHYTONADIONE
PIFELTRO, DORAVIRINE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PIMECROLIMUS, PIMECROLIMUS
PIMOZIDE, PIMOZIDE
PIMTREA, DESOGESTREL
PINDOLOL, PINDOLOL
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PIPERACILLIN, PIPERACILLIN SODIUM
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PIQRAY, ALPELISIB
PIRFENIDONE, PIRFENIDONE
PIROXICAM, PIROXICAM

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** P **

PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 PITOCIN, OXYTOCIN
 PLAN B ONE-STEP, LEVONORGESTREL (OTC)
 PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
 PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PLAVIX, CLOPIDOGREL BISULFATE
 PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PLENVU, ASCORBIC ACID
 PLERIXAFOR, PLERIXAFOR
 PLUVICTO, LUTETIUM LU-177 VIPIVOTIDE TETRAKETAN
 PODOFILOX, PODOFILOX
 POKONZA, POTASSIUM CHLORIDE
 POLMON, DEXCHLORPHENIRAMINE MALEATE
 POLOCAINE, MEPIVACAINE HYDROCHLORIDE
 POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POMALYST, POMALIDOMIDE
 PONSTEL, MEFENAMIC ACID
 PONVORY, PONESIMOD
 PORTIA-28, ETHINYL ESTRADIOL
 POSACONAZOLE, POSACONAZOLE
 POSIMIR, BUPIVACAINE
 POSLUMA, FLOTUFOLASTAT F-18 GALLIUM
 POTASSIUM ACETATE, POTASSIUM ACETATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE

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POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 10MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 POVIDONE IODINE, POVIDONE-IODINE (OTC)
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAMOSONE, HYDROCORTISONE ACETATE

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PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRASUGREL HYDROCHLORIDE, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZIQUANTEL, PRAZIQUANTEL
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PRE-OP, HEXACHLOROPHENE
 PRE-OP II, HEXACHLOROPHENE
 PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
 PRED FORTE, PREDNISOLONE ACETATE
 PRED MILD, PREDNISOLONE ACETATE
 PREDNICARBATE, PREDNICARBATE
 PREDNISOLONE, PREDNISOLONE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISON, PREDNISON
 PREDNISON INTENSOL, PREDNISON
 PREGABALIN, PREGABALIN
 PRELONE, PREDNISOLONE
 PREMARIN, ESTROGENS, CONJUGATED
 PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMPHASE 14/14, ESTROGENS, CONJUGATED
 PREMPRO, ESTROGENS, CONJUGATED
 PREPIDIL, DINOPROSTONE
 PRESTALIA, AMLODIPINE BESYLATE
 PRETOMANID, PRETOMANID
 PREVACID, LANSOPRAZOLE
 PREVACID 24 HR, LANSOPRAZOLE (OTC)
 PREVALITE, CHOLESTYRAMINE
 PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 PREVDUO, GLYCOPYRROLATE
 PREVIFEM, ETHINYL ESTRADIOL
 PREVMIS, LETERMOVIR
 PREZCOBIX, COBICISTAT
 PREZISTA, DARUNAVIR
 PRIALT, ZICONOTIDE ACETATE
 PRIFTIN, RIFAPENTINE
 PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE
 PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
 PRILOSEC, OMEPRAZOLE MAGNESIUM
 PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
 PRIMAQUINE, PRIMAQUINE PHOSPHATE
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
 PRIMATENE MIST, EPINEPHRINE (OTC)
 PRIMAXIN, CILASTATIN SODIUM
 PRIMIDONE, PRIMIDONE
 PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISTIQ, DESVENLAFAXINE SUCCINATE
 PROAIR DIGIHALER, ALBUTEROL SULFATE
 PROAIR HFA, ALBUTEROL SULFATE
 PROAIR RESPICLICK, ALBUTEROL SULFATE
 PROBALAN, PROBENECID
 PROBENECID, PROBENECID
 PROBENECID AND COLCHICINE, COLCHICINE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 PROCARDIA, NIFEDIPINE
 PROCARDIA XL, NIFEDIPINE
 PROCHLORPERAZINE, PROCHLORPERAZINE

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PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROCOMP, PROCHLORPERAZINE MALEATE
 PROCTOFOAM HC, HYDROCORTISONE ACETATE
 PROCYSBI, CYSTEAMINE BITARTRATE
 PROGESTERONE, PROGESTERONE
 PROGLYCEM, DIAZOXIDE
 PROGRAF, TACROLIMUS
 PROHANCE, GADOTERIDOL
 PROHANCE MULTIPACK, GADOTERIDOL
 PROLENSA, BROMFENAC SODIUM
 PROMACTA, ELTROMBOPAG OLAMINE
 PROMACTA KIT, ELTROMBOPAG OLAMINE
 PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
 PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
 PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
 PROMETRIUM, PROGESTERONE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 PROPECIA, FINASTERIDE
 PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 PROSCAR, FINASTERIDE
 PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PROTONIX, PANTOPRAZOLE SODIUM
 PROTONIX IV, PANTOPRAZOLE SODIUM
 PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
 PROTOPIC, TACROLIMUS
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 PROVAYBLUE, METHYLENE BLUE
 PROVENTIL-HFA, ALBUTEROL SULFATE
 PROVERA, MEDROXYPROGESTERONE ACETATE
 PROVIGIL, MODAFINIL
 PROVOCHOLINE, METHACHOLINE CHLORIDE
 PROZAC, FLUOXETINE HYDROCHLORIDE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 PULMICORT FLEXHALER, BUDESONIDE
 PULMICORT RESPULES, BUDESONIDE
 PUR-WASH, PURIFIED WATER (OTC)
 PURIFIED CORTROPHIN GEL, CORTICOTROPIN
 PURINETHOL, MERCAPTOPYRINE
 PURIXAN, MERCAPTOPYRINE
 PYLARIFY, PIFLUFOLASTAT F-18
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 PYRIMETHAMINE, PYRIMETHAMINE
 PYRUKYND, MITAPIVAT SULFATE
 PYTEST, UREA, C-14

** Q **

QALSODY, TOFERSEN

APPENDIX A - PRODUCT NAME INDEX

** Q **

QBRELIS, LISINAPRIL
 QBREXZA, GLYCOPYRRONIUM TOSYLATE
 QDOLO, TRAMADOL HYDROCHLORIDE
 QELBREE, VILOXAZINE HYDROCHLORIDE
 QINLOCK, RIPRETINIB
 QLOSI, PILOCARPINE HYDROCHLORIDE
 QNASL, BECLOMETHASONE DIPROPIONATE
 QOLIANA, BRIMONIDINE TARTRATE
 QSYMIA, PHENTERMINE HYDROCHLORIDE
 QTERN, DAPAGLIFLOZIN
 QUALAQUIN, QUININE SULFATE
 QUARTETTE, ETHINYL ESTRADIOL
 QUASENSE, ETHINYL ESTRADIOL
 QUDEXY XR, TOPIRAMATE
 QUELICIN, SUCCINYLCHOLINE CHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE
 QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 QUININE SULFATE, QUININE SULFATE
 QULIPTA, ATOGEPANT
 QUTENZA, CAPSAICIN
 QUVIVIQ, DARIDOREXANT HYDROCHLORIDE
 QUZYTIR, CETIRIZINE HYDROCHLORIDE
 QVAR REDHALER, BECLOMETHASONE DIPROPIONATE

** R **

R-GENE 10, ARGININE HYDROCHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RADICAVA, EDARAVONE
 RADICAVA ORS, EDARAVONE
 RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RAMELTEON, RAMELTEON
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 RAPAFLO, SILODOSIN
 RAPAMUNE, SIROLIMUS
 RAPIVAB, PERAMIVIR
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RASUVO, METHOTREXATE
 RAVICTI, GLYCEROL PHENYL BUTYRATE
 RAYALDEE, CALCIFEDIOL
 RAYOS, PREDNISONE
 READI-CAT 2, BARIUM SULFATE
 READI-CAT 2 SMOOTHIE, BARIUM SULFATE
 READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)
 RECARBRIO, CILASTATIN SODIUM
 RECLAST, ZOLEDRONIC ACID
 RECORLEV, LEVOKETOCONAZOLE
 RECTIV, NITROGLYCERIN
 REGADENOSON, REGADENOSON
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 REGONOL, PYRIDOSTIGMINE BROMIDE
 RELENZA, ZANAMIVIR
 RELEXXII, METHYLPHENIDATE HYDROCHLORIDE
 RELISTOR, METHYLNALTREXONE BROMIDE
 RELPAX, ELETRIPTAN HYDROBROMIDE
 RELYVRIO, SODIUM PHENYL BUTYRATE
 REMERON, MIRTAZAPINE

APPENDIX A - PRODUCT NAME INDEX

** R **

REMERON SOLTAB, MIRTAZAPINE
REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
REMODULIN, TREPROSTINIL
RENACIDIN, CITRIC ACID
RENAGEL, SEVELAMER HYDROCHLORIDE
RENOVA, TRETINOIN
RENVELA, SEVELAMER CARBONATE
REPAGLINIDE, REPAGLINIDE
RESTASIS, CYCLOSPORINE
RESTASIS MULTIDOSE, CYCLOSPORINE
RESTORIL, TEMAZEPAM
RETEVMO, SELPERCATINIB
RETIN-A, TRETINOIN
RETIN-A MICRO, TRETINOIN
RETIN-A-MICRO, TRETINOIN
RETISERT, FLUOCINOLONE ACETONIDE
RETROVIR, ZIDOVUDINE
REVATIO, SILDENAFIL CITRATE
REVLIMID, LENALIDOMIDE
REVONTO, DANTROLENE SODIUM
REXULTI, BREXPIPIRAZOLE
REYATAZ, ATAZANAVIR SULFATE
REYVOW, LASMIDITAN SUCCINATE
REZIPRES, EPHEDRINE HYDROCHLORIDE
REZLIDHIA, OLUTASIDENIB
REZUROCK, BELUMOSUDIL MESYLATE
REZZAYO, REZAFUNGIN ACETATE
RHINOCORT ALLERGY, BUDESONIDE (OTC)
RHOFADE, OXYMETAZOLINE HYDROCHLORIDE
RHOPRESSA, NETARSUDIL MESYLATE
RIBAVIRIN, RIBAVIRIN
RIDAURA, AURANOFIN
RIFABUTIN, RIFABUTIN
RIFADIN, RIFAMPIN
RIFAMPIN, RIFAMPIN
RILUTEK, RILUZOLE
RILUZOLE, RILUZOLE
RIMACTANE, RIFAMPIN
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
RIMSO-50, DIMETHYL SULFOXIDE
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
RINVOQ, UPADACITINIB
RIOCIGUAT, RIOCIGUAT
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RISPERDAL, RISPERIDONE
RISPERDAL CONSTA, RISPERIDONE
RISPERIDONE, RISPERIDONE
RITALIN, METHYLPHENIDATE HYDROCHLORIDE
RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
RITONAVIR, RITONAVIR
RIVASTIGMINE, RIVASTIGMINE
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
RIVFLOZA, NEDOSIRAN SODIUM
RIVIVE, NALOXONE HYDROCHLORIDE (OTC)
RIZAFILM, RIZATRIPTAN BENZOATE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
ROBAXIN, METHOCARBAMOL
ROBINUL, GLYCOPYRROLATE
ROBINUL FORTE, GLYCOPYRROLATE
ROCALTROL, CALCITRIOL
ROCKLATAN, LATANOPROST
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
ROFLUMILAST, ROFLUMILAST
ROGAINE (FOR MEN), MINOXIDIL (OTC)
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)

APPENDIX A - PRODUCT NAME INDEX

** R **

ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 ROMIDEPSIN, ROMIDEPSIN
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 ROWASA, MESALAMINE
 ROXICODONE, OXYCODONE HYDROCHLORIDE
 ROXYBOND, OXYCODONE HYDROCHLORIDE
 ROZEREM, RAMELTEON
 ROZLYTREK, ENTRECTINIB
 RUBRACA, RUCAPARIB CAMSYLATE
 RUBY-FILL, RUBIDIUM CHLORIDE RB-82
 RUFINAMIDE, RUFINAMIDE
 RUKOBIA, FOSTEMSAVIR TROMETHAMINE
 RYALTRIS, MOMETASONE FUROATE
 RYANODEX, DANTROLENE SODIUM
 RYBELSUS, SEMAGLUTIDE
 RYDAPT, MIDOSTAURIN
 RYKINDO, RISPERIDONE
 RYTARY, CARBIDOPA
 RYZUMVI, PHENTOLAMINE MESYLATE

** S **

SABRIL, VIGABATRIN
 SAFYRAL, DROSPIRENONE
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 SALONPAS, MENTHOL (OTC)
 SAMSCA, TOLVAPTAN
 SANCUSO, GRANISETRON
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SAPHRIS, ASENAPINE MALEATE
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SAVAYSA, EDOXABAN TOSYLATE
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 SAXENDA, LIRAGLUTIDE RECOMBINANT
 SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
 SCEMBLIX, ASCIMINIB HYDROCHLORIDE
 SCENESSE, AFAMELANOTIDE
 SCLEROSOL, TALC
 SCOPOLAMINE, SCOPOLAMINE
 SEASONALE, ETHINYL ESTRADIOL
 SEASONIQUE, ETHINYL ESTRADIOL
 SECUADO, ASENAPINE
 SEGLENTIS, CELECOXIB
 SEGLUOMET, ERTUGLIFLOZIN
 SEIZALAM, MIDAZOLAM HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SELENIOS ACID, SELENIOS ACID
 SELENIUM SULFIDE, SELENIUM SULFIDE
 SELEXIPAG, SELEXIPAG
 SELZENTRY, MARAVIROC
 SENSIPAR, CINACALCET HYDROCHLORIDE
 SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 SEPTOCAINE, ARTICAINE HYDROCHLORIDE
 SEPTRA, SULFAMETHOXAZOLE
 SEPTRA DS, SULFAMETHOXAZOLE
 SEREVENT, SALMETEROL XINAFOATE
 SERNIVO, BETAMETHASONE DIPROPIONATE
 SEROMYCIN, CYCLOSERINE
 SEROQUEL, QUETIAPINE FUMARATE
 SEROQUEL XR, QUETIAPINE FUMARATE

APPENDIX A - PRODUCT NAME INDEX

** S **

SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SETLAKIN, ETHINYL ESTRADIOL
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SEVOFLURANE, SEVOFLURANE
 SEYSARA, SARECYCLINE HYDROCHLORIDE
 SEZABY, PHENOBARBITAL SODIUM
 SFROWASA, MESALAMINE
 SIGNIFOR, PASIREOTIDE DIASPARTATE
 SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
 SIKLOS, HYDROXYUREA
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILENOR, DOXEPIN HYDROCHLORIDE
 SILODOSIN, SILODOSIN
 SILVADENE, SILVER SULFADIAZINE
 SIMBRINZA, BRIMONIDINE TARTRATE
 SIMLIYA, DESOGESTREL
 SIMPESSSE, ETHINYL ESTRADIOL
 SIMVASTATIN, SIMVASTATIN
 SINCALIDE, SINCALIDE
 SINE-AID IB, IBUPROFEN (OTC)
 SINEMET, CARBIDOPA
 SINGULAIR, MONTELUKAST SODIUM
 SINUVA, MOMETASONE FUROATE
 SIROLIMUS, SIROLIMUS
 SIRTURO, BEDAQUILINE FUMARATE
 SITAVIG, ACYCLOVIR
 SIVEXTRO, TEDIZOLID PHOSPHATE
 SKLICE, IVERMECTIN (OTC)
 SKYCLARYS, OMAVELOXOLONE
 SKYLA, LEVONORGESTREL
 SLYND, DROSPIRENONE
 SMOFLIPID 20%, FISH OIL
 SOANZ, TORSEMIDE
 SODIUM ACETATE, SODIUM ACETATE
 SODIUM BICARBONATE, SODIUM BICARBONATE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, FERRIC OXYHYDROXIDE
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
 SODIUM IODIDE I 131, SODIUM IODIDE I-131
 SODIUM NITRITE, SODIUM NITRITE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 SODIUM PHOSPHATES, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 SODIUM PICOSULFATE, MAGNESIUM OXIDE AND ANHYDROUS CITRIC ACID, CITRIC ACID
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 SODIUM TETRADECYL SULFATE, SODIUM TETRADECYL SULFATE
 SODIUM THIOSULFATE, SODIUM THIOSULFATE
 SOHONOS, PALOVAROTENE
 SOJOURN, SEVOFLURANE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SOLODYN, MINOCYCLINE HYDROCHLORIDE
 SOLOSEC, SECNIDAZOLE
 SOLTAMOX, TAMOXIFEN CITRATE

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** S **

SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
 SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
 SOLUPREP S, CHLORHEXIDINE GLUCONATE (OTC)
 SOMA, CARISOPRODOL
 SOMATULINE DEPOT, LANREOTIDE ACETATE
 SONATA, ZALEPLON
 SOOLANTRA, IVERMECTIN
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
 SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
 SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
 SORILUX, CALCIPOTRIENE
 SORINE, SOTALOL HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SOTRADECOL, SODIUM TETRADECYL SULFATE
 SOTYKTU, DEUCRAVACITINIB
 SOTYLIZE, SOTALOL HYDROCHLORIDE
 SOVALDI, SOFOSBUVIR
 SOVUNA, HYDROXYCHLOROQUINE SULFATE
 SPINRAZA, NUSINERSEN SODIUM
 SPIRIVA, TIOTROPIUM BROMIDE
 SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPORANOX, ITRACONAZOLE
 SPRAVATO, ESKETAMINE HYDROCHLORIDE
 SPRINTEC, ETHINYL ESTRADIOL
 SPRITAM, LEVETIRACETAM
 SPRIX, KETOROLAC TROMETHAMINE
 SPRYCEL, DASATINIB
 SPS, SODIUM POLYSTYRENE SULFONATE
 SPY AGENT GREEN KIT, INDOCYANINE GREEN
 SSD, SILVER SULFADIAZINE
 STALEVO 100, CARBIDOPA
 STALEVO 125, CARBIDOPA
 STALEVO 150, CARBIDOPA
 STALEVO 200, CARBIDOPA
 STALEVO 50, CARBIDOPA
 STALEVO 75, CARBIDOPA
 STEGLATRO, ERTUGLIFLOZIN
 STEGLUJAN, ERTUGLIFLOZIN
 STENDRA, AVANAFIL
 STERILE WATER, STERILE WATER FOR IRRIGATION
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 STERITALC, TALC
 STIE-CORT, HYDROCORTISONE
 STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
 STIVARGA, REGORAFENIB
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 STRIBILD, COBICISTAT
 STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE
 STROMEKTOL, IVERMECTIN
 STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89
 SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUCRALFATE, SUCRALFATE
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 SUFLAVE, MAGNESIUM SULFATE

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** S **

SUGAMMADEX SODIUM, SUGAMMADEX SODIUM
 SULAR, NISOLDIPINE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 SULFADIAZINE, SULFADIAZINE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULFAMYLON, MAFENIDE ACETATE
 SULFASALAZINE, SULFASALAZINE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 SUMATRIPTAN, SUMATRIPTAN
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUNITINIB MALATE, SUNITINIB MALATE
 SUNLENCA, LENACAPAVIR SODIUM
 SUNOSI, SOLRIAMFETOL HYDROCHLORIDE
 SUPPRELIN LA, HISTRELIN ACETATE
 SUPRANE, DESFLURANE
 SUPRAX, CEFIXIME
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE
 SUSTOL, GRANISETRON
 SUTAB, MAGNESIUM SULFATE
 SUTENT, SUNITINIB MALATE
 SYEDA, DROSPIRENONE
 SYFOVRE, PEGCETACOPLAN
 SYMBICORT, BUDESONIDE
 SYMBICORT AEROSPHERE, BUDESONIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 SYMDEKO (COPACKAGED), IVACAFTOR
 SYMFI, EFAVIRENZ
 SYMFI LO, EFAVIRENZ
 SYMJEPI, EPINEPHRINE
 SYMLIN, PRAMLINTIDE ACETATE
 SYMPAZAN, CLOBAZAM
 SYMPROIC, NALDEMEDINE TOSYLATE
 SYMTUZA, COBICISTAT
 SYNALAR, FLUOCINOLONE ACETONIDE
 SYNAREL, NAFARELIN ACETATE
 SYNDROS, DRONABINOL
 SYNERCID, DALFOPRISTIN
 SYNJARDY, EMPAGLIFLOZIN
 SYNJARDY XR, EMPAGLIFLOZIN
 SYNRIPO, OMACETAXINE MEPESUCCINATE
 SYNTHROID, LEVOTHYROXINE SODIUM **
 SYPRINE, TRIENTINE HYDROCHLORIDE

** T **

TABRECTA, CAPMATINIB HYDROCHLORIDE
 TACLONEX, BETAMETHASONE DIPROPIONATE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TADLIQ, TADALAFIL
 TAFINLAR, DABRAFENIB MESYLATE
 TAFLUPROST, TAFLUPROST
 TAGAMET HB, CIMETIDINE (OTC)
 TAGITOL V, BARIUM SULFATE
 TAGRISSO, OSIMERTINIB MESYLATE
 TALC, TALC
 TALICIA, AMOXICILLIN
 TALZENNA, TALAZOPARIB TOSYLATE
 TAMIFLU, OSELTAMIVIR PHOSPHATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TARGRETIN, BEXAROTENE
 TARPEYO, BUDESONIDE

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** T **

TASCENSO ODT, FINGOLIMOD LAURYL SULFATE
TASIGNA, NILOTINIB HYDROCHLORIDE
TASIMELTEON, TASIMELTEON
TASMAR, TOLCAPONE
TAUVID, FLORTAUCIPIR F-18
TAVABOROLE, TAVABOROLE
TAVALISSE, FOSTAMATINIB DISODIUM
TAVNEOS, AVACOPAN
TAXOTERE, DOCETAXEL
TAYTULLA, ETHINYL ESTRADIOL
TAZAROTENE, TAZAROTENE
TAZICEF, CEFTAZIDIME
TAZORAC, TAZAROTENE
TAZTIA XT, DILTIAZEM HYDROCHLORIDE
TAZVERIK, TAZEMETOSTAT HYDROBROMIDE
TECFIDERA, DIMETHYL FUMARATE
TECHNEGAS KIT, TECHNETIUM TC-99M LABELED CARBON
TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT
TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
TECHNETIUM TC 99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT
TECHNETIUM TC-99M MEDRONATE KIT, TECHNETIUM TC-99M MEDRONATE KIT
TECHNETIUM TC-99M SULFUR COLLOID KIT, TECHNETIUM TC-99M SULFUR COLLOID KIT
TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
TEFLARO, CEFTAROLINE FOSAMIL
TEGRETOL, CARBAMAZEPINE
TEGRETOL-XR, CARBAMAZEPINE
TEGSEDI, INOTERSEN SODIUM
TEKTRUNA, ALISKIREN HEMIFUMARATE
TELMISARTAN, TELMISARTAN
TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TEMAZEPAM, TEMAZEPAM
TEMBEXA, BRINCIDOFIVIR
TEMODAR, TEMOZOLOMIDE
TEMOZOLOMIDE, TEMOZOLOMIDE
TEMSIROLIMUS, TEMSIROLIMUS
TENOFVIR ALAFENAMIDE, TENOFVIR ALAFENAMIDE FUMARATE
TENOFVIR DISOPROXIL FUMARATE, TENOFVIR DISOPROXIL FUMARATE
TENORETIC 100, ATENOLOL
TENORETIC 50, ATENOLOL
TENORMIN, ATENOLOL
TEPADINA, THIOTEPA
TEPMETKO, TEPOTINIB HYDROCHLORIDE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
TERBUTALINE SULFATE, TERBUTALINE SULFATE
TERCONAZOLE, TERCONAZOLE
TERIFLUNOMIDE, TERIFLUNOMIDE
TERIL, CARBAMAZEPINE
TERIPARATIDE, TERIPARATIDE
TERLIVAZ, TERLIPRESSIN ACETATE
TESSALON, BENZONATATE
TESTIM, TESTOSTERONE
TESTOPEL, TESTOSTERONE
TESTOSTERONE, TESTOSTERONE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
TETRABENAZINE, TETRABENAZINE
TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** T **

TEXACORT, HYDROCORTISONE
 THALITONE, CHLORTHALIDONE
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 THALOMID, THALIDOMIDE
 THAM, TROMETHAMINE
 THEO-24, THEOPHYLLINE
 THEOCHRON, THEOPHYLLINE
 THEOPHYLLINE, THEOPHYLLINE
 THERMAZENE, SILVER SULFADIAZINE
 THEROXIDIL, MINOXIDIL (OTC)
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOGUANINE, THIOGUANINE
 THIOLA, TIOPRONIN
 THIOLA EC, TIOPRONIN
 THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 THIOTHIXENE, THIOTHIXENE
 THYQUIDITY, LEVOTHYROXINE SODIUM
 THYRO-TABS, LEVOTHYROXINE SODIUM **
 THYROSAFE, POTASSIUM IODIDE (OTC)
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 TIAZAC, DILTIAZEM HYDROCHLORIDE
 TIBSOVO, IVOSIDENIB
 TICAGRELOR, TICAGRELOR
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TIGLUTIK KIT, RILUZOLE
 TIKOSYN, DOFETILIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIMOPTIC, TIMOLOL MALEATE
 TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
 TIMOPTIC-XE, TIMOLOL MALEATE
 TINDAMAX, TINIDAZOLE
 TINIDAZOLE, TINIDAZOLE
 TIOCONAZOLE, TIOCONAZOLE (OTC)
 TIOPRONIN, TIOPRONIN
 TIOTROPIUM BROMIDE, TIOTROPIUM BROMIDE
 TIPIRACIL HYDROCHLORIDE AND TRIFLURIDINE, TIPIRACIL HYDROCHLORIDE
 TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
 TIROSINT, LEVOTHYROXINE SODIUM
 TIROSINT-SOL, LEVOTHYROXINE SODIUM
 TIS-U-SOL, MAGNESIUM SULFATE
 TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 TISSUEBLUE, BRILLIANT BLUE G
 TIVICAY, DOLUTEGRAVIR SODIUM
 TIVICAY PD, DOLUTEGRAVIR SODIUM
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TLANDO, TESTOSTERONE UNDECANOATE
 TOBI, TOBRAMYCIN
 TOBI PODHALER, TOBRAMYCIN
 TOBRADEX, DEXAMETHASONE
 TOBRADEX ST, DEXAMETHASONE
 TOBRAMYCIN, TOBRAMYCIN
 TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
 TOBEX, TOBRAMYCIN
 TOFACITINIB CITRATE, TOFACITINIB CITRATE
 TOFRANIL, IMIPRAMINE HYDROCHLORIDE
 TOLAK, FLUOROURACIL
 TOLCAPONE, TOLCAPONE
 TOLMETIN SODIUM, TOLMETIN SODIUM
 TOLSURA, ITRACONAZOLE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

APPENDIX A - PRODUCT NAME INDEX

** T **

TOLVAPTAN, TOLVAPTAN
TOPAMAX, TOPIRAMATE
TOPICORT, DESOXIMETASONE
TOPIRAMATE, TOPIRAMATE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TOPROL-XL, METOPROLOL SUCCINATE
TOREMIFENE CITRATE, TOREMIFENE CITRATE
TORISEL, TEMSIROLIMUS
TORSEMIDE, TORSEMIDE
TOSYMRA, SUMATRIPTAN
TOVIAZ, FESOTERODINE FUMARATE
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
TPOXX, TECOVIRIMAT
TRACLEER, BOSENTAN
TRADJENTA, LINAGLIPTIN
TRALEMENT, CUPRIC SULFATE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRANDOLAPRIL, TRANDOLAPRIL
TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
TRANEXAMIC ACID, TRANEXAMIC ACID
TRANSDERM SCOP, SCOPOLAMINE
TRANXENE, CLORAZEPATE DIPOTASSIUM
TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVATAN Z, TRAVOPROST
TRAVOPROST, TRAVOPROST
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TREANDA, BENDAMUSTINE HYDROCHLORIDE
TRECATOR, ETHIONAMIDE
TRELEGY ELLIPTA, FLUTICASONE FUROATE
TRELSTAR, TRIPTORELIN PAMOATE
TREPROSTINIL, TREPROSTINIL
TRETINOIN, TRETINOIN
TRETINOIN MICROSPHERE, TRETINOIN
Trexall, METHOTREXATE SODIUM
Treximet, NAPROXEN SODIUM
Trezix, ACETAMINOPHEN
TRI LO SPRINTEC, ETHINYL ESTRADIOL
TRI-ESTARYLLA, ETHINYL ESTRADIOL
TRI-LEGEST FE, ETHINYL ESTRADIOL
TRI-LINYAH, ETHINYL ESTRADIOL
TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
TRI-LO-LINYAH, ETHINYL ESTRADIOL
TRI-LO-MILI, ETHINYL ESTRADIOL
TRI-LUMA, FLUOCINOLONE ACETONIDE
TRI-MILI, ETHINYL ESTRADIOL
TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
TRI-SPRINTEC, ETHINYL ESTRADIOL
TRIACIN-C, CODEINE PHOSPHATE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRIAMTERENE, TRIAMTERENE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRIAZOLAM, TRIAZOLAM
TRIBENZOR, AMLODIPINE BESYLATE
TRICOR, FENOFIBRATE
TRIDERM, TRIAMCINOLONE ACETONIDE
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
TRIESENCE, TRIAMCINOLONE ACETONIDE
TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
TRIFLURIDINE, TRIFLURIDINE
TRIGLIDE, FENOFIBRATE

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** T **

TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIJARDY XR, EMPAGLIFLOZIN
 TRIKAFTA (COPACKAGED), ELEXACAFTOR, IVACAFTOR, TEZACAFTOR
 TRILEPTAL, OXCARBAZEPINE
 TRILIPIX, CHOLINE FENOFIBRATE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 TRIPTODUR KIT, TRIPTORELIN PAMOATE
 TRISENOX, ARSENIC TRIOXIDE
 TRIUMEQ, ABACAVIR SULFATE
 TRIUMEQ PD, ABACAVIR SULFATE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 TRIVORA-28, ETHINYL ESTRADIOL
 TROKENDI XR, TOPIRAMATE
 TROPHAMINE, AMINO ACIDS
 TROPHAMINE 10%, AMINO ACIDS
 TROPICACYL, TROPICAMIDE
 TROPICAMIDE, TROPICAMIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 TRUDHESA, DIHYDROERGOTAMINE MESYLATE
 TRULANCE, PLECANATIDE
 TRUQAP, CAPIVASERTIB
 TRUVADA, EMTRICITABINE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 TUKYSA, TUCATINIB
 TURALIO, PEXIDARTINIB HYDROCHLORIDE
 TURQOZ, ETHINYL ESTRADIOL
 TUXARIN ER, CHLORPHENIRAMINE MALEATE
 TWIRLA, ETHINYL ESTRADIOL
 TWYNEO, BENZOYL PEROXIDE
 TYBLUME, ETHINYL ESTRADIOL
 TYBOST, COBICISTAT
 TYDEMY, DROSPIRENONE
 TYGACIL, TIGECYCLINE
 TYKERB, LAPATINIB DITOSYLATE
 TYLENOL, ACETAMINOPHEN (OTC)
 TYMLOS, ABALOPARATIDE
 TYRVAYA, VARENICLINE TARTRATE
 TYVASO, TREPROSTINIL
 TYVASO DPI, TREPROSTINIL
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

** U **

UBRELVY, UBROGEPANT
 UCERIS, BUDESONIDE
 ULORIC, FEBUXOSTAT
 ULSPIRA, NITRIC OXIDE
 ULTANE, SEVOFLURANE
 ULTIVA, REMIFENTANIL HYDROCHLORIDE
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 ULTRAVATE, HALOBETASOL PROPIONATE
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 UNASYN, AMPICILLIN SODIUM
 UNISOM, DOXYLAMINE SUCCINATE (OTC)
 UNITHROID, LEVOTHYROXINE SODIUM **
 UPNEEQ, OXYMETAZOLINE HYDROCHLORIDE
 UPTRAVI, SELEXIPAG
 UREX, METHENAMINE HIPPURATE
 UROCIT-K, POTASSIUM CITRATE

APPENDIX A - PRODUCT NAME INDEX

** U **

UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 URSODIOL, URSODIOL
 UVADEX, METHOXSALEN
 UZEDY, RISPERIDONE

** V **

VABOMERE, MEROPENEM
 VAGIFEM, ESTRADIOL
 VAGISTAT-1, TIOCONAZOLE (OTC)
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE
 VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALIUM, DIAZEPAM
 VALNAC, BETAMETHASONE VALERATE
 VALPROATE SODIUM, VALPROATE SODIUM
 VALPROIC ACID, VALPROIC ACID
 VALRUBICIN, VALRUBICIN
 VALSARTAN, VALSARTAN
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSTAR PRESERVATIVE FREE, VALRUBICIN
 VALTOCO, DIAZEPAM
 VALTRES, VALACYCLOVIR HYDROCHLORIDE
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
 VANDAZOLE, METRONIDAZOLE
 VANFLYTA, QUIZARTINIB DIHYDROCHLORIDE
 VANOS, FLUOCINONIDE
 VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VARIBAR HONEY, BARIUM SULFATE
 VARIBAR NECTAR, BARIUM SULFATE
 VARIBAR PUDDING, BARIUM SULFATE
 VARIBAR THIN HONEY, BARIUM SULFATE
 VARIBAR THIN LIQUID, BARIUM SULFATE
 VARITHENA, POLIDOCANOL
 VARUBI, ROLAPITANT HYDROCHLORIDE
 VASCEPA, ICOSAPENT ETHYL
 VASERETIC, ENALAPRIL MALEATE
 VASOPRESSIN, VASOPRESSIN
 VASOPRESSIN IN SODIUM CHLORIDE 0.9%, VASOPRESSIN
 VASOSTRICT, VASOPRESSIN
 VASOTEC, ENALAPRIL MALEATE
 VAZALORE, ASPIRIN (OTC)
 VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE
 VECTICAL, CALCITRIOL
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VEKLURY, REMDESIVIR
 VELCADE, BORTEZOMIB
 VELETRI, EPOPROSTENOL SODIUM
 VELIVET, DESOGESTREL
 VELPHORO, FERRIC OXYHYDROXIDE
 VELSIPITY, ETASIMOD ARGININE
 VELTASSA, PATIROMER SORBITE CALCIUM
 VELTIN, CLINDAMYCIN PHOSPHATE
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VENCLEXTA, VENETOCLAX
 VENLAFAXINE BESYLATE, VENLAFAXINE BESYLATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VENOFER, FERRIC OXYHYDROXIDE

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** v **

VENTAVIS, ILOPROST
VENTOLIN HFA, ALBUTEROL SULFATE
VEOZAH, FEZOLINETANT
VERAMYST, FLUTICASONE FUROATE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VEREGEN, SINECATECHINS
VERELAN, VERAPAMIL HYDROCHLORIDE
VERELAN PM, VERAPAMIL HYDROCHLORIDE
VERKAZIA, CYCLOSPORINE
VERQUVO, VERICIGUAT
VERSACLOZ, CLOZAPINE
VERZENIO, ABEMACICLIB
VESICARE, SOLIFENACIN SUCCINATE
VESICARE LS, SOLIFENACIN SUCCINATE
VEVYE, CYCLOSPORINE
VFEND, VORICONAZOLE
VIAGRA, SILDENAFIL CITRATE
VIBATIV, TELAVANCIN HYDROCHLORIDE
VIBERZI, ELUXADOLINE
VIBISONE, CYANOCOBALAMIN
VIBRAMYCIN, DOXYCYCLINE
VIBRAMYCIN, DOXYCYCLINE CALCIUM
VIBRAMYCIN, DOXYCYCLINE HYCLATE
VICTOZA, LIRAGLUTIDE RECOMBINANT
VIDAZA, AZACITIDINE
VIENVA, ETHINYL ESTRADIOL
VIGABATRIN, VIGABATRIN
VIGADRONE, VIGABATRIN
VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
VIGPODER, VIGABATRIN
VIIBRYD, VILAZODONE HYDROCHLORIDE
VIJOICE, ALPELISIB
VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
VILTEPSO, VILTOLARSEN
VIMPAT, LACOSAMIDE
VINBLASTINE SULFATE, VINBLASTINE SULFATE
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
VIORELE, DESOGESTREL
VIRACEPT, NELFINAVIR MESYLATE
VIRAMUNE, NEVIRAPINE
VIRAMUNE XR, NEVIRAPINE
VIRAZOLE, RIBAVIRIN
VIREAD, TENOFOVIR DISOPROXIL FUMARATE
VIROPTIC, TRIFLURIDINE
VISINE, NAPHAZOLINE HYDROCHLORIDE (OTC)
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISIONBLUE, TRYPAN BLUE
VISIPAQUE 270, IODIXANOL
VISIPAQUE 320, IODIXANOL
VISTARIL, HYDROXYZINE PAMOATE
VISTOGARD, URIDINE TRIACETATE
VISUDYNE, VERTEPORFIN
VITAMIN D, ERGOCALCIFEROL
VITAMIN K1, PHYTONADIONE
VITRAKVI, LAROTRECTINIB SULFATE
VIVELLE-DOT, ESTRADIOL
VIVIMUSTA, BENDAMUSTINE HYDROCHLORIDE
VIVITROL, NALTREXONE
VIVJOA, OTESECONAZOLE
VIZAMYL, FLUTEMETAMOL F-18
VIZIMPRO, DACOMITINIB
VOCABRIA, CABOTEGRAVIR SODIUM
VOGELXO, TESTOSTERONE
VOLNEA, DESOGESTREL

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** V **

VOLTAREN ARTHRITIS PAIN, DICLOFENAC SODIUM (OTC)
 VONJO, PACRITINIB CITRATE
 VOQUEZNA, VONOPRAZAN FUMARATE
 VOQUEZNA DUAL PAK, AMOXICILLIN
 VOQUEZNA TRIPLE PAK, AMOXICILLIN
 VORICONAZOLE, VORICONAZOLE
 VOSEVI, SOFOSBUVIR
 VOSOL, ACETIC ACID, GLACIAL
 VOSOL HC, ACETIC ACID, GLACIAL
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 VOXZOGO, VOSORITIDE
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE
 VTAMA, TAPINAROF
 VUIITY, PILOCARPINE HYDROCHLORIDE
 VUMERITY, DIROXIMEL FUMARATE
 VUSION, MICONAZOLE NITRATE
 VYDUO, NEBIVOLOL HYDROCHLORIDE
 VYFEMLA, ETHINYL ESTRADIOL
 VYLEESI (AUTOINJECTOR), BREMELANOTIDE ACETATE
 VYNDAMAX, TAFAMIDIS
 VYNDAQEL, TAFAMIDIS MEGLUMINE
 VYONDYS 53, GOLODIRSEN
 VYTORIN, EZETIMIBE
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE
 VYXEOS, CYTARABINE
 VYZULTA, LATANOPROSTENE BUNOD

** W **

WAINUA, EPLONTERSEN SODIUM
 WAKIX, PITOLISANT HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 WEGOVY, SEMAGLUTIDE
 WELCHOL, COLESEVELAM HYDROCHLORIDE
 WELIREG, BELZUTIFAN
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 WERA, ETHINYL ESTRADIOL
 WINLEVI, CLASCOTERONE
 WIXELA INHUB, FLUTICASONE PROPIONATE
 WOMEN'S ROGAINE, MINOXIDIL (OTC)
 WYNZORA, BETAMETHASONE DIPROPIONATE

** X **

XACDURO (COPACKAGED), DURLOBACTAM SODIUM
 XACIATO, CLINDAMYCIN PHOSPHATE
 XADAGO, SAFINAMIDE MESYLATE
 XALATAN, LATANOPROST
 XALKORI, CRIZOTINIB
 XANAX, ALPRAZOLAM
 XANAX XR, ALPRAZOLAM
 XARACOLL, BUPIVACAINE HYDROCHLORIDE
 XARELTO, RIVAROXABAN
 XATMEP, METHOTREXATE SODIUM
 XCOPRI, CENOBAMATE
 XDEMVI, LOTILANER
 XELJANZ, TOFACITINIB CITRATE
 XELJANZ XR, TOFACITINIB CITRATE
 XELODA, CAPECITABINE
 XELPROS, LATANOPROST
 XELSTRYM, DEXTROAMPHETAMINE
 XENAZINE, TETRABENAZINE
 XENICAL, ORLISTAT
 XENLETA, LEFAMULIN ACETATE
 XENON XE 133, XENON XE-133
 XENOVIEW, XENON XE-129 HYPERPOLARIZED

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** X **

XEPI, OZENOXACIN
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE
 XERESE, ACYCLOVIR
 XERMELO, TELOTRISTAT ETIPRATE
 XHANCE, FLUTICASON PROPIONATE
 XIFAXAN, RIFAXIMIN
 XIGDUO XR, DAPAGLIFLOZIN
 XIIDRA, LIFITEGRAST
 XIPERE, TRIAMCINOLONE ACETONIDE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 XOFLUZA, BALOXAVIR MARBOXIL
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 XOPENEX HFA, LEVALBUTEROL TARTRATE
 XOSPATA, GILTERITINIB FUMARATE
 XPHOZAH, TENAPANOR HYDROCHLORIDE
 XPOVIO, SELINEXOR
 XTAMPZA ER, OXYCODONE
 XTANDI, ENZALUTAMIDE
 XULANE, ETHINYL ESTRADIOL
 XURIDEN, URIDINE TRIACETATE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE
 XYREM, SODIUM OXYBATE
 XYWAV, CALCIUM OXYBATE
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

** Y **

YAELA, DROSPIRENONE
 YARGESA, MIGLUSTAT
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE
 YCANTH, CANTHARIDIN
 YONDELIS, TRABECTEDIN
 YONSA, ABIRATERONE ACETATE
 YUPELRI, REVEFENACIN
 YUTIQ, FLUOCINOLONE ACETONIDE

** Z **

ZADITOR, KETOTIFEN FUMARATE (OTC)
 ZAFIRLUKAST, ZAFIRLUKAST
 ZALEPLON, ZALEPLON
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE
 ZANOSAR, STREPTOZOCIN
 ZARONTIN, ETHOSUXIMIDE
 ZAVESCA, MIGLUSTAT
 ZAVZPRET, ZAVEGEPANT HYDROCHLORIDE
 ZEGALOGUE, DASIGLUCAGON HYDROCHLORIDE
 ZEGALOGUE (AUTOINJECTOR), DASIGLUCAGON HYDROCHLORIDE
 ZEGERID, OMEPRAZOLE
 ZEGERID OTC, OMEPRAZOLE (OTC)
 ZEJULA, NIRAPARIB TOSYLATE
 ZELAPAR, SELEGILINE HYDROCHLORIDE
 ZELBORAF, VEMURAFENIB
 ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE
 ZEMDRI, PLAZOMICIN SULFATE
 ZEMPLAR, PARICALCITOL
 ZENATANE, ISOTRETINOIN
 ZEPATIER, ELBASVIR
 ZEPBOUND, TIRZEPATIDE
 ZEPOSIA, OZANIMOD HYDROCHLORIDE
 ZEPZELCA, LURBINECTEDIN
 ZERBAXA, CEFTOLOZANE SULFATE
 ZERVIAE, CETIRIZINE HYDROCHLORIDE
 ZESTORETIC, HYDROCHLOROTHIAZIDE

APPENDIX A - PRODUCT NAME INDEX

** Z **

ZESTRIL, LISINOPRIL
ZETIA, EZETIMIBE
ZETONNA, CICLESONIDE
ZIAC, BISOPROLOL FUMARATE
ZIAGEN, ABACAVIR SULFATE
ZIANA, CLINDAMYCIN PHOSPHATE
ZIDOVUDINE, ZIDOVUDINE
ZILBRYSQ, ZILUCOPLAN SODIUM
ZILEUTON, ZILEUTON
ZILRETTA, TRIAMCINOLONE ACETONIDE
ZILXI, MINOCYCLINE HYDROCHLORIDE
ZIMHI, NALOXONE HYDROCHLORIDE
ZINC CHLORIDE, ZINC CHLORIDE
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
ZINC SULFATE, ZINC SULFATE
ZINGO, LIDOCAINE HYDROCHLORIDE
ZIOPTAN, TAFLUPROST
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
ZIPSOR, DICLOFENAC POTASSIUM
ZIRGAN, GANCICLOVIR
ZITHROMAX, AZITHROMYCIN
ZITUVIMET, METFORMIN HYDROCHLORIDE
ZITUVIO, SITAGLIPTIN
ZOCOR, SIMVASTATIN
ZOKINVY, LONAFARNIB
ZOLADEX, GOSERELIN ACETATE
ZOLEDRONIC, ZOLEDRONIC ACID
ZOLEDRONIC ACID, ZOLEDRONIC ACID
ZOLINZA, VORINOSTAT
ZOLMITRIPTAN, ZOLMITRIPTAN
ZOLOFT, SERTRALINE HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOMIG, ZOLMITRIPTAN
ZONALON, DOXEPIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE
ZONISADE, ZONISAMIDE
ZONISAMIDE, ZONISAMIDE
ZONTIVITY, VORAPAXAR SULFATE
ZORTRESS, EVEROLIMUS
ZORYVE, ROFLUMILAST
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOVIA 1/50E-28, ETHINYL ESTRADIOL
ZOVIRAX, ACYCLOVIR
ZTALMY, GANAXOLONE
ZTLIDO, LIDOCAINE
ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE
ZULRESSO, BREXANOLONE
ZUMANDIMINE, DROSPIRENONE
ZURAGARD, ISOPROPYL ALCOHOL (OTC)
ZURZUVAE, ZURANOLONE
ZYCLARA, IMIQIMOD
ZYDELIG, IDELALISIB
ZYFLO, ZILEUTON
ZYKADIA, CERITINIB
ZYLET, LOTEHPREDNOL ETABONATE
ZYLOPRIM, ALLOPURINOL
ZYMAXID, GATIFLOXACIN
ZYNRELEF KIT, BUPIVACAINE
ZYPITAMAG, PITAVASTATIN MAGNESIUM
ZYPREXA, OLANZAPINE
ZYPREXA RELPREVV, OLANZAPINE PAMOATE
ZYPREXA ZYDIS, OLANZAPINE
ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

APPENDIX A - PRODUCT NAME INDEX

**** Z ****

ZYTIGA, ABIRATERONE ACETATE
ZYVOX, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** 3 ******3D IMAGING DRUG**

- * 3D IMAGING DRUG DESIGN AND DEVELOPMENT LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

3M

- * 3M CO
PERIDEX, CHLORHEXIDINE GLUCONATE
- * 3M HEALTH CARE INC
AVAGARD, ALCOHOL (OTC)
DURAPREP, IODINE POVACRYLEX (OTC)

3M HEALTH CARE

- * 3M HEALTH CARE INFECTION PREVENTION DIV
SOLUPREP S, CHLORHEXIDINE GLUCONATE (OTC)

**** 6 ******60 DEGREES PHARMS**

- * 60 DEGREES PHARMACEUTICALS INC
ARAKODA, TAFENOQUINE SUCCINATE

**** A ******AAA USA INC**

- * ADVANCED ACCELERATOR APPLICATIONS USA INC
LUTATHERA, LUTETIUM LU 177 DOTATATE
NETSPOT, GALLIUM DOTATATE GA-68

AADI

- * AADI BIOSCIENCE INC
FYARRO, SIROLIMUS

AAIPHARMA LLC

- * AAIPHARMA LLC
AZASAN, AZATHIOPRINE

ABBVIE

- * ABBVIE
OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
- * ABBVIE INC
ACULAR LS, KETOROLAC TROMETHAMINE
ACULAR, KETOROLAC TROMETHAMINE
ACUVAIL, KETOROLAC TROMETHAMINE
ACZONE, DAPSONE
ALPHAGAN P, BRIMONIDINE TARTRATE
BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
BENTYL, DICYCLOMINE HYDROCHLORIDE
CANASA, MESALAMINE
CARAFATE, SUCRALFATE
CELEXA, CITALOPRAM HYDROBROMIDE
COMBIGAN, BRIMONIDINE TARTRATE
CYCLOSPORINE, CYCLOSPORINE
DALVANCE, DALBAVANCIN HYDROCHLORIDE
DELZICOL, MESALAMINE
DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
DUOPA, CARBIDOPA
DURYSTA, BIMATOPROST
FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
FML FORTE, FLUOROMETHOLONE
FML, FLUOROMETHOLONE
GENGRAF, CYCLOSPORINE
KALETRA, LOPINAVIR
LASTACAFIT, ALCAFTADINE (OTC)
LATISSE, BIMATOPROST
LEXAPRO, ESCITALOPRAM OXALATE
LINZESS, LINACLOTIDE
LUMIGAN, BIMATOPROST
MAVYRET, GLECAPREVIR
NAMENDA XR, MEMANTINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ABBVIE INC**

NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 NIMBEX, CISATRACURIUM BESYLATE
 NORVIR, RITONAVIR
 ORIAHNN (COPACKAGED), ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE
 ORILISSA, ELAGOLIX SODIUM
 OZURDEX, DEXAMETHASONE
 PRED FORTE, PREDNISOLONE ACETATE
 PRED MILD, PREDNISOLONE ACETATE
 QULIPTA, ATOGEPANT
 RECTIV, NITROGLYCERIN
 RESTASIS MULTIDOSE, CYCLOSPORINE
 RESTASIS, CYCLOSPORINE
 RINVOQ, UPADACITINIB
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 SYNTHROID, LEVOTHYROXINE SODIUM **
 TEFLARO, CEFTAROLINE FOSAMIL
 TRICOR, FENOFIBRATE
 TRILIPIX, CHOLINE FENOFIBRATE
 UBRELVY, UBROGEPANT
 ULTANE, SEVOFLURANE
 VENCLEXTA, VENETOCLAX
 VIBERZI, ELUXADOLINE
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE
 VUITY, PILOCARPINE HYDROCHLORIDE
 ZEMPLAR, PARICALCITOL
 ZYMAXID, GATIFLOXACIN

ABBVIE ENDOCRINE INC*** ABBVIE ENDOCRINE INC**

LUPRON DEPOT, LEUPROLIDE ACETATE
 LUPRON DEPOT-PED KIT, LEUPROLIDE ACETATE

ABHAI INC*** ABHAI INC**

DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ABHAI LLC*** ABHAI LLC**

ATOVAQUONE, ATOVAQUONE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LEFLUNOMIDE, LEFLUNOMIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 URSODIOL, URSODIOL

ABON PHARMS LLC*** ABON PHARMACEUTICALS LLC**

ATOVAQUONE, ATOVAQUONE
 CLOFARABINE, CLOFARABINE

ABRAXIS PHARM*** ABRAXIS PHARMACEUTICAL PRODUCTS**

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE

ACACIA*** ACACIA PHARMA LTD**

BARHEMSYS, AMISULPRIDE
 BYFAVO, REMIMAZOLAM BESYLATE

ACADIA PHARMS INC*** ACADIA PHARMACEUTICALS INC**

DAYBUE, TROFINETIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACADIA PHARMACEUTICALS INC
 NUPLAZID, PIMAVANSERIN TARTRATE

ACCELRX LABS

* ACCELRX LABS LLC
 CARISOPRODOL, CARISOPRODOL

ACCORD

* ACCORD BIOPHARMA INC
 CAMCEVI KIT, LEUPROLIDE MESYLATE

ACCORD HLTHCARE

* ACCORD HEALTHCARE INC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ALLOPURINOL, ALLOPURINOL
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ANASTROZOLE, ANASTROZOLE
 APIXABAN, APIXABAN
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATROPINE SULFATE, ATROPINE SULFATE
 AZACITIDINE, AZACITIDINE
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BIVALIRUDIN, BIVALIRUDIN
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CABAZITAXEL, CABAZITAXEL
 CAPECITABINE, CAPECITABINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBOPLATIN, CARBOPLATIN
 CARMUSTINE, CARMUSTINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOZAPINE, CLOZAPINE
 DALFAMPRIDINE, DALFAMPRIDINE
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DOCETAXEL, DOCETAXEL
 DODEX, CYANOCOBALAMIN
 DOFETILIDE, DOFETILIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 EPLERENONE, EPLERENONE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ETOPOSIDE, ETOPOSIDE
 EZETIMIBE, EZETIMIBE
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUOROURACIL, FLUOROURACIL
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACCORD HEALTHCARE INC
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LISINAPRIL, LISINAPRIL
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MITOMYCIN, MITOMYCIN
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 OXYBUTYNYNIN CHLORIDE, OXYBUTYNYNIN CHLORIDE
 PACLITAXEL, PACLITAXEL
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIRFENIDONE, PIRFENIDONE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PROGESTERONE, PROGESTERONE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 REGADENOSON, REGADENOSON
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TOPIRAMATE, TOPIRAMATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ACCORD HLTHCARE INC

* ACCORD HEALTHCARE INC USA
 BUSULFAN, BUSULFAN
 TIGECYCLINE, TIGECYCLINE

ACELLA

* ACELLA PHARMACEUTICALS LLC
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BENZONATATE, BENZONATATE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CICLOPIROX, CICLOPIROX
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DUTASTERIDE, DUTASTERIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACELLA PHARMACEUTICALS LLC
NIFEDIPINE, NIFEDIPINE
PHENYTOIN SODIUM, PHENYTOIN SODIUM

ACELLA PHARMS LLC

* ACELLA PHARMACEUTICALS LLC
GABAPENTIN, GABAPENTIN

ACER

* ACER THERAPEUTICS INC
OLPRUVA, SODIUM PHENYL BUTYRATE

ACERUS

* ACERUS PHARMACEUTICALS CORP
NATESTO, TESTOSTERONE

ACI

* ACI HEALTHCARE LTD
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
AZITHROMYCIN, AZITHROMYCIN
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
LEVETIRACETAM, LEVETIRACETAM
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

ACIC PHARMS

* ACIC PHARMACEUTICALS INC
BACLOFEN, BACLOFEN
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
TRANEXAMIC ACID, TRANEXAMIC ACID
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

ACORDA

* ACORDA THERAPEUTICS INC
AMPYRA, DALFAMPRIDINE
INBRIJA, LEVODOPA

ACROTECH BIOPHARMA

* ACROTECH BIOPHARMA INC
BELEODAQ, BELINOSTAT
EVOMELA, MELPHALAN HYDROCHLORIDE
FOLOTYN, PRALATREXATE
FUSILEV, LEVOLEUCOVORIN CALCIUM
KHAPZORY, LEVOLEUCOVORIN

ACS DOBFAR

* ACS DOBFAR SPA
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
CEFAZOLIN SODIUM, CEFZAZOLIN SODIUM
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFOXITIN, CEFOXITIN SODIUM
CEFTAZIDIME, CEFTAZIDIME
CEFTRIAXONE, CEFTRIAXONE SODIUM
IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
MEROPENEM, MEROPENEM

ACS DOBFAR SPA

* ACS DOBFAR SPA
AMPICILLIN SODIUM, AMPICILLIN SODIUM
CEFUROXIME SODIUM, CEFUROXIME SODIUM
ERTAPENEM SODIUM, ERTAPENEM SODIUM
MEROPENEM, MEROPENEM
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

ACTAVIS

* ACTAVIS LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
DOCETAXEL, DOCETAXEL
HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
OXALIPLATIN, OXALIPLATIN
PEMETREXED, PEMETREXED

ACTAVIS ELIZABETH

* ACTAVIS ELIZABETH LLC
ALBENDAZOLE, ALBENDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS ELIZABETH LLC
 ALPRAZOLAM, ALPRAZOLAM
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 GABAPENTIN, GABAPENTIN
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 INDAPAMIDE, INDAPAMIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMOTRIGINE, LAMOTRIGINE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LOVASTATIN, LOVASTATIN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 RANOLAZINE, RANOLAZINE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
- * ACTAVIS ELIZABETH LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ALPRAZOLAM, ALPRAZOLAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

ACTAVIS INC

- * ACTAVIS INC
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE

ACTAVIS LABS

- * ACTAVIS LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 PERMETHRIN, PERMETHRIN

ACTAVIS LABS FL

- * ACTAVIS LABORATORIES FL INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 ISOTRETINOIN, ISOTRETINOIN
 MESALAMINE, MESALAMINE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE

ACTAVIS LABS FL INC

- * ACTAVIS LABORATORIES FL INC
 BUDESONIDE, BUDESONIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACTAVIS LABORATORIES FL INC
 CARTIA XT, DILTIAZEM HYDROCHLORIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DALFAMPRIDINE, DALFAMPRIDINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 GEFITINIB, GEFITINIB
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 LEVETIRACETAM, LEVETIRACETAM
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OMEPRAZOLE, OMEPRAZOLE
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISONE, PREDNISONE
 RAMELTEON, RAMELTEON
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TETRABENAZINE, TETRABENAZINE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

ACTAVIS LABS UT INC

* ACTAVIS LABORATORIES UT INC
 AZELAIC ACID, AZELAIC ACID
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLONIDINE, CLONIDINE
 FIORICET W/ CODEINE, ACETAMINOPHEN
 LIDOCAINE, LIDOCAINE
 SCOPOLAMINE, SCOPOLAMINE
 TESTOSTERONE, TESTOSTERONE

* ACTAVIS LABORATORIES UT INC INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 PIMECROLIMUS, PIMECROLIMUS
 TESTOSTERONE, TESTOSTERONE

ACTAVIS LLC

* ACTAVIS LLC
 AZACITIDINE, AZACITIDINE
 DAPSONE, DAPSONE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE

ACTAVIS MID ATLANTIC

* ACTAVIS MID ATLANTIC LLC
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE, ADAPALENE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS MID ATLANTIC LLC
HYDROCORTISONE, HYDROCORTISONE
LEVETIRACETAM, LEVETIRACETAM
MESALAMINE, MESALAMINE
NITROFURANTOIN, NITROFURANTOIN
NYSTATIN, NYSTATIN
VALNAC, BETAMETHASONE VALERATE
- * ACTAVIS MID ATLANTIC LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
IBUPROFEN, IBUPROFEN
PERMETHRIN, PERMETHRIN (OTC)

ACTAVIS PHARMA

- * ACTAVIS PHARMA INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
MICONAZOLE NITRATE, MICONAZOLE NITRATE

ACTAVIS TOTOWA

- * ACTAVIS TOTOWA LLC
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
PACLITAXEL, PACLITAXEL
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
VINORELBINE TARTRATE, VINORELBINE TARTRATE

ACTELION

- * ACTELION PHARMACEUTICALS US INC
OPSUMIT, MACITENTAN
TRACLEER, BOSENTAN
UPTRAVI, SELEXIPAG
VELETRI, EPOPROSTENOL SODIUM
VENTAVIS, ILOPROST
ZAVESCA, MIGLUSTAT

ADAMAS OPERATIONS

- * ADAMAS OPERATIONS LLC
GOCOVRI, AMANTADINE HYDROCHLORIDE
OSMOLEX ER, AMANTADINE HYDROCHLORIDE

ADAMIS PHARMS CORP

- * ADAMIS PHARMACEUTICALS CORP
SYMJEPI, EPINEPHRINE
ZIMHI, NALOXONE HYDROCHLORIDE

ADAPTIS

- * ADAPTIS PHARMA PRIVATE LTD
BICALUTAMIDE, BICALUTAMIDE
DUTASTERIDE, DUTASTERIDE
PREGABALIN, PREGABALIN
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

ADARE PHARMS INC

- * ADARE PHARMACEUTICALS INC
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

ADHERA

- * ADHERA THERAPEUTICS INC
PRESTALIA, AMLODIPINE BESYLATE

ADIENNE SA

- * ADIENNE SA
TEPADINA, THIOTEPA

AET PHARMA

- * AET PHARMA US INC
LEFLUNOMIDE, LEFLUNOMIDE
POSACONAZOLE, POSACONAZOLE

AFAXYS

- * AFAXYS PHARMA LLC
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

AGEPHA PHARMA FZ

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AGEPHA PHARMA FZ LLC
LODOCO, COLCHICINE

AGILE

* AGILE THERAPEUTICS INC
TWIRLA, ETHINYL ESTRADIOL

AGIOS PHARMS INC

* AGIOS PHARMACEUTICALS INC
PYRUKYND, MITAPIVAT SULFATE

AGNITIO

* AGNITIO INC
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
ETHACRYNIC ACID, ETHACRYNIC ACID
PHYTONADIONE, PHYTONADIONE
TRIAMTERENE, TRIAMTERENE

AGOURON PHARMS

* AGOURON PHARMACEUTICALS LLC
VIRACEPT, NELFINAVIR MESYLATE

AILEX PHARMS LLC

* AILEX PHARMACEUTICALS LLC
CROMOLYN SODIUM, CROMOLYN SODIUM
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

AIPING PHARM INC

* AIPING PHARMACEUTICAL INC
LEVETIRACETAM, LEVETIRACETAM
SORINE, SOTALOL HYDROCHLORIDE

AIRGAS THERAP

* AIRGAS THERAPEUTICS LLC
ULSPIRA, NITRIC OXIDE

AIRIS PHARMA PVT LTD

* AIRIS PHARMA PRIVATE LTD
MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM

AIZANT

* AIZANT DRUG RESEARCH SOLUTIONS PRIVATE LTD
ALBUTEROL SULFATE, ALBUTEROL SULFATE
PIRFENIDONE, PIRFENIDONE
ZILEUTON, ZILEUTON
* AIZANT DRUG RESEARCH SOLUTIONS PVT LTD
ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE

AJANTA PHARMA LTD

* AJANTA PHARMA LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ARIPIRAZOLE, ARIPIRAZOLE
CAPTOPRIL, CAPTOPRIL
CHLORTHALIDONE, CHLORTHALIDONE
CHOLESTYRAMINE, CHOLESTYRAMINE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DROXIDOPA, DROXIDOPA
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
ENTACAPONE, ENTACAPONE
FAMOTIDINE, FAMOTIDINE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AJANTA PHARMA LTD**

MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 RANOLAZINE, RANOLAZINE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TOPIRAMATE, TOPIRAMATE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLMITRIPTAN, ZOLMITRIPTAN

AJENAT PHARMS*** AJENAT PHARMACEUTICALS LLC**

DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
 TRANXENE, CLORAZEPATE DIPOTASSIUM

AKARX INC*** AKARX INC**

DOPTELET, AVATROMBOPAG MALEATE

AKCEA THERAPS*** AKCEA THERAPEUTICS INC**

TEGSEDI, INOTERSEN SODIUM

AKORN*** AKORN OPERATING CO LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FAMOTIDINE, FAMOTIDINE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

ALBIREO*** ALBIREO AB**

BYLVAY, ODEVIXIBAT

ALCON*** ALCON LABORATORIES INC**

BSS PLUS, CALCIUM CHLORIDE
 BSS, CALCIUM CHLORIDE
 MIOSTAT, CARBACHOL
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

*** ALCON RESEARCH LLC**

PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

ALCON LABS*** ALCON LABORATORIES LTD**

TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE

ALCON LABS INC*** ALCON LABORATORIES INC**

ALCAINE, PROPARACAINE HYDROCHLORIDE
 CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 EYSUVIS, LOTEHPREDNOL ETABONATE
 FLUORESCITE, FLUORESCIN SODIUM
 INVELTYS, LOTEHPREDNOL ETABONATE
 ISOPTO ATROPINE, ATROPINE SULFATE
 MYDRIACYL, TROPICAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALCON LABORATORIES INC**

PATADAY ONCE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 PATADAY TWICE DAILY RELIEF, OLOPATADINE HYDROCHLORIDE (OTC)
 RHOPRESSA, NETARSUDIL MESYLATE
 ROCKLATAN, LATANOPROST
 SIMBRINZA, BRIMONIDINE TARTRATE

ALCON PHARMS LTD*** ALCON PHARMACEUTICALS LTD**

BETADINE, POVIDONE-IODINE
 ZADITOR, KETOTIFEN FUMARATE (OTC)

ALEMBIC*** ALEMBIC PHARMACEUTICALS LTD**

ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ADAPALENE, ADAPALENE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
 BIMATOPROST, BIMATOPROST
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CARMUSTINE, CARMUSTINE
 CELECOXIB, CELECOXIB
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DAPSONE, DAPSONE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DEFERASIROX, DEFERASIROX
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESONIDE, DESONIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DOCETAXEL, DOCETAXEL
 DOCOSANOL, DOCOSANOL (OTC)
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 FEBUXOSTAT, FEBUXOSTAT
 FENOFIBRATE (MICRONIZED), FENOFIBRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALEMBIC PHARMACEUTICALS LTD
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 FULVESTRANT, FULVESTRANT
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ITRACONAZOLE, ITRACONAZOLE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METRONIDAZOLE, METRONIDAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MODAFINIL, MODAFINIL
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MUPIROCIN, MUPIROCIN CALCIUM
 NADOLOL, NADOLOL
 NIFEDIPINE, NIFEDIPINE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PACLITAXEL, PACLITAXEL
 PIRFENIDONE, PIRFENIDONE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SELEXIPAG, SELEXIPAG
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TAVABOROLE, TAVABOROLE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TERIFLUNOMIDE, TERIFLUNOMIDE
 THEOPHYLLINE, THEOPHYLLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRAVOPROST, TRAVOPROST
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ALEMBIC GLOBAL

* ALEMBIC GLOBAL HOLDING SA
 TREPROSTINIL, TREPROSTINIL

ALEMBIC LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALEMBIC LTD**

LITHIUM CARBONATE, LITHIUM CARBONATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

ALEMBIC PHARMS LTD*** ALEMBIC PHARMACEUTICALS LTD**

CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DESVENLAFAXINE, DESVENLAFAXINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FAMOTIDINE, FAMOTIDINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEPROBAMATE, MEPROBAMATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE, METRONIDAZOLE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

ALEXZA PHARMS*** ALEXZA PHARMACEUTICALS INC**

ADASUVE, LOXAPINE

ALIGNSCIENCE PHARMA*** ALIGNSCIENCE PHARMA INC**

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

ALIMERA SCIENCES INC*** ALIMERA SCIENCES INC**

ILUVIEN, FLUOCINOLONE ACETONIDE
 YUTIQ, FLUOCINOLONE ACETONIDE

ALKALOIDA ZRT*** ALKALOIDA CHEMICAL CO ZRT**

HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE

ALKEM*** ALKEM LABORATORIES LTD**

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 GABAPENTIN, GABAPENTIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ALKEM LABS LTD*** ALKEM LABORATORIES LTD**

AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 APREMILAST, APREMILAST
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BREXPIPRAZOLE, BREXPIPRAZOLE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CAPECITABINE, CAPECITABINE
 CEFDINIR, CEFDINIR
 CEFIXIME, CEFIXIME
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALKEM LABORATORIES LTD**

COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DALFAMPRIDINE, DALFAMPRIDINE
 DEFERASIROX, DEFERASIROX
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ERYTHROMYCIN, ERYTHROMYCIN
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 EVEROLIMUS, EVEROLIMUS
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FINASTERIDE, FINASTERIDE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 GABAPENTIN, GABAPENTIN
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 IBUPROFEN, IBUPROFEN
 ITRACONAZOLE, ITRACONAZOLE
 LACOSAMIDE, LACOSAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MARINOL, DRONABINOL
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NIFEDIPINE, NIFEDIPINE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANOLAZINE, RANOLAZINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 RUFINAMIDE, RUFINAMIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SIROLIMUS, SIROLIMUS
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TACROLIMUS, TACROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALKEM LABORATORIES LTD
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
 TICAGRELOR, TICAGRELOR
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TOLVAPTAN, TOLVAPTAN
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN, VALSARTAN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN

ALKERMES

* ALKERMES INC
 VIVITROL, NALTREXONE

ALKERMES INC

* ALKERMES INC
 ARISTADA INITIO KIT, ARIPIPIRAZOLE LAUROXIL
 ARISTADA, ARIPIPIRAZOLE LAUROXIL
 LYBALVI, OLANZAPINE

ALLEGIANCE HLTHCARE

* ALLEGIANCE HEALTHCARE CORP
 POVIDONE IODINE, POVIDONE-IODINE (OTC)

ALLERGAN

* ALLERGAN INC
 AVAGE, TAZAROTENE
 OCUFLOX, OFLOXACIN
 TAZORAC, TAZAROTENE

* ALLERGAN PHARMACEUTICAL
 BETAGAN, LEVOBUNOLOL HYDROCHLORIDE

* ALLERGAN SALES LLC
 AVYCAZ, AVIBACTAM SODIUM
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE
 CONDYLOX, PODOFILOX
 CRINONE, PROGESTERONE
 ESTRACE, ESTRADIOL
 INFED, FERRIC OXYHYDROXIDE
 OXYTROL, OXYBUTYNIN
 RAPAFLO, SILODOSIN
 SAPHRIS, ASENAPINE MALEATE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL

ALMAJECT

* ALMAJECT INC
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

ALMATICA

* ALMATICA PHARMA INC
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 GRALISE, GABAPENTIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* ALMATICA PHARMA LLC
 LOREEV XR, LORAZEPAM
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 VENLAFAXINE BESYLATE, VENLAFAXINE BESYLATE
 ZESTORETIC, HYDROCHLOROTHIAZIDE

ALMIRALL

* ALMIRALL LLC
 ACTICLATE, DOXYCYCLINE HYCLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALMIRALL LLC**

ACZONE, DAPSONE
 ALTABAX, RETAPAMULIN
 AZELEX, AZELAIC ACID
 CORDRAN, FLURANDRENOLIDE
 KLISYRI, TIRBANIBULIN
 SEYSARA, SARECYCLINE HYDROCHLORIDE
 VELTIN, CLINDAMYCIN PHOSPHATE

ALNYLAM PHARMS INC

* ALNYLAM PHARMACEUTICALS INC
 AMVUTTRA, VUTRISIRAN SODIUM
 GIVLAARI, GIVOSIRAN SODIUM
 ONPATTRO, PATISIRAN SODIUM
 OXLUMO, LUMASIRAN SODIUM

ALTAIRE PHARMS INC

* ALTAIRE PHARMACEUTICALS INC
 ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OFLOXACIN, OFLOXACIN

ALTATHERA PHARMS LLC

* ALTATHERA PHARMACEUTICALS LLC
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

ALTHERA PHARMS

* ALTHERA PHARMACEUTICALS LLC
 LYPQOZET, ATORVASTATIN CALCIUM

ALVOGEN

* ALVOGEN INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 BONSIITY, TERIPARATIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE, BUPRENORPHINE
 CARBIDOPA, CARBIDOPA
 DEXAMETHASONE, DEXAMETHASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DISULFIRAM, DISULFIRAM
 ESTRADIOL, ESTRADIOL
 FELBAMATE, FELBAMATE
 FLUOCINONIDE, FLUOCINONIDE
 HYDROCODONE BITARTRATE, HYDROCODONE BITARTRATE
 LIDEX, FLUOCINONIDE
 MELPHALAN, MELPHALAN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 PYRIMETHAMINE, PYRIMETHAMINE
 RIVASTIGMINE, RIVASTIGMINE
 THYRO-TABS, LEVOTHYROXINE SODIUM **
 UREX, METHENAMINE HIPPURATE

AM REGENT

* AMERICAN REGENT INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ATROPINE SULFATE, ATROPINE SULFATE
 BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CUPRIC SULFATE, CUPRIC SULFATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DROPERIDOL, DROPERIDOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AMERICAN REGENT INC**

ESTRADIOL VALERATE, ESTRADIOL VALERATE
 FOMEPIZOLE, FOMEPIZOLE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INJECTAFER, FERRIC CARBOXYMALTOSIDE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 LEVOCARNITINE, LEVOCARNITINE
 METHOCARBAMOL, METHOCARBAMOL
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 MULTRYS, CUPRIC SULFATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NITROGLYCERIN, NITROGLYCERIN
 OLANZAPINE, OLANZAPINE
 PACLITAXEL, PACLITAXEL
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 SELENIOS ACID, SELENIOS ACID
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TRALEMENT, CUPRIC SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VASOPRESSIN, VASOPRESSIN
 VENOFER, FERRIC OXYHYDROXIDE
 ZINC SULFATE, ZINC SULFATE

AMARIN PHARMS

*** AMARIN PHARMACEUTICALS IRELAND LTD**
 VASCEPA, ICOSAPENT ETHYL

AMGEN

*** AMGEN INC**
 SENSIPAR, CINACALCET HYDROCHLORIDE

AMGEN INC

*** AMGEN INC**
 CORLANOR, IVABRADINE
 CORLANOR, IVABRADINE HYDROCHLORIDE
 LUMAKRAS, SOTORASIB
 OTEZLA, APREMILAST

AMICI

*** AMICI PHARMACEUTICALS LLC**
 CATAFLAM, DICLOFENAC POTASSIUM
 DIGOXIN, DIGOXIN

AMICUS THERAP US

*** AMICUS THERAPEUTICS US LLC**
 GALAFOLD, MIGALASTAT HYDROCHLORIDE
 OPFOLDA, MIGLUSTAT

AMIVAS

*** AMIVAS INC**
 ARTESUNATE, ARTESUNATE

AMNEAL

*** AMNEAL EU LTD**
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZACITIDINE, AZACITIDINE
 BUSULFAN, BUSULFAN
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CARMUSTINE, CARMUSTINE
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CLOFARABINE, CLOFARABINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DARUNAVIR, DARUNAVIR
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AMNEAL EU LTD**

DEXAMETHASONE, DEXAMETHASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIFLUPREDNATE, DIFLUPREDNATE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DOCETAXEL, DOCETAXEL
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETRAVIRINE, ETRAVIRINE
 FLUOROMETHOLONE, FLUOROMETHOLONE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FULVESTRANT, FULVESTRANT
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 METYROSINE, METYROSINE
 NELARABINE, NELARABINE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OFLOXACIN, OFLOXACIN
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PIRFENIDONE, PIRFENIDONE
 PLERIXAFOR, PLERIXAFOR
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 PREDNISONE, PREDNISONE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 THEOPHYLLINE, THEOPHYLLINE
 THIOTHIXENE, THIOTHIXENE
 TIOPRONIN, TIOPRONIN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VASOPRESSIN, VASOPRESSIN
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

*** AMNEAL PHARMACEUTICALS LLC**

ACTIVELLA, ESTRADIOL
 ACYCLOVIR, ACYCLOVIR
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 BEXAROTENE, BEXAROTENE
 BUPRENORPHINE, BUPRENORPHINE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DAPSONE, DAPSONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS LLC
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ELURYNG, ETHINYL ESTRADIOL
 ESTRADIOL, ESTRADIOL
 ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL
 FENOFIBRATE, FENOFIBRATE
 FLUOCINONIDE, FLUOCINONIDE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LIDOCAINE, LIDOCAINE
 LIORESAL, BACLOFEN
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LUBIPROSTONE, LUBIPROSTONE
 LYVISPAH, BACLOFEN
 MUPIROCIN, MUPIROCIN CALCIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 POSACONAZOLE, POSACONAZOLE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SIROLIMUS, SIROLIMUS
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCRALFATE, SUCRALFATE
 TAVABOROLE, TAVABOROLE
 TESTOSTERONE, TESTOSTERONE
 TIGECYCLINE, TIGECYCLINE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOMIG, ZOLMITRIPTAN

AMNEAL PHARM

* AMNEAL PHARMACEUTICAL
 ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FOLIC ACID, FOLIC ACID
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE

AMNEAL PHARMS

* AMNEAL PHARMACEUTICALS
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATOVAQUONE, ATOVAQUONE
 CALCITRIOL, CALCITRIOL
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ENTECAVIR, ENTECAVIR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AMNEAL PHARMACEUTICALS
 - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 - ESTRADIOL, ESTRADIOL
 - FELBAMATE, FELBAMATE
 - GABAPENTIN, GABAPENTIN
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - INDOMETHACIN, INDOMETHACIN
 - ITRACONAZOLE, ITRACONAZOLE
 - LEVETIRACETAM, LEVETIRACETAM
 - LIDOCAINE, LIDOCAINE
 - LORAZEPAM, LORAZEPAM
 - MEROPENEM, MEROPENEM
 - METAXALONE, METAXALONE
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - MONTELUKAST SODIUM, MONTELUKAST SODIUM
 - NIACIN, NIACIN
 - NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 - NITROFURANTOIN, NITROFURANTOIN
 - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - OXCARBAZEPINE, OXCARBAZEPINE
 - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 - PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE
 - PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 - QUININE SULFATE, QUININE SULFATE
 - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 - RISPERIDONE, RISPERIDONE
 - SILDENAFIL CITRATE, SILDENAFIL CITRATE
 - TELMISARTAN, TELMISARTAN
 - TEMAZEPAM, TEMAZEPAM
 - TEMOZOLOMIDE, TEMOZOLOMIDE
 - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 - VORICONAZOLE, VORICONAZOLE
 - WARFARIN SODIUM, WARFARIN SODIUM
- * AMNEAL PHARMACEUTICALS HOLDINGS GMBH
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
- * AMNEAL PHARMACEUTICALS OF NEW YORK LLC
 - ABIRATERONE ACETATE, ABIRATERONE ACETATE
 - ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 - AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 - ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 - BUDESONIDE, BUDESONIDE
 - BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 - CARBAMAZEPINE, CARBAMAZEPINE
 - CELECOXIB, CELECOXIB
 - DUTASTERIDE, DUTASTERIDE
 - ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 - GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 - GUAIFENESIN, GUAIFENESIN (OTC)
 - IBUPROFEN, IBUPROFEN (OTC)
 - IRBESARTAN, IRBESARTAN
 - LACOSAMIDE, LACOSAMIDE
 - LAMOTRIGINE, LAMOTRIGINE
 - LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 - METHOTREXATE SODIUM, METHOTREXATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AMNEAL PHARMACEUTICALS OF NEW YORK LLC**

METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PALIPERIDONE, PALIPERIDONE
 PARICALCITOL, PARICALCITOL
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 RIVASTIGMINE, RIVASTIGMINE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCRALFATE, SUCRALFATE
 TOBRAMYCIN, TOBRAMYCIN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN

AMNEAL PHARMS CO*** AMNEAL PHARMACEUTICALS CO GMBH**

ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARGATROBAN, ARGATROBAN
 BUMETANIDE, BUMETANIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CLOBAZAM, CLOBAZAM
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FUROSEMIDE, FUROSEMIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 NADOLOL, NADOLOL
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 PARICALCITOL, PARICALCITOL
 PHYTONADIONE, PHYTONADIONE
 PREGABALIN, PREGABALIN
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 URSODIOL, URSODIOL

AMNEAL PHARMS NY*** AMNEAL PHARMACEUTICALS NY LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ALPRAZOLAM, ALPRAZOLAM
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 GABAPENTIN, GABAPENTIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS NY LLC
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

* AMNEAL PHARMACEUTICALS OF NY LLC
 BEXAROTENE, BEXAROTENE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISOTRETINOIN, ISOTRETINOIN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PROGESTERONE, PROGESTERONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

AMNEALS PHARMS

* AMNEALS PHARMACEUTICALS LLC
 FAMOTIDINE, FAMOTIDINE

AMPHASTAR PHARM

* AMPHASTAR PHARMACEUTICAL INC
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM

AMPHASTAR PHARMS INC

* AMPHASTAR PHARMACEUTICALS INC
 BAQSIMI, GLUCAGON
 CORTROSYN, COSYNTROPIN
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GLUCAGON, GLUCAGON
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 VASOPRESSIN, VASOPRESSIN

AMRING PHARMS

* AMRING PHARMACEUTICALS INC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 LATANOPROST, LATANOPROST
 LYSTEDA, TRANEXAMIC ACID
 MESALAMINE, MESALAMINE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE

AMRYT

* AMRYT PHARMACEUTICALS DAC
 FILSUVEZ, BIRCH TRITERPENES
 JUXTAPID, LOMITAPIDE MESYLATE

AMTA

* AMTA LABS LTD
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE

AMVLYX

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMYLYX PHARMACEUTICALS INC
RELYVRIO, SODIUM PHENYL BUTYRATE

ANACOR PHARMS INC

* ANACOR PHARMACEUTICALS INC
EUCRISA, CRISABOROLE
KERYDIN, TAVABOROLE

ANBEX

* ANBEX INC
IOSAT, POTASSIUM IODIDE (OTC)

ANBISON LAB

* ANBISON LABORATORY CO LTD
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CARBAMAZEPINE, CARBAMAZEPINE
MONTELUKAST SODIUM, MONTELUKAST SODIUM

ANCHEN PHARMS

* ANCHEN PHARMACEUTICALS INC
ALISKIREN HEMIFUMARATE, ALISKIREN HEMIFUMARATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
TRETINOIN, TRETINOIN

* ANCHEN PHARMACEUTICALS, INC
ALPRAZOLAM, ALPRAZOLAM
CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN

ANDA REPOSITORY

* ANDA REPOSITORY LLC
AMCINONIDE, AMCINONIDE
CEFDINIR, CEFDINIR
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM, DIATRIZOATE MEGLUMINE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LEVETIRACETAM, LEVETIRACETAM
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRIMIDONE, PRIMIDONE

ANDOR PHARMS

* ANDOR PHARMACEUTICALS LLC
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ANDRX LABS LLC

* ANDRX LABS LLC
FORTAMET, METFORMIN HYDROCHLORIDE

ANI PHARMS

* ANI PHARMACEUTICALS INC
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
ARIMIDEX, ANASTROZOLE
ATACAND HCT, CANDESARTAN CILEXETIL
ATACAND, CANDESARTAN CILEXETIL
BACLOFEN, BACLOFEN
BENZAEPRIIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENZAEPRIIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BEXAROTENE, BEXAROTENE
BRETHINE, TERBUTALINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ANI PHARMACEUTICALS INC
 CASODEX, BICALUTAMIDE
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 CORTENEMA, HYDROCORTISONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ETODOLAC, ETODOLAC
 FELBAMATE, FELBAMATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCONAZOLE, FLUCONAZOLE
 GLIPIZIDE, GLIPIZIDE
 INDAPAMIDE, INDAPAMIDE
 INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 LITHOBID, LITHIUM CARBONATE
 LUVOX, FLUVOXAMINE MALEATE
 MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIGLUSTAT, MIGLUSTAT
 MISOPROSTOL, MISOPROSTOL
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NILUTAMIDE, NILUTAMIDE
 OXISTAT, OXICONAZOLE NITRATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PANDEL, HYDROCORTISONE PROBUTATE
 PENICILLAMINE, PENICILLAMINE
 PINDOLOL, PINDOLOL
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PURIFIED CORTROPHIN GEL, CORTICOTROPIN
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VALPROIC ACID, VALPROIC ACID
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VEREGEN, SINECATECHINS

ANIMA

* ANIMA PHARMACEUTICALS PVT LTD
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DEXAMETHASONE, DEXAMETHASONE
 TRIACIN-C, CODEINE PHOSPHATE

ANNORA

* ANNORA PHARMA PRIVATE LTD
 APREMILAST, APREMILAST
 DROXIDOPA, DROXIDOPA
 LAMIVUDINE, LAMIVUDINE

ANNORA PHARMA

* ANNORA PHARMA PRIVATE LTD
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DIATRIZOATE MEGLUMINE AND DIATRIZOATE SODIUM, DIATRIZOATE MEGLUMINE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 EPLERENONE, EPLERENONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ANNORA PHARMA PRIVATE LTD**

FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 GLYCOPYRROLATE, GLYCOPYRROLATE
 ITRACONAZOLE, ITRACONAZOLE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 OXCARBAZEPINE, OXCARBAZEPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 THEOPHYLLINE, THEOPHYLLINE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZAFIRLUKAST, ZAFIRLUKAST
 ZILEUTON, ZILEUTON

ANTARES PHARMA INC*** ANTARES PHARMA INC**

TLANDO, TESTOSTERONE UNDECANOATE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE

ANTIBIOTICE*** ANTIBIOTICE SA**

AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

ANXIN*** ANXIN PHARMA INC**

SEVELAMER CARBONATE, SEVELAMER CARBONATE

APELLIS PHARMS*** APELLIS PHARMACEUTICALS INC**

EMPAVELI, PEGCETACOPLAN
 SYFOVRE, PEGCETACOPLAN

APGDI*** ASTELLAS PHARMA GLOBAL DEVELOPMENT INC**

MYRBETRIQ GRANULES, MIRABEGRON
 MYRBETRIQ, MIRABEGRON

APIL*** ALLERGAN PHARMACEUTICALS INTERNATIONAL LTD**

ACTONEL, RISEDRONATE SODIUM
 ATELVIA, RISEDRONATE SODIUM
 LO LOESTRIN FE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 TAYTULLA, ETHINYL ESTRADIOL

APNAR PHARMA LP*** APNAR PHARMA LP**

BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

APOTEX*** APOTEX INC**

ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACYCLOVIR, ACYCLOVIR
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 ARIPIPRAZOLE, ARIPIPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTEX INC
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATOVAQUONE, ATOVAQUONE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
 BICALUTAMIDE, BICALUTAMIDE
 BIMATOPROST, BIMATOPROST
 BORTEZOMIB, BORTEZOMIB
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BUSULFAN, BUSULFAN
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CELECOXIB, CELECOXIB
 CIMETIDINE, CIMETIDINE (OTC)
 CYCLOSPORINE, CYCLOSPORINE
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DEXAMETHASONE, DEXAMETHASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ETODOLAC, ETODOLAC
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 GEFITINIB, GEFITINIB
 GEMFIBROZIL, GEMFIBROZIL
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LACOSAMIDE, LACOSAMIDE
 LAMIVUDINE, LAMIVUDINE
 LENALIDOMIDE, LENALIDOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 MACITENTAN, MACITENTAN
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MICAfungin SODIUM, MICAfungin SODIUM
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MODAFINIL, MODAFINIL
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NILOTINIB HYDROCHLORIDE, NILOTINIB HYDROCHLORIDE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTEX INC
 OMEPRAZOLE, OMEPRAZOLE
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PAXIL CR, PAROXETINE HYDROCHLORIDE
 PAXIL, PAROXETINE HYDROCHLORIDE
 PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PENICILLAMINE, PENICILLAMINE
 PENTOXIFYLLINE, PENTOXIFYLLINE
 PIRFENIDONE, PIRFENIDONE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREGABALIN, PREGABALIN
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 REGADENOSON, REGADENOSON
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SIROLIMUS, SIROLIMUS
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TASIMELTEON, TASIMELTEON
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TERIPARATIDE, TERIPARATIDE
 TETRABENAZINE, TETRABENAZINE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLVAPTAN, TOLVAPTAN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAVOPROST, TRAVOPROST
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

APOTEX INC

* APOTEX INC
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFPROZIL, CEFPROZIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMIQUIMOD, IMIQUIMOD
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTEX INC
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* APOTEX INC ETOBICOKE SITE
 ACYCLOVIR, ACYCLOVIR
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CILOSTAZOL, CILOSTAZOL
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 ETODOLAC, ETODOLAC
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 LEFLUNOMIDE, LEFLUNOMIDE
 LORATADINE, LORATADINE (OTC)
 MIRTAZAPINE, MIRTAZAPINE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

* APOTEX INC RICHMOND HILL
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BUDESONIDE, BUDESONIDE (OTC)
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

APOTHECON

* APOTHECON INC DIV BRISTOL MYERS SQUIBB
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENALOG-80, TRIAMCINOLONE ACETONIDE

APP PHARMS

* APP PHARMACEUTICALS LLC
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE

APPCO

* APPCO PHARMA LLC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 METAXALONE, METAXALONE
 MODAFINIL, MODAFINIL
 PERPHENAZINE, PERPHENAZINE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

APRECIA PHARMS

* APRECIA PHARMACEUTICALS LLC
 SPRITAM, LEVETIRACETAM

APTAPHARMA INC

* APTAPHARMA INC
 CHLORZOXAZONE, CHLORZOXAZONE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN (OTC)
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AQUESTIVE**

* AQUESTIVE THERAPEUTICS INC
EXSERVAN, RILUZOLE

ARBOR PHARMS LLC

* ARBOR PHARMACEUTICALS LLC
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
ERY-TAB, ERYTHROMYCIN
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN, ERYTHROMYCIN
EVEKEO, AMPHETAMINE SULFATE
SKLICE, IVERMECTIN (OTC)

ARCUTIS

* ARCUTIS BIOTHERAPEUTICS INC
ZORYVE, ROFLUMILAST

ARDELYX INC

* ARDELYX INC
IBSRELA, TENAPANOR HYDROCHLORIDE
XPHOZAH, TENAPANOR HYDROCHLORIDE

AREVA PHARMS

* AREVA PHARMACEUTICALS INC
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FUROSEMIDE, FUROSEMIDE
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

ARMSTRONG PHARMS

* ARMSTRONG PHARMACEUTICALS INC
PRIMATENE MIST, EPINEPHRINE (OTC)

ARNA PHARMA

* ARNA PHARMA PTY LTD
ARIDOL KIT, MANNITOL

ARRAY BIOPHARMA INC

* ARRAY BIOPHARMA INC
BRAFTOVI, ENCORAFENIB
MEKTOVI, BINIMETINIB

ARROW INTL

* ARROW INTERNATIONAL LTD
LENALIDOMIDE, LENALIDOMIDE

ARTHUR GRP

* ARTHUR GROUP LLC
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
SEVELAMER CARBONATE, SEVELAMER CARBONATE

ASCEND THERAPS US

* ASCEND THERAPEUTICS US LLC
ESTROGEL, ESTRADIOL

ASCENT PHARMS INC

* ASCENT PHARMACEUTICALS INC
BENZONATATE, BENZONATATE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DRONABINOL, DRONABINOL
DUTASTERIDE, DUTASTERIDE
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)
GABAPENTIN, GABAPENTIN
GEMFIBROZIL, GEMFIBROZIL
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
IBUPROFEN, IBUPROFEN (OTC)
ICOSAPENT ETHYL, ICOSAPENT ETHYL
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ASCENT PHARMACEUTICALS INC**

LISINOPRIL, LISINOPRIL
 MELOXICAM, MELOXICAM
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PALIPERIDONE, PALIPERIDONE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

ASPEN*** ASPEN PHARMA USA INC**

ACETYLCYSTEINE, ACETYLCYSTEINE
 GLYCOPYRROLATE, GLYCOPYRROLATE

ASPEN GLOBAL INC*** ASPEN GLOBAL INC**

BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CYCLESSA, DESOGESTREL

ASPIRO*** ASPIRO PHARMA LTD**

ACETAMINOPHEN, ACETAMINOPHEN
 DAPTOMYCIN, DAPTOMYCIN
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUGAMMADEX SODIUM, SUGAMMADEX SODIUM
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ASSERTIO*** ASSERTIO THERAPEUTICS INC**

CAMBIA, DICLOFENAC POTASSIUM
 ZIPSOR, DICLOFENAC POTASSIUM

ASTELLAS*** ASTELLAS PHARMA US INC**

AMBISOME, AMPHOTERICIN B
 ASTAGRAF XL, TACROLIMUS
 CRESEMBA, ISAVUCONAZONIUM SULFATE
 LEXISCAN, REGADENOSON
 MYCAMINE, MICAFUNGIN SODIUM
 PROGRAF, TACROLIMUS
 VEOZAH, FEZOLINETANT
 VESICARE LS, SOLIFENACIN SUCCINATE
 VESICARE, SOLIFENACIN SUCCINATE
 XOSPATA, GILTERITINIB FUMARATE
 XTANDI, ENZALUTAMIDE

ASTRAL*** ASTRAL STERITECH PVT LTD**

AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

ASTRAZENECA*** ASTRAZENECA LP**

PULMICORT FLEXHALER, BUDESONIDE
 SYMBICORT, BUDESONIDE

*** ASTRAZENECA PHARMACEUTICALS LP**

AIRSUPRA, ALBUTEROL SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ASTRAZENECA PHARMACEUTICALS LP
 - BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
 - BRILINTA, TICAGRELOR
 - DALIRESP, ROFLUMILAST
 - FASLODEX, FULVESTRANT
 - IRESSA, GEFITINIB
 - KOSELUGO, SELUMETINIB SULFATE
 - LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
 - LYNPARZA, OLAPARIB
 - NEXIUM IV, ESOMEPRAZOLE SODIUM
 - NEXIUM, ESOMEPRAZOLE MAGNESIUM
 - PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
 - PULMICORT RESPULES, BUDESONIDE
 - SEROQUEL, QUETIAPINE FUMARATE
 - SYMBICORT AEROSPHERE, BUDESONIDE
 - TAGRISSE, OSIMERTINIB MESYLATE
 - TRUQAP, CAPIVASERTIB
- * ASTRAZENECA UK LTD
 - CALQUENCE, ACALABRUTINIB
 - CALQUENCE, ACALABRUTINIB MALEATE
 - SEROQUEL XR, QUETIAPINE FUMARATE

ASTRAZENECA AB

- * ASTRAZENECA AB
 - BREZTRI AEROSPHERE, BUDESONIDE
 - BYDUREON BCISE, EXENATIDE SYNTHETIC
 - BYETTA, EXENATIDE SYNTHETIC
 - FARXIGA, DAPAGLIFLOZIN
 - QTERN, DAPAGLIFLOZIN
 - SYMLIN, PRAMLINTIDE ACETATE
 - XIGDUO XR, DAPAGLIFLOZIN

ASTRAZENECA LP

- * ASTRAZENECA LP
 - NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)

ATHEM

- * ATHEM HOLDINGS LLC
 - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 - FOLIC ACID, FOLIC ACID

ATHENA

- * ATHENA BIOSCIENCE LLC
 - NEXICLON XR, CLONIDINE
 - QDOLO, TRAMADOL HYDROCHLORIDE

ATLAS PHARMS LLC

- * ATLAS PHARMACEUTICALS LLC
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ATNAHS PHARMA US

- * ATNAHS PHARMA US LTD
 - ANAPROX DS, NAPROXEN SODIUM
 - EC-NAPROSYN, NAPROXEN
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - NAPROSYN, NAPROXEN

ATON

- * ATON PHARMA INC
 - LODOSYN, CARBIDOPA

AUCTA

- * AUCTA PHARMACEUTICALS INC
 - DEFERASIROX, DEFERASIROX
 - FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 - MOTPOLY XR, LACOSAMIDE
 - OXCARBAZEPINE, OXCARBAZEPINE
 - VIGADRONE, VIGABATRIN

AURINIA

- * AURINIA PHARMACEUTICALS INC
 - LUPKYNIS, VOCLOSPORIN

AUROBINDO

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AUROBINDO PHARMA LTD**

AMOXICILLIN, AMOXICILLIN
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 NEVIRAPINE, NEVIRAPINE
 ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA*** AUROBINDO PHARMA LTD**

ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACYCLOVIR, ACYCLOVIR
 AFIRMELLE, ETHINYL ESTRADIOL
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALPRAZOLAM, ALPRAZOLAM
 AMBRISENTAN, AMBRISENTAN
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATENOLOL, ATENOLOL
 ATHENTIA NEXT, LEVONORGESTREL (OTC)
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 AUROVELA 1.5/30, ETHINYL ESTRADIOL
 AUROVELA 1/20, ETHINYL ESTRADIOL
 AUROVELA 24 FE, ETHINYL ESTRADIOL
 AUROVELA FE 1.5/30, ETHINYL ESTRADIOL
 AUROVELA FE 1/20, ETHINYL ESTRADIOL
 AYUNA, ETHINYL ESTRADIOL
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CARBIDOPA, CARBIDOPA
 CARISOPRODOL, CARISOPRODOL
 CARVEDILOL, CARVEDILOL
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CELECOXIB, CELECOXIB
 CEPHALEXIN, CEPHALEXIN
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 CLOZAPINE, CLOZAPINE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYONANZ, ETHINYL ESTRADIOL
 DALFAMPRIDINE, DALFAMPRIDINE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DIMETHYL FUMARATE, DIMETHYL FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMZAHH, NORETHINDRONE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENTECAVIR, ENTECAVIR
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESZOPICLONE, ESZOPICLONE
 EZETIMIBE, EZETIMIBE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FENOFIBRATE, FENOFIBRATE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 INCASSIA, NORETHINDRONE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 KALLIGA, DESOGESTREL
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LO SIMPESE, ETHINYL ESTRADIOL
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MELOXICAM, MELOXICAM
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 NADOLOL, NADOLOL
 NAPROXEN, NAPROXEN
 NEXESTA FE, ETHINYL ESTRADIOL
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NIFEDIPINE, NIFEDIPINE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN
 NITROGLYCERIN, NITROGLYCERIN
 NYLIA 1/35, ETHINYL ESTRADIOL
 NYLIA 7/7/7, ETHINYL ESTRADIOL
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 OSHIH, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREGABALIN, PREGABALIN
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 RIBAVIRIN, RIBAVIRIN
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RUFINAMIDE, RUFINAMIDE
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIMLIYA, DESOGESTREL
 SIMPESSA, ETHINYL ESTRADIOL
 SIMVASTATIN, SIMVASTATIN
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TOPIRAMATE, TOPIRAMATE
 TORSEMIDE, TORSEMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRANDOLAPRIL, TRANDOLAPRIL
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRI-LO-MILI, ETHINYL ESTRADIOL
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZAFIRLUKAST, ZAFIRLUKAST
 ZALEPLON, ZALEPLON
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

AUROBINDO PHARMA LTD

* AUROBINDO PHARMA LIMITED
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

* AUROBINDO PHARMA LTD
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ARMODAFINIL, ARMODAFINIL
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFIXIME, CEFIXIME
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORZOXAZONE, CHLORZOXAZONE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 DARUNAVIR, DARUNAVIR
 DEFERASIROX, DEFERASIROX
 DIAZEPAM, DIAZEPAM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DIGOXIN, DIGOXIN
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOFETILDE, DOFETILIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 EFAVIRENZ, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EMTRICITABINE, EMTRICITABINE
 ENTACAPONE, ENTACAPONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FELODIPINE, FELODIPINE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
 FINASTERIDE, FINASTERIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCYTOSINE, FLUCYTOSINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GEMFIBROZIL, GEMFIBROZIL
 GLIMEPIRIDE, GLIMEPIRIDE
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 HYDROCORTISONE, HYDROCORTISONE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 ICLEVIA, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LACOSAMIDE, LACOSAMIDE
 LACTULOSE, LACTULOSE
 LANSOPRAZOLE, LANSOPRAZOLE
 LO-ZUMANDIMINE, DROSPIRENONE
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHOCARBAMOL, METHOCARBAMOL
 METRONIDAZOLE, METRONIDAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MILI, ETHINYL ESTRADIOL
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 MODAFINIL, MODAFINIL
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NIACIN, NIACIN
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 OLANZAPINE, OLANZAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 PREDNISON, PREDNISON
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PYRIMETHAMINE, PYRIMETHAMINE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RAMELTEON, RAMELTEON
 RAMIPRIL, RAMIPRIL
 REPAGLINIDE, REPAGLINIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRI-MILI, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AUROBINDO PHARMA LTD
 - URSODIOL, URSODIOL
 - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - VALSARTAN, VALSARTAN
 - VIGABATRIN, VIGABATRIN
 - VORICONAZOLE, VORICONAZOLE
 - ZONISAMIDE, ZONISAMIDE
 - ZUMANDIMINE, DROSPIRENONE
- * AUROBINDO PHARMA LTD INC
 - ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA USA

- * AUROBINDO PHARMA USA INC
 - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 - BUDESONIDE, BUDESONIDE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLONAZEPAM, CLONAZEPAM
 - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 - ELIMITE, PERMETHRIN
 - GLIPIZIDE, GLIPIZIDE
 - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 - MAXZIDE, HYDROCHLOROTHIAZIDE
 - MAXZIDE-25, HYDROCHLOROTHIAZIDE
 - MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 - NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 - PINDOLOL, PINDOLOL
 - RASAGILINE MESYLATE, RASAGILINE MESYLATE
 - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 - TRANEXAMIC ACID, TRANEXAMIC ACID

AUROLIFE PHARMA LLC

- * AUROLIFE PHARMA LLC
 - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 - AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 - LORAZEPAM, LORAZEPAM
 - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 - PHEENTERMINE HYDROCHLORIDE, PHEENTERMINE HYDROCHLORIDE

AUSTARPHARMA

- * AUSTARPHARMA LLC
 - FENOFIBRATE, FENOFIBRATE
 - SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 - TADALAFIL, TADALAFIL

AUSTARPHARMA LLC

- * AUSTARPHARMA LLC
 - METHOCARBAMOL, METHOCARBAMOL

AUXILIUM PHARMS INC

- * AUXILIUM PHARMACEUTICALS INC
 - TESTOPEL, TESTOSTERONE

AUXILIUM PHARMS LLC

- * AUXILIUM PHARMACEUTICALS LLC
 - EDEX, ALPROSTADIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUXILIUM PHARMACEUTICALS LLC
 TESTIM, TESTOSTERONE
 THEO-24, THEOPHYLLINE

AVADEL CNS

* AVADEL CNS PHARMACEUTICALS LLC
 LUMRYZ, SODIUM OXYBATE

AVANIR PHARMS

* AVANIR PHARMACEUTICALS INC
 NUEDEXTA, DEXTROMETHORPHAN HYDROBROMIDE

AVANTHI INC

* AVANTHI INC
 CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 INDOMETHACIN, INDOMETHACIN
 LOMAIRA, PHENTERMINE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE

AVEMA PHARMA

* AVEMA PHARMA SOLUTIONS
 IBUPROFEN, IBUPROFEN (OTC)

AVENT

* AVENT INC
 PYTEST, UREA, C-14

AVEO PHARMS

* AVEO PHARMACEUTICALS INC
 FOTIVDA, TIVOZANIB HYDROCHLORIDE

AVERITAS

* AVERITAS PHARMA INC
 QUTENZA, CAPSAICIN

AVET

* AVET PHARMACEUTICALS INC
 DOXERCALCIFEROL, DOXERCALCIFEROL

AVET LIFESCIENCES

* AVET LIFESCIENCES LTD
 ACARBOSE, ACARBOSE
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ADENOSINE, ADENOSINE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 BICNU, CARMUSTINE
 CIDOFOVIR, CIDOFOVIR
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 FUROSEMIDE, FUROSEMIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPOFOL, PROPOFOL
 TRANEXAMIC ACID, TRANEXAMIC ACID
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

* AVET LIFESCIENCES PRIVATE LTD
 FOSCARNET SODIUM, FOSCARNET SODIUM
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE

AVEVA

* AVEVA DRUG DELIVERY SYSTEMS INC
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE, BUPRENORPHINE
 CLONIDINE, CLONIDINE
 FENTANYL-100, FENTANYL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AVEVA DRUG DELIVERY SYSTEMS INC**

FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-37, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL
 FENTANYL-87, FENTANYL
 NICOTINE, NICOTINE (OTC)

AVID RADIOPHARMS INC

*** AVID RADIOPHARMACEUTICALS INC**
 AMYVID, FLORBETAPIR F-18
 TAUVID, FLORTAUCIPIR F-18

AVION PHARMS

*** AVION PHARMACEUTICALS LLC**
 BALCOLTRA, ETHINYL ESTRADIOL
 DHIVY, CARBIDOPA
 PONSTEL, MEFENAMIC ACID

AVONDALE PHARMS

*** AVONDALE PHARMACEUTICALS LLC**
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 MELOXICAM, MELOXICAM
 NIACOR, NIACIN

AXSOME

*** AXSOME THERAPEUTICS INC**
 AUVELITY, BUPROPION HYDROCHLORIDE

AXSOME MALTA

*** AXSOME MALTA LTD**
 SUNOSI, SOLRIAMFETOL HYDROCHLORIDE

AYANA PHARMA LTD

*** AYANA PHARMA LTD**
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE

AYTU

*** AYTU BIOSCIENCE INC**
 KARBINAL ER, CARBINOXAMINE MALEATE

AYTU BIOPHARMA

*** AYTU BIOPHARMA INC**
 METADATE CD, METHYLPHENIDATE HYDROCHLORIDE

AZURITY

*** AZURITY PHARMACEUTICALS INC**
 BIDIL, HYDRALAZINE HYDROCHLORIDE
 E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
 EDARBI, AZILSARTAN KAMEDOXOMIL
 EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
 EPANED, ENALAPRIL MALEATE
 EPRONTIA, TOPIRAMATE
 ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
 EVEKEO ODT, AMPHETAMINE SULFATE
 FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE
 FLEQSUVY, BACLOFEN
 GLIADEL, CARMUSTINE
 HORIZANT, GABAPENTIN ENACARBIL
 KATERZIA, AMLODIPINE BENZOATE
 KONVOMEF, OMEPRAZOLE
 NYMALIZE, NIMODIPINE
 QBRELIS, LISINOPRIL
 SOTYLIZE, SOTALOL HYDROCHLORIDE
 THYQUIDITY, LEVOTHYROXINE SODIUM
 TRIPTODUR KIT, TRIPTORELIN PAMOATE
 XATMEP, METHOTREXATE SODIUM
 ZONISADE, ZONISAMIDE

**** B ******B BRAUN**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* B BRAUN MEDICAL INC**

ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 AMINO ACIDS, AMINO ACIDS
 BALANCED SALT, CALCIUM CHLORIDE
 CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
 CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
 CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
 DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
 DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
 METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
 NUTRILIPID 10%, SOYBEAN OIL
 NUTRILIPID 20%, SOYBEAN OIL
 PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* B BRAUN MEDICAL INC**

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 TROPHAMINE 10%, AMINO ACIDS
 TROPHAMINE, AMINO ACIDS

B BRAUN MEDICAL INC*** B BRAUN MEDICAL INC**

ACETAMINOPHEN, ACETAMINOPHEN
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 HEPARIN SODIUM, HEPARIN SODIUM
 LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM

BAJAJ

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BAJAJ MEDICAL**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
LACTULOSE, LACTULOSE

*** BAJAJ MEDICAL LLC**

CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

BAMF*** BAMF HEALTH INC**

SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

BANNER LIFE SCIENCES*** BANNER LIFE SCIENCES LLC**

BAFIERTAM, MONOMETHYL FUMARATE

BARR*** BARR LABORATORIES INC**

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
ARANELLE, ETHINYL ESTRADIOL
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
BALZIVA-28, ETHINYL ESTRADIOL
CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
DANAZOL, DANAZOL
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIPYRIDAMOLE, DIPYRIDAMOLE
DUTASTERIDE, DUTASTERIDE
ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
ESTRADIOL AND NORGESTIMATE, ESTRADIOL
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
HYDROXYUREA, HYDROXYUREA
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
ISONIAZID, ISONIAZID
JUNEL 1.5/30, ETHINYL ESTRADIOL
JUNEL 1/20, ETHINYL ESTRADIOL
JUNEL FE 1.5/30, ETHINYL ESTRADIOL
JUNEL FE 1/20, ETHINYL ESTRADIOL
KARIVA, DESOGESTREL
KELNOR, ETHINYL ESTRADIOL
LESSINA-28, ETHINYL ESTRADIOL
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEGESTROL ACETATE, MEGESTROL ACETATE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
NIACIN, NIACIN
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORTREL 0.5/35-28, ETHINYL ESTRADIOL
NORTREL 1/35-21, ETHINYL ESTRADIOL
NORTREL 1/35-28, ETHINYL ESTRADIOL
NORTREL 7/7/7, ETHINYL ESTRADIOL
PORTIA-28, ETHINYL ESTRADIOL
SPRINTEC, ETHINYL ESTRADIOL
Trexall, METHOTREXATE SODIUM
TRI-LEGEST FE, ETHINYL ESTRADIOL
TRI-SPRINTEC, ETHINYL ESTRADIOL
WARFARIN SODIUM, WARFARIN SODIUM

*** BARR LABORATORIES INC A WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC**

LANTHANUM CARBONATE, LANTHANUM CARBONATE

*** BARR PHARMACEUTICALS**

LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM

BARR LABS INC*** BARR LABORATORIES INC**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BARR LABORATORIES INC
 ACITRETIN, ACITRETIN
 CLOZAPINE, CLOZAPINE
 ESTRADIOL, ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 TRETINOIN, TRETINOIN
 TRI LO SPRINTEC, ETHINYL ESTRADIOL

BAUSCH

* BAUSCH HEALTH AMERICAS INC
 ACANYA, BENZOYL PEROXIDE
 BRYHALI, HALOBETASOL PROPIONATE
 DUOBRII, HALOBETASOL PROPIONATE
 EDECRIN, ETHACRYNATE SODIUM
 EDECRIN, ETHACRYNIC ACID
 EFUDEX, FLUOROURACIL
 JUBLIA, EFINACONAZOLE
 LOCID, HYDROCORTISONE BUTYRATE
 ONEXTON, BENZOYL PEROXIDE
 OXSORALEN-ULTRA, METHOXSALEN
 SYPRINE, TRIENTINE HYDROCHLORIDE

* BAUSCH HEALTH IRELAND LTD
 TARGRETIN, BEXAROTENE

* BAUSCH HEALTH US LLC
 AMMONUL, SODIUM BENZOATE
 ANCOBON, FLUCYTOSINE
 APLENZIN, BUPROPION HYDROBROMIDE
 ARAZLO, TAZAROTENE
 ATIVAN, LORAZEPAM
 CABTREO, ADAPALENE
 CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
 CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
 CARDIZEM, DILTIAZEM HYDROCHLORIDE
 CESAMET, NABILONE
 CLINDAGEL, CLINDAMYCIN PHOSPHATE
 DEMSER, METYROSINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIASTAT ACUDIAL, DIAZEPAM
 DIASTAT, DIAZEPAM
 ELIDEL, PIMECROLIMUS
 FLUNISOLIDE, FLUNISOLIDE
 GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZED
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 ISORDIL, ISOSORBIDE DINITRATE
 KLARON, SULFACETAMIDE SODIUM
 LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LOCID, HYDROCORTISONE BUTYRATE
 LOPROX, CICLOPIROX
 LUZU, LULICONAZOLE
 MESTINON, PYRIDOSTIGMINE BROMIDE
 METROGEL-VAGINAL, METRONIDAZOLE
 MIGRANAL, DIHYDROERGOTAMINE MESYLATE
 MINOCIN, MINOCYCLINE HYDROCHLORIDE
 NORITATE, METRONIDAZOLE
 RETIN-A MICRO, TRETINOIN
 RETIN-A, TRETINOIN
 RETIN-A-MICRO, TRETINOIN
 SOLODYN, MINOCYCLINE HYDROCHLORIDE
 TASMAR, TOLCAPONE
 TIAZAC, DILTIAZEM HYDROCHLORIDE
 VANOS, FLUOCINONIDE
 VASERETIC, ENALAPRIL MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BAUSCH HEALTH US LLC**

VASOTEC, ENALAPRIL MALEATE
 VIRAZOLE, RIBAVIRIN
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 XENAZINE, TETRABENAZINE
 XERESE, ACYCLOVIR
 ZELAPAR, SELEGILINE HYDROCHLORIDE
 ZIANA, CLINDAMYCIN PHOSPHATE
 ZOVIRAX, ACYCLOVIR
 ZYCLARA, IMIQUIMOD

BAUSCH AND LOMB*** BAUSCH AND LOMB INC**

ALAWAY, KETOTIFEN FUMARATE (OTC)
 ALREX, LOTEPIEDNOL ETABONATE
 BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
 BRINZOLAMIDE, BRINZOLAMIDE
 CHILDREN'S ALAWAY, KETOTIFEN FUMARATE (OTC)
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 ISTALOL, TIMOLOL MALEATE
 LATANOPROST, LATANOPROST
 LOTEMAX, LOTEPIEDNOL ETABONATE
 MIOCHOL-E, ACETYLCHOLINE CHLORIDE
 OFLOXACIN, OFLOXACIN
 OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 PROLENSA, BROMFENAC SODIUM
 RETISERT, FLUOCINOLONE ACETONIDE
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TROPICAMIDE, TROPICAMIDE
 VYZULTA, LATANOPROSTENE BUNOD
 ZIRGAN, GANCICLOVIR
 ZYLET, LOTEPIEDNOL ETABONATE

*** BAUSCH AND LOMB PHARMACEUTICALS INC**

BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXASPORIN, DEXAMETHASONE
 ERYTHROMYCIN, ERYTHROMYCIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 OFLOXACIN, OFLOXACIN
 OTICAIR, HYDROCORTISONE
 PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
 TOBRAMYCIN, TOBRAMYCIN
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TROPICAMIDE, TROPICAMIDE

BAUSCH AND LOMB INC*** BAUSCH AND LOMB INC**

ALAWAY, KETOTIFEN FUMARATE (OTC)
 ATROPINE SULFATE, ATROPINE SULFATE
 BEPREVE, BEPOTASTINE BESILATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BAUSCH AND LOMB INC**

LACRISERT, HYDROXYPROPYL CELLULOSE
 LOTEMAX SM, LOTEPRDNOL ETABONATE
 LOTEMAX, LOTEPRDNOL ETABONATE
 LUMIFY, BRIMONIDINE TARTRATE (OTC)
 METHAZOLAMIDE, METHAZOLAMIDE
 MIEBO, PERFLUOROHEXYLOCTANE
 TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
 TIMOPTIC, TIMOLOL MALEATE
 TIMOPTIC-XE, TIMOLOL MALEATE
 XIPERE, TRIAMCINOLONE ACETONIDE

BAUSCH LOMB IRELAND*** BAUSCH AND LOMB IRELAND LTD**

FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE, BENOXINATE HYDROCHLORIDE
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE

BAXTER HLTHCARE*** BAXTER HEALTHCARE CORP**

ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
 BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
 BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC, ESMOLOL HYDROCHLORIDE
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFTRIAZONE IN PLASTIC CONTAINER, CEFTRIAZONE SODIUM
 CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC
 CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC
 CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYTOXAN, CYCLOPHOSPHAMIDE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 EXTRANEAL, ICODEXTRIN
 FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FORANE, ISOFLURANE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 IFEX, IFOSFAMIDE
 LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 MESNEX, MESNA
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 NALLPEN IN PLASTIC CONTAINER, NAFICILLIN SODIUM
 NEXTERONE, AMIODARONE HYDROCHLORIDE
 NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
 OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 10% IN WATER, MANNITOL
 OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 15% IN WATER, MANNITOL
 OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 20% IN WATER, MANNITOL
 OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 5% IN WATER, MANNITOL
 PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
 PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SEVOFLURANE, SEVOFLURANE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 STERILE WATER, STERILE WATER FOR IRRIGATION
 SUPRANE, DESFLURANE
 TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 TIS-U-SOL, MAGNESIUM SULFATE
 TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
 VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

* BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV

PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

BAXTER HLTHCARE CORP

* BAXTER HEALTHCARE CORP

ACETAMINOPHEN, ACETAMINOPHEN
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BORTEZOMIB, BORTEZOMIB
 CEFAZOLIN IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLINOLIPID 20%, OLIVE OIL
 DAPTOMYCIN IN 0.9% SODIUM CHLORIDE, DAPTOMYCIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 FLUMAZENIL, FLUMAZENIL
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOFLOXACIN, LEVOFLOXACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE CORP
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MICAFUNGIN IN SODIUM CHLORIDE 0.9%, MICAFUNGIN SODIUM
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE IN 5% DEXTROSE, NOREPINEPHRINE BITARTRATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 REGADENOSON, REGADENOSON
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANSDERM SCOP, SCOPOLAMINE
 VASOPRESSIN IN SODIUM CHLORIDE 0.9%, VASOPRESSIN
 ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
- * BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
 PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE

BAYER

- * BAYER HEALTHCARE LLC
 ALEVE, NAPROXEN SODIUM (OTC)

BAYER HEALTHCARE

- * BAYER HEALTHCARE PHARMACEUTICALS INC
 LAMPIT, NIFURTIMOX
 NUBEQA, DAROLUTAMIDE
 VITRAKVI, LAROTRECTINIB SULFATE

BAYER HEALTHCARE LLC

- * BAYER HEALTHCARE LLC
 CHILDREN'S CLARITIN, LORATADINE (OTC)
 CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
 CLARITIN HIVES RELIEF, LORATADINE (OTC)
 CLARITIN REDITABS, LORATADINE (OTC)
 CLARITIN, LORATADINE (OTC)
 CLARITIN-D 24 HOUR, LORATADINE (OTC)
 CLARITIN-D, LORATADINE (OTC)
 LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
 MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)

BAYER HLTHCARE

- * BAYER HEALTHCARE CONSUMER CARE
 ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
 CHILDREN'S ASTEPRO ALLERGY, AZELASTINE HYDROCHLORIDE (OTC)
- * BAYER HEALTHCARE PHARMACEUTICALS INC
 ADEMPAS, RIOCIQUAT
 ANGELIQ, DROSPIRENONE
 BEYAZ, DROSPIRENONE
 BILTRICIDE, PRAZIQUANTEL
 CIPRO, CIPROFLOXACIN
 CIPRO, CIPROFLOXACIN HYDROCHLORIDE
 CLIMARA PRO, ESTRADIOL
 CLIMARA, ESTRADIOL
 EOVIIST, GADOXETATE DISODIUM
 GADAVIST, GADOBUTROL
 KERENDIA, FINERENONE
 KYLEENA, LEVONORGESTREL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BAYER HEALTHCARE PHARMACEUTICALS INC**

MENOSTAR, ESTRADIOL
 MIRENA, LEVONORGESTREL
 NATAZIA, DIENOGEST
 NEXAVAR, SORAFENIB TOSYLATE
 SAFYRAL, DROSPIRENONE
 SKYLA, LEVONORGESTREL
 STIVARGA, REGORAFENIB
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 VITRAKVI, LAROTRECTINIB SULFATE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE

BAYSHORE PHARMS LLC*** BAYSHORE PHARMACEUTICALS LLC**

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 ETODOLAC, ETODOLAC
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METOLAZONE, METOLAZONE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PINDOLOL, PINDOLOL
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE

BDSI*** BIODELIVERY SCIENCES INTERNATIONAL INC**

BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 SYMPROIC, NALDEMEDINE TOSYLATE

BE PHARMS*** BE PHARMACEUTICALS AG**

DAPTOMYCIN, DAPTOMYCIN
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 HEPARIN SODIUM, HEPARIN SODIUM
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

BECTON DICKINSON*** BECTON DICKINSON AND CO**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)

BECTON DICKINSON CO*** BECTON DICKINSON AND CO**

CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)

BEIGENE*** BEIGENE USA INC**

BRUKINSA, ZANUBRUTINIB

BEIJING*** BEIJING SCIECURE PHARMACEUTICAL CO LTD**

FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 FOSCARNET SODIUM, FOSCARNET SODIUM

BEIJING TIDE PHARM*** BEIJING TIDE PHARMACEUTICAL CO LTD**

COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE

BEIJING YILING*** BEIJING YILING BIO-ENGINEERING AND TECHNOLOGY CO LTD**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BEIJING YILING BIO-ENGINEERING AND TECHNOLOGY CO LTD
ANASTROZOLE, ANASTROZOLE
LETROZOLE, LETROZOLE

BELCHER

* BELCHER PHARMACEUTICALS LLC
AMINOCAPROIC ACID, AMINOCAPROIC ACID
CEFIXIME, CEFIXIME
CHLORZOXAZONE, CHLORZOXAZONE
GABAPENTIN, GABAPENTIN
LEVETIRACETAM, LEVETIRACETAM
MEFENAMIC ACID, MEFENAMIC ACID
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
TACROLIMUS, TACROLIMUS

BELCHER PHARMS

* BELCHER PHARMACEUTICALS LLC
CEPHALEXIN, CEPHALEXIN
DESLORATADINE, DESLORATADINE

BELOTECA

* BELOTECA INC
DAPSONE, DAPSONE
DIAZEPAM, DIAZEPAM
THIOTEPA, THIOTEPA

BENUVIA OPERATIONS

* BENUVIA OPERATIONS LLC
SYNDROS, DRONABINOL

BEOWULF ASSET

* BEOWULF ASSET MANAGEMENT LP
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
NYSTATIN, NYSTATIN
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

BESINS HLTHCARE

* BESINS HEALTHCARE IRELAND LTD
ANDROGEL, TESTOSTERONE

BEXIMCO PHARMS USA

* BEXIMCO PHARMACEUTICALS USA INC
BACLOFEN, BACLOFEN
CARBIDOPA, CARBIDOPA
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
NADOLOL, NADOLOL
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

BEXIMCO USA

* BEXIMCO PHARMACEUTICALS USA INC
CARVEDILOL, CARVEDILOL

BIOCODEX SA

* BIOCODEX SA
DIACOMIT, STIRIPENTOL

BIOCON PHARMA

* BIOCON PHARMA INC
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
* BIOCON PHARMA LTD
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
EVEROLIMUS, EVEROLIMUS
MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
POSACONAZOLE, POSACONAZOLE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BIOCON PHARMA LTD
SIMVASTATIN, SIMVASTATIN
TACROLIMUS, TACROLIMUS
TERIFLUNOMIDE, TERIFLUNOMIDE

BIOCRYST

* BIOCRYST PHARMACEUTICALS INC
ORLADEYO, BEROTRALSTAT HYDROCHLORIDE
RAPIVAB, PERAMIVIR

BIOFRONTERA

* BIOFRONTERA BIOSCIENCE GMBH
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE

BIOGEN IDEC

* BIOGEN IDEC INC
SPINRAZA, NUSINERSEN SODIUM

BIOGEN INC

* BIOGEN INC
TECFIDERA, DIMETHYL FUMARATE
VUMERITY, DIROXIMEL FUMARATE
ZURZUVAE, ZURANOLONE

BIOGEN MA

* BIOGEN MA INC
QALSODY, TOFERSEN

BIOLINERX LTD

* BIOLINERX LTD
APHEXDA, MOTIXAFORTIDE ACETATE

BIOMARIN PHARM

* BIOMARIN PHARMACEUTICAL INC
KUVAN, SAPROPTERIN DIHYDROCHLORIDE
VOXZOGO, VOSORITIDE

BIOMEDCL RES FDN

* BIOMEDICAL RESEARCH FOUNDATION NORTHWEST LOUISIANA
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

BIONPHARMA

* BIONPHARMA INC
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
ATOVAQUONE, ATOVAQUONE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
AZITHROMYCIN, AZITHROMYCIN
BENZONATATE, BENZONATATE
BEXAROTENE, BEXAROTENE
CALCITRIOL, CALCITRIOL
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
CLOBAZAM, CLOBAZAM
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
DEFERASIROX, DEFERASIROX
DEXAMETHASONE, DEXAMETHASONE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DOFETILIDE, DOFETILIDE
DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
DROXIDOPA, DROXIDOPA
DUTASTERIDE, DUTASTERIDE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ETHOSUXIMIDE, ETHOSUXIMIDE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BIONPHARMA INC
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVETIRACETAM, LEVETIRACETAM
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 METHIMAZOLE, METHIMAZOLE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MIDOL LIQUID GELS, IBUPROFEN (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PARICALCITOL, PARICALCITOL
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROGESTERONE, PROGESTERONE
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 TETRABENAZINE, TETRABENAZINE
 THEOPHYLLINE, THEOPHYLLINE
 VALPROIC ACID, VALPROIC ACID
 VITAMIN D, ERGOCALCIFEROL
 ZONISAMIDE, ZONISAMIDE

BIOXCEL

* BIOXCEL THERAPEUTICS INC
 IGALMI, DEXMEDETOMIDINE HYDROCHLORIDE

BLUE EARTH

* BLUE EARTH DIAGNOSTICS LTD
 AXUMIN, FLUCICLOVINE F-18
 POSLUMA, FLOTUFOLASTAT F-18 GALLIUM

BLUE WATER BIOTECH

* BLUE WATER BIOTECH INC
 ENTADFI, FINASTERIDE

BLUEPHARMA INDUSTRIA

* BLUEPHARMA INDUSTRIA FARMACEUTICA SA
 DROXIDOPA, DROXIDOPA

BLUEPRINT MEDICINES

* BLUEPRINT MEDICINES CORP
 AYVAKIT, AVAPRITINIB

BOEHRINGER INGELHEIM

* BOEHRINGER INGELHEIM
 GILOTRIF, AFATINIB DIMALEATE
 GLYXAMBI, EMPAGLIFLOZIN
 MICARDIS HCT, HYDROCHLOROTHIAZIDE
 MICARDIS, TELMISARTAN

* BOEHRINGER INGELHEIM PHARMACEUTICALS INC
 APTIVUS, TIPRANAVIR
 ATROVENT HFA, IPRATROPIUM BROMIDE
 COMBIVENT RESPIMAT, ALBUTEROL SULFATE
 JARDIANCE, EMPAGLIFLOZIN
 JENTADUETO XR, LINAGLIPTIN
 JENTADUETO, LINAGLIPTIN
 MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE
 MOBIC, MELOXICAM
 OFEV, NINTEDANIB ESYLATE
 PERSANTINE, DIPYRIDAMOLE
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE
 SPIRIVA, TIOTROPIUM BROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BOEHRINGER INGELHEIM PHARMACEUTICALS INC
 STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
 STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE
 SYNJARDY XR, EMPAGLIFLOZIN
 SYNJARDY, EMPAGLIFLOZIN
 TRADJENTA, LINAGLIPTIN
 TRIJARDY XR, EMPAGLIFLOZIN
 VIRAMUNE XR, NEVIRAPINE
 VIRAMUNE, NEVIRAPINE

BPI LABS

* BPI LABS LLC
 ABLYSINOL, ALCOHOL
 EPINEPHRINE, EPINEPHRINE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

BRACCO

* BRACCO DIAGNOSTICS INC
 CARADIOGEN-82, RUBIDIUM CHLORIDE RB-82
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 E-Z-HD, BARIUM SULFATE
 E-Z-PAQUE, BARIUM SULFATE
 ENTERO VU 24%, BARIUM SULFATE
 GASTROGRAFIN, DIATRIZOATE MEGLUMINE
 ISOVUE-200, IOPAMIDOL
 ISOVUE-250, IOPAMIDOL
 ISOVUE-300, IOPAMIDOL
 ISOVUE-370, IOPAMIDOL
 ISOVUE-M 200, IOPAMIDOL
 ISOVUE-M 300, IOPAMIDOL
 KINEVAC, SINCALIDE
 LIQUID E-Z-PAQUE, BARIUM SULFATE
 LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
 MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
 MULTIHANCE, GADOBENATE DIMEGLUMINE
 PROHANCE MULTIPACK, GADOTERIDOL
 PROHANCE, GADOTERIDOL
 READI-CAT 2 SMOOTHIE, BARIUM SULFATE
 READI-CAT 2, BARIUM SULFATE
 TAGITOL V, BARIUM SULFATE
 VARIBAR HONEY, BARIUM SULFATE
 VARIBAR NECTAR, BARIUM SULFATE
 VARIBAR PUDDING, BARIUM SULFATE
 VARIBAR THIN HONEY, BARIUM SULFATE
 VARIBAR THIN LIQUID, BARIUM SULFATE

BRAEBURN

* BRAEBURN INC
 BRIXADI, BUPRENORPHINE

BRAINTREE

* BRAINTREE LABORATORIES INC
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350

BRAINTREE LABS

* BRAINTREE LABORATORIES INC
 SUFLAVE, MAGNESIUM SULFATE
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE
 SUTAB, MAGNESIUM SULFATE

BRECKENRIDGE

* BRECKENRIDGE PHARMACEUTICAL INC
 ALPRAZOLAM, ALPRAZOLAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BRECKENRIDGE PHARMACEUTICAL INC
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 CLOBAZAM, CLOBAZAM
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPLERENONE, EPLERENONE
 EVEROLIMUS, EVEROLIMUS
 EXEMESTANE, EXEMESTANE
 LAMIVUDINE, LAMIVUDINE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MIGLUSTAT, MIGLUSTAT
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OMEPRAZOLE, OMEPRAZOLE
 PENICILLAMINE, PENICILLAMINE
 RIVASTIGMINE, RIVASTIGMINE
 ROFLUMILAST, ROFLUMILAST
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

BRECKENRIDGE PHARM

* BRECKENRIDGE PHARMACEUTICAL INC
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

BRIGHAM WOMENS

* BRIGHAM AND WOMENS HOSP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

BRIGHAM WOMENS HOSP

* BRIGHAM AND WOMENS HOSP INC
 AMMONIA N 13, AMMONIA N-13

BRIGHTGENE

* BRIGHTGENE BIO-MEDICAL TECHNOLOGY CO LTD
 ENTECAVIR, ENTECAVIR

BRISTOL

* BRISTOL MYERS SQUIBB CO
 AUGTYRO, REPOTRECTINIB
 CAMZYOS, MAVACAMTEN
 EVOTAZ, ATAZANAVIR SULFATE
 ONUREG, AZACITIDINE
 POMALYST, POMALIDOMIDE
 SOTYKTU, DEUCRAVACITINIB

BRISTOL MYERS SQUIBB

* BRISTOL MYERS SQUIBB
 AZACTAM, AZTREONAM
 BARACLUDE, ENTECAVIR
 IDHIFA, ENASIDENIB MESYLATE
 REVLIMID, LENALIDOMIDE

* BRISTOL MYERS SQUIBB CO
 REYATAZ, ATAZANAVIR SULFATE
 SPRYCEL, DASATINIB

* BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
 ELIQUIS, APIXABAN

BRISTOL-MYERS

* BRISTOL-MYERS SQUIBB CO
 ABRAKANE, PACLITAXEL
 ISTODAX, ROMIDEPSIN
 THALOMID, THALIDOMIDE
 VIDAZA, AZACITIDINE

BROOKS STERISCIENCE

* BROOKS STERISCIENCE LTD
 MEROPENEM, MEROPENEM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******BTG INTL**

* BTG INTERNATIONAL INC
 CYANOKIT, HYDROXOCOBALAMIN
 THYROSAFE, POTASSIUM IODIDE (OTC)
 VISTOGARD, URIDINE TRIACETATE
 XURIDEN, URIDINE TRIACETATE

BWXT ITG

* BWXT ITG CANADA INC
 INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE

**** C ******CADILA**

* CADILA HEALTHCARE LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DUTASTERIDE, DUTASTERIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FELBAMATE, FELBAMATE
 FLUOCINONIDE, FLUOCINONIDE
 INDOMETHACIN, INDOMETHACIN
 LEVOFLOXACIN, LEVOFLOXACIN
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MODAFINIL, MODAFINIL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 PIROXICAM, PIROXICAM
 RANOLAZINE, RANOLAZINE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

CADILA PHARMS LTD

* CADILA PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ERYTHROMYCIN, ERYTHROMYCIN
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 GEMFIBROZIL, GEMFIBROZIL
 GLYBURIDE, GLYBURIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 NATEGLINIDE, NATEGLINIDE
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* CADILA PHARMACEUTICALS LTD**

OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TELMISARTAN, TELMISARTAN
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

CALL INC

*** CALL INC DBA ROCHESTER PHARMACEUTICALS**
 ADAPALENE, ADAPALENE

CALLIDITAS

*** CALLIDITAS THERAPEUTICS AB**
 TARPEYO, BUDESONIDE

CAPELLON PHARMS LLC

*** CAPELLON PHARMACEUTICALS LLC**
 POLMON, DEXCHLORPHENIRAMINE MALEATE

CAPLIN

*** CAPLIN STERILES LTD**
 ARGATROBAN, ARGATROBAN
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 ETOMIDATE, ETOMIDATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OFLOXACIN, OFLOXACIN
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

CARA THERAP

*** CARA THERAPEUTICS INC**
 KORSUVA, DIFELIKEFALIN ACETATE

CARDINAL HEALTH 414

*** CARDINAL HEALTH 414 LLC CARDINAL HEALTH NUCLEAR PHARMACY SERVICES**
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEDRONATE KIT, TECHNETIUM TC-99M MEDRONATE KIT

CARDINAL HEALTH 418

*** CARDINAL HEALTH 418 INC**
 SODIUM IODIDE I 123, SODIUM IODIDE I-123

CARDINAL HLTH 414

*** CARDINAL HEALTH 414 LLC**
 AMMONIA N 13, AMMONIA N-13

CARIBE HOLDINGS

*** CARIBE HOLDINGS CAYMAN CO LTD DBA PURACAP CARIBE**
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 GEMFIBROZIL, GEMFIBROZIL

CARLSBAD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

- * CARLSBAD TECHNOLOGY INC
 ACYCLOVIR, ACYCLOVIR
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 FAMOTIDINE, FAMOTIDINE
 GLIMEPIRIDE, GLIMEPIRIDE
 LOVASTATIN, LOVASTATIN
- CARLSBAD TECHNOLOGY**
- * CARLSBAD TECHNOLOGY INC
 ACYCLOVIR, ACYCLOVIR
- CARNEGIE**
- * CARNEGIE PHARMACEUTICALS LLC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 ETRAVIRINE, ETRAVIRINE
 FAMOTIDINE, FAMOTIDINE
 LATANOPROST, LATANOPROST
 LETROZOLE, LETROZOLE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
- CASI PHARMS INC**
- * CASI PHARMACEUTICALS INC
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
- CASPER PHARMA LLC**
- * CASPER PHARMA LLC
 ANTIVERT, MECLIZINE HYDROCHLORIDE
 AQUASOL A, VITAMIN A PALMITATE
 CASPORYN HC, HYDROCORTISONE
 EDETATE CALCIUM DISODIUM, EDETATE CALCIUM DISODIUM
 FURADANTIN, NITROFURANTOIN
 LUMI-SPORYN, BACITRACIN ZINC
 ROBINUL FORTE, GLYCOPYRROLATE
 ROBINUL, GLYCOPYRROLATE
 THALITONE, CHLORTHALIDONE
 ZYLOPRIM, ALLOPURINOL
- CATALENT**
- * CATALENT PHARMA SOLUTIONS LLC
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 VALPROIC ACID, VALPROIC ACID
- CATALYST PHARMS**
- * CATALYST PHARMACEUTICALS INC
 AGAMREE, VAMOROLONE
 FIRDAPSE, AMIFAMPRIDINE PHOSPHATE
 FYCOMPA, PERAMPANEL
- CEDIPROF INC**
- * CEDIPROF INC
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 LEVO-T, LEVOTHYROXINE SODIUM **
- CELATOR PHARMS**
- * CELATOR PHARMACEUTICALS INC
 VYXEOS, CYTARABINE
- CELGENE INTL**
- * CELGENE INTERNATIONAL II SARL
 ZEPOSIA, OZANIMOD HYDROCHLORIDE
- CELLTRION**
- * CELLTRION INC
 LEVOFLOXACIN, LEVOFLOXACIN
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
- CEPHALON**
- * CEPHALON INC
 GABITRIL, TIAGABINE HYDROCHLORIDE
 NUVIGIL, ARMODAFINIL
 TREANDA, BENDAMUSTINE HYDROCHLORIDE
 TRISENOX, ARSENIC TRIOXIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* CEPHALON LLC**

ACTIQ, FENTANYL CITRATE
 FENTORA, FENTANYL CITRATE
 PROVIGIL, MODAFINIL

CEROVENE INC*** CEROVENE INC**

AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 PYRIMETHAMINE, PYRIMETHAMINE
 SEROMYCIN, CYCLOSERINE

CHANGZHOU PHARM*** CHANGZHOU PHARMACEUTICAL FACTORY**

CAPTOPRIL, CAPTOPRIL
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 PREGABALIN, PREGABALIN
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

CHARTWELL*** CHARTWELL LIFE MOLECULES LLC**

ALLOPURINOL, ALLOPURINOL
 AMOXICILLIN, AMOXICILLIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CIPROFLOXACIN, CIPROFLOXACIN
 CLARITHROMYCIN, CLARITHROMYCIN
 COLOCORT, HYDROCORTISONE
 FLUCONAZOLE, FLUCONAZOLE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

*** CHARTWELL PHARMA SCIENCE LLC**

SULFASALAZINE, SULFASALAZINE

*** CHARTWELL SCHEDULED LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE

CHARTWELL INJECTABLE*** CHARTWELL INJECTABLES LLC**

DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 TERBUTALINE SULFATE, TERBUTALINE SULFATE

CHARTWELL MOLECULAR*** CHARTWELL MOLECULAR HOLDINGS LLC**

ACETAZOLAMIDE, ACETAZOLAMIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CALCIUM ACETATE, CALCIUM ACETATE
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXAMETHASONE, DEXAMETHASONE
 DIAZEPAM, DIAZEPAM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FOLIC ACID, FOLIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHARTWELL MOLECULAR HOLDINGS LLC
 GLIMEPIRIDE, GLIMEPIRIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 OLANZAPINE, OLANZAPINE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 RAMIPRIL, RAMIPRIL
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 THEOPHYLLINE, THEOPHYLLINE
 ZALEPLON, ZALEPLON
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

CHARTWELL MOLECULES

* CHARTWELL MOLECULES LLC
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DISULFIRAM, DISULFIRAM
 GEMFIBROZIL, GEMFIBROZIL
 INDOMETHACIN, INDOMETHACIN
 NABUMETONE, NABUMETONE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

CHARTWELL RX

* CHARTWELL RX SCIENCES LLC
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATOVAQUONE, ATOVAQUONE
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCIUM ACETATE, CALCIUM ACETATE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARISOPRODOL, CARISOPRODOL
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLOROTHALIDONE, CHLOROTHALIDONE
 CICLOPIROX, CICLOPIROX
 CILOSTAZOL, CILOSTAZOL
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** C **

* CHARTWELL RX SCIENCES LLC
 DEFERASIROX, DEFERASIROX
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DUVOID, BETHANECHOL CHLORIDE
 EPLERENONE, EPLERENONE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE, FENOFIBRATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FULVICIN P/G 165, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G 330, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN P/G, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 FULVICIN-U/F, GRISEOFULVIN, MICROCRYSTALLINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HALCINONIDE, HALCINONIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INDOMETHACIN, INDOMETHACIN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACTULOSE, LACTULOSE
 LEVETIRACETAM, LEVETIRACETAM
 LINEZOLID, LINEZOLID
 LISINOPRIL, LISINOPRIL
 LOCHOLEST LIGHT, CHOLESTYRAMINE
 LOCHOLEST, CHOLESTYRAMINE
 LOVASTATIN, LOVASTATIN
 METHIMAZOLE, METHIMAZOLE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METRONIDAZOLE, METRONIDAZOLE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MONODOX, DOXYCYCLINE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISOLONE, PREDNISOLONE
 PREGABALIN, PREGABALIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 REPAGLINIDE, REPAGLINIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHARTWELL RX SCIENCES LLC
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TAVABOROLE, TAVABOROLE
 TELMISARTAN, TELMISARTAN
 TETRABENAZINE, TETRABENAZINE
 TINIDAZOLE, TINIDAZOLE
 TOBRAMYCIN, TOBRAMYCIN
 TRAVOPROST, TRAVOPROST
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 URSODIOL, URSODIOL
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

CHARTWELL TETRA

* CHARTWELL TETRA LLC
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

CHATTEM

* CHATTEM INC
 UNISOM, DOXYLAMINE SUCCINATE (OTC)

CHATTEM SANOFI

* CHATTEM INC DBA SANOFI CONSUMER HEALTHCARE
 ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
 NICODERM CQ, NICOTINE (OTC)
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

CHEMI SPA

* CHEMI SPA
 DECITABINE, DECITABINE
 TEMOZOLOMIDE, TEMOZOLOMIDE

CHEMISCH FBRK KRSSLR

* CHEMISCHE FABRIK KREUSSLER & CO. GMBH
 ASCLERA, POLIDOCANOL

CHEMISTRY HLTH

* CHEMISTRY AND HEALTH FZ LLC
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE

CHEMO RESEARCH SL

* CHEMO RESEARCH SL
 BENZNIDAZOLE, BENZNIDAZOLE
 ESTRADIOL, ESTRADIOL
 NUVESSA, METRONIDAZOLE

CHEMOCENTRYX

* CHEMOCENTRYX INC
 TAVNEOS, AVACOPAN

CHENGDU SHUODE

* CHENGDU SHUODE PHARMACEUTICAL CO LTD
 NALMEFENE HYDROCHLORIDE, NALMEFENE HYDROCHLORIDE

CHEPLAPHARM

* CHEPLAPHARM ARZNEIMITTEL GMBH
 DROXIA, HYDROXYUREA
 ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 HYDREA, HYDROXYUREA
 KLONOPIN, CLONAZEPAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHEPLAPHARM ARZNEIMITTEL GMBH
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
XELODA, CAPECITABINE
XENICAL, ORLISTAT

* CHEPLAPHARM REGISTRATION GMBH
ZYPREXA RELPREVV, OLANZAPINE PAMOATE
ZYPREXA ZYDIS, OLANZAPINE
ZYPREXA, OLANZAPINE

CHIA TAI TIANQING

* CHIA TAI TIANQING PHARMACEUTICAL GROUP CO LTD
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
FULVESTRANT, FULVESTRANT

CHIESI

* CHIESI FARMACEUTICI SPA
MYCAPSSA, OCTREOTIDE ACETATE

* CHIESI USA INC
BETHKIS, TOBRAMYCIN
BRONCHITOL, MANNITOL
CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CLEVIPREX, CLEVIDIPINE
FERRIPROX, DEFERIPRONE
KENGREAL, CANGRELOR
ZYFLO, ZILEUTON

CHILDRENS HOSP MI

* CHILDRENS HOSP MICHIGAN
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

CHINA RESOURCES

* CHINA RESOURCES SAIKE PHARMACEUTICAL CO LTD
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

CHIRHOCLIN

* CHIRHOCLIN INC
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN

CIDARA THERAPS

* CIDARA THERAPEUTICS INC
REZZAYO, REZAFUNGIN ACETATE

CINTEX SVCS

* CINTEX SERVICES LLC
DESONIDE, DESONIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

CIPHER PHARMS INC

* CIPHER PHARMACEUTICALS INC
CONZIP, TRAMADOL HYDROCHLORIDE
LIPOFEN, FENOFIBRATE

CIPLA

* CIPLA LTD
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
ABACAVIR SULFATE, ABACAVIR SULFATE
ACYCLOVIR, ACYCLOVIR
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
AMBRISANTAN, AMBRISANTAN
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ANASTROZOLE, ANASTROZOLE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
AZACITIDINE, AZACITIDINE
BUDESONIDE, BUDESONIDE
CELECOXIB, CELECOXIB
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
DARUNAVIR, DARUNAVIR
DEFERASIROX, DEFERASIROX

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* CIPLA LTD**

DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLUPREDNATE, DIFLUPREDNATE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE, EMTRICITABINE
 ENTECAVIR, ENTECAVIR
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 EXEMESTANE, EXEMESTANE
 FENOFIBRATE, FENOFIBRATE
 FINASTERIDE, FINASTERIDE
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LENALIDOMIDE, LENALIDOMIDE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 MELOXICAM, MELOXICAM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 PHYTONADIONE, PHYTONADIONE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREGABALIN, PREGABALIN
 RITONAVIR, RITONAVIR
 SUMATRIPTAN, SUMATRIPTAN
 TADALAFIL, TADALAFIL
 TAVABOROLE, TAVABOROLE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE, TESTOSTERONE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VASOPRESSIN, VASOPRESSIN
 ZIDOVUDINE, ZIDOVUDINE

CIPLA LTD*** CIPLA LTD**

DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 ZIDOVUDINE, ZIDOVUDINE

CIPLA USA*** CIPLA USA INC**

ZEMDRI, PLAZOMICIN SULFATE

CISEN*** CISEN PHARMACEUTICAL CO LTD**

ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM

CLINIGEN*** CLINIGEN INC**

ETHYOL, AMIFOSTINE

CLINIGEN HLTHCARE*** CLINIGEN HEALTHCARE LTD**

FOSCAVIR, FOSCARNET SODIUM

CLIVUNEL INC*** CLINUNEL INC**

SCENESSE, AFAMELANOTIDE

CMP DEV LLC*** CMP DEVELOPMENT LLC**

ATORVALIQ, ATORVASTATIN CALCIUM
 CAROSPIR, SPIRONOLACTONE
 LIQREV, SILDENAFIL CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* CMP DEVELOPMENT LLC**

NORLIQVA, AMLODIPINE BESYLATE
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 TADLIQ, TADALAFIL

CMP PHARMA INC*** CMP PHARMA INC**

AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 HYDROCORTISONE IN ABSORBABLE, HYDROCORTISONE
 ISONIAZID, ISONIAZID
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SPS, SODIUM POLYSTYRENE SULFONATE

COLGATE-PALMOLIVE CO*** COLGATE-PALMOLIVE CO**

PERIOGARD, CHLORHEXIDINE GLUCONATE

COLLEGIUM PHARM INC*** COLLEGIUM PHARMACEUTICAL INC**

NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
 NUCYNTA, TAPENTADOL HYDROCHLORIDE
 XTAMPZA ER, OXYCODONE

COMBE*** COMBE INC**

VAGISTAT-1, TIOCONAZOLE (OTC)

COMMAVE THERAP*** COMMAVE THERAPEUTICS SA**

AZSTARIS, DEXMETHYLPHENIDATE HYDROCHLORIDE

CONCORD BIOTECH LTD*** CONCORD BIOTECH LTD**

MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 TACROLIMUS, TACROLIMUS

CONCORDIA*** CONCORDIA PHARMACEUTICALS INC**

DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
 DYRENIUM, TRIAMTERENE
 LANOXIN, DIGOXIN
 NILANDRON, NILUTAMIDE
 PANRETIN, ALITRETINOIN
 PARNATE, TRANYLCPROMINE SULFATE
 PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 ZONEGRAN, ZONISAMIDE

CONCORDIA PHARMS INC*** CONCORDIA PHARMACEUTICALS INC**

ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE

CONTRACT PHARMACAL*** CONTRACT PHARMACAL CORP**

CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 FAMOTIDINE, FAMOTIDINE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

COOPERSURGICAL*** COOPERSURGICAL INC**

PARAGARD T 380A, COPPER

CORCEPT THERAP*** CORCEPT THERAPEUTICS INC**

KORLYM, MIFEPRISTONE

CORDEN PHARMA*** CORDEN PHARMA LATINA SPA**

GLEOSTINE, LOMUSTINE

COREPHARMA*** COREPHARMA LLC**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* COREPHARMA LLC**

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CAPTOPRIL, CAPTOPRIL
 CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORZOXAZONE, CHLORZOXAZONE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 LISINOPRIL, LISINOPRIL
 LOVASTATIN, LOVASTATIN
 MELOXICAM, MELOXICAM
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TORSEMIDE, TORSEMIDE

CORIUM*** CORIUM INC**

ADLARITY, DONEPEZIL HYDROCHLORIDE

CORMEDIX*** CORMEDIX INC**

DEFENCATH, HEPARIN SODIUM

COSETTE*** COSETTE PHARMACEUTICALS INC**

ACEPHEN, ACETAMINOPHEN (OTC)
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AZOR, AMLODIPINE BESYLATE
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 BETA-VAL, BETAMETHASONE VALERATE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CICLOPIROX, CICLOPIROX
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
 DAPSONE, DAPSONE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DOXYCYCLINE, DOXYCYCLINE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 INDOMETHACIN, INDOMETHACIN
 LIDOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MIGERGOT, CAFFEINE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MYKACET, NYSTATIN
 NYSTATIN, NYSTATIN
 PROCHLORPERAZINE, PROCHLORPERAZINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* COSETTE PHARMACEUTICALS INC
 TAZAROTENE, TAZAROTENE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIBENZOR, AMLODIPINE BESYLATE
 WELCHOL, COLESEVELAM HYDROCHLORIDE

COVIS

* COVIS PHARMA GMBH
 ALTOPREV, LOVASTATIN
 ALVESCO, CICLESONIDE
 DUAKLIR PRESSAIR, ACLIDINIUM BROMIDE
 FERAHEME, FERUMOXYTOL
 LANOXIN PEDIATRIC, DIGOXIN
 LANOXIN, DIGOXIN
 OMNARIS, CICLESONIDE
 PRILOSEC, OMEPRAZOLE MAGNESIUM
 RILUTEK, RILUZOLE
 SULAR, NISOLDIPINE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 ZETONNA, CICLESONIDE

CPPI CV

* CP PHARMACEUTICALS INTERNATIONAL CV
 SUTENT, SUNITINIB MALATE

CREEKWOOD PHARMS

* CREEKWOOD PHARMACEUTICALS LLC
 FENOFIBRATE, FENOFIBRATE
 PREGABALIN, PREGABALIN
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILODOSIN, SILODOSIN

CROSSMEDIKA SA

* CROSSMEDIKA SA
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

CROWN LABS

* CROWN LABORATORIES INC
 ALA-CORT, HYDROCORTISONE
 TRIDERM, TRIAMCINOLONE ACETONIDE

CROWN LABS INC

* CROWN LABORATORIES INC
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN

CSPC OUYI

* CSPC OUYI PHARMACEUTICAL CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 CARBAMAZEPINE, CARBAMAZEPINE
 CELECOXIB, CELECOXIB
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 GABAPENTIN, GABAPENTIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

CSPC OUYI PHARM CO

* CSPC OUYI PHARMACEUTICAL CO LTD
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

CSPC-NBP PHARM

* CSPC-NBP PHARMACEUTICAL CO LTD
 BENZONATATE, BENZONATATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

CTI BIOPHARMA CORP

* CTI BIOPHARMA CORP
 VONJO, PACRITINIB CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CUBIST PHARMS LLC**

* CUBIST PHARMACEUTICALS LLC
 DIFICID, FIDAXOMICIN
 SIVEXTRO, TEDIZOLID PHOSPHATE
 ZERBAXA, CEFTOLOZANE SULFATE

CUMBERLAND

* CUMBERLAND PHARMACEUTICALS INC
 SANCUSO, GRANISETRON
 VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
 VIBATIV, TELAVANCIN HYDROCHLORIDE

CUMBERLAND PHARMS

* CUMBERLAND PHARMACEUTICALS INC
 ACETADOTE, ACETYLCYSTEINE
 CALDOLOR, IBUPROFEN
 LACTULOSE, LACTULOSE

CURIUM

* CURIUM US LLC
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
 IOFLUPANE I-123, IOFLUPANE I-123
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
 TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
 TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 XENON XE 133, XENON XE-133

CURRAX

* CURRAX PHARMACEUTICALS LLC
 ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
 SILENOR, DOXEPIN HYDROCHLORIDE
 TREXIMET, NAPROXEN SODIUM

CUSTOPHARM INC

* CUSTOPHARM INC
 CALCITONIN-SALMON, CALCITONIN SALMON

CYCLE

* CYCLE PHARMACEUTICALS LTD
 NITYR, NITISINONE
 TASCENSO ODT, FINGOLIMOD LAURYL SULFATE

CYCLOMEDICA

* CYCLOMEDICA AUSTRALIA PTY LTD
 TECHNEGAS KIT, TECHNETIUM TC-99M LABELED CARBON

**** D ******DAEWOONG PHARM CO**

* DAEWOONG PHARMACEUTICAL CO LTD
 MEROPENEM, MEROPENEM

DAIICHI SANKYO INC

* DAIICHI SANKYO INC
 SAVAYSA, EDOXABAN TOSYLATE
 TURALIO, PEXIDARTINIB HYDROCHLORIDE
 VANFLYTA, QUIZARTINIB DIHYDROCHLORIDE

DAITO

* DAITO PHARMACEUTICAL CO LTD
 METHOTREXATE SODIUM, METHOTREXATE SODIUM

DANCO LABS LLC

* DANCO LABORATORIES LLC
 MIFEPREX, MIFEPRISTONE

DARE

* DARE BIOSCIENCE INC
 XACIATO, CLINDAMYCIN PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******DASH PHARMS**

* DASH PHARMACEUTICALS LLC
TESTOSTERONE, TESTOSTERONE

DASH PHARMS NATCO

* DASH PHARMACEUTICALS LLC A FULLY OWNED SUB OF NATCO PHARMA LTD
ALBUTEROL SULFATE, ALBUTEROL SULFATE

DAVA PHARMS INC

* DAVA PHARMACEUTICALS INC
AMOXICILLIN, AMOXICILLIN
MORPHINE SULFATE, MORPHINE SULFATE
PROPYLTHIOURACIL, PROPYLTHIOURACIL

DAVIS AND GECK

* DAVIS AND GECK DIV AMERICAN CYANAMID CO
PRE-OP II, HEXACHLOROPHENE
PRE-OP, HEXACHLOROPHENE

DECATUR

* DECATUR MEMORIAL HOSP
AMMONIA N 13, AMMONIA N-13
CHOLINE C-11, CHOLINE C-11
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

DECIPHERA PHARMS

* DECIPHERA PHARMACEUTICALS LLC
QINLOCK, RIPRETINIB

DEL CATH SYSTEMS INC

* DELCATH SYSTEMS INC
HEPZATO, MELPHALAN HYDROCHLORIDE

DENTSPLY PHARM

* DENTSPLY PHARMACEUTICAL INC
CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
ORAQIX, LIDOCAINE

DEPROCO

* DEPROCO INC
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
SEPTOCAINE, ARTICAIN HYDROCHLORIDE

DERMAVANT SCI

* DERMAVANT SCIENCES INC
VTAMA, TAPINAROF

DEVA HOLDING AS

* DEVA HOLDING AS
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TEMOZOLOMIDE, TEMOZOLOMIDE

DEXCEL

* DEXCEL PHARMA TECHNOLOGIES LTD
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
HEMADY, DEXAMETHASONE
LANSOPRAZOLE, LANSOPRAZOLE (OTC)
OMEPRAZOLE, OMEPRAZOLE (OTC)
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VIGABATRIN, VIGABATRIN

DEXCEL LTD

* DEXCEL LTD
DICLOFENAC SODIUM, DICLOFENAC SODIUM
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

DEXCEL PHARMA

* DEXCEL PHARMA TECHNOLOGIES LTD
OMEPRAZOLE, OMEPRAZOLE (OTC)

DIALYSIS SUPS

* DIALYSIS SUPPLIES INC
NORMOCARB HF 25, MAGNESIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DIALYSIS SUPPLIES INC
NORMOCARB HF 35, MAGNESIUM CHLORIDE

DORC

* DORC INTERNATIONAL BV
VISIONBLUE, TRYPAN BLUE

DOW PHARM

* DOW PHARMACEUTICAL SCIENCES
ALTRENO, TRETINOIN
ATRALIN, TRETINOIN

DR REDDYS

* DR REDDYS LABORATORIES INC
FENOFIBRATE, FENOFIBRATE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
ICOSAPENT ETHYL, ICOSAPENT ETHYL
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
NITROGLYCERIN, NITROGLYCERIN
PROGESTERONE, PROGESTERONE
PROPOFOL, PROPOFOL
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* DR REDDYS LABORATORIES LTD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ALBENDAZOLE, ALBENDAZOLE
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
AZACITIDINE, AZACITIDINE
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
BIVALIRUDIN, BIVALIRUDIN
BORTEZOMIB, BORTEZOMIB
CABAZITAXEL, CABAZITAXEL
CAPECITABINE, CAPECITABINE
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
CARFILZOMIB, CARFILZOMIB
CARMUSTINE, CARMUSTINE
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN
CLOFARABINE, CLOFARABINE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
COLCHICINE, COLCHICINE
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DAPTOMYCIN, DAPTOMYCIN
DARUNAVIR, DARUNAVIR
DECITABINE, DECITABINE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DIFLUPREDNATE, DIFLUPREDNATE
DOCETAXEL, DOCETAXEL
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
ENALAPRILAT, ENALAPRILAT
EPHEDRINE SULFATE, EPHEDRINE SULFATE
ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
ESZOPICLONE, ESZOPICLONE
FEBUXOSTAT, FEBUXOSTAT
FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
FULVESTRANT, FULVESTRANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES LTD
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 IMATINIB MESYLATE, IMATINIB MESYLATE
 LANSOPRAZOLE, LANSOPRAZOLE
 LENALIDOMIDE, LENALIDOMIDE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LIDOCAINE, LIDOCAINE
 LUBIPROSTONE, LUBIPROSTONE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NELARABINE, NELARABINE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PENICILLAMINE, PENICILLAMINE
 PHYTONADIONE, PHYTONADIONE
 PLERIXAFOR, PLERIXAFOR
 POSACONAZOLE, POSACONAZOLE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREGABALIN, PREGABALIN
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 REGADENOSON, REGADENOSON
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIROLIMUS, SIROLIMUS
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUNITINIB MALATE, SUNITINIB MALATE
 TADALAFIL, TADALAFIL
 TETRABENAZINE, TETRABENAZINE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOPIRAMATE, TOPIRAMATE
 TREPROSTINIL, TREPROSTINIL
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN
 VINORELBINE TARTRATE, VINORELBINE TARTRATE

DR REDDYS LA

* DR REDDYS LABORATORIES LOUISIANA LLC
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LOPURIN, ALLOPURINOL
 SSD, SILVER SULFADIAZINE

DR REDDYS LABS EU

* DR REDDYS LABORATORIES EU LTD
 PERMETHRIN, PERMETHRIN

DR REDDYS LABS INC

* DR REDDYS LABORATORIES INC
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

- * DR REDDYS LABORATORIES INC
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LEVOFLOXACIN, LEVOFLOXACIN
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SIMVASTATIN, SIMVASTATIN
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
- DR REDDYS LABS LTD**
- * DR REDDYS LABORATORIES LIMITED
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- * DR REDDYS LABORATORIES LTD
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******* DR REDDYS LABORATORIES LTD**

NATEGLINIDE, NATEGLINIDE
 NIZATIDINE, NIZATIDINE
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 TACROLIMUS, TACROLIMUS
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZAFIRLUKAST, ZAFIRLUKAST
 ZENATANE, ISOTRETINOIN
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

DR REDDYS LABS SA*** DR REDDYS LABORATORIES SA**

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BIORPHEN, PHENYLEPHRINE HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CAMILA, NORETHINDRONE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CHLORZOXAZONE, CHLORZOXAZONE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLONIDINE, CLONIDINE
 CYCLOSPORINE, CYCLOSPORINE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOFETILIDE, DOFETILIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ERRIN, NORETHINDRONE
 ERYC, ERYTHROMYCIN
 ESTAZOLAM, ESTAZOLAM
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FLUOROURACIL, FLUOROURACIL
 HABITROL, NICOTINE (OTC)
 HALOETTE, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 MERCAPTOPYRIMIDINE, MERCAPTOPYRIMIDINE
 METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 RAMELTEON, RAMELTEON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES SA
 REZIPRES, EPHEDRINE HYDROCHLORIDE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TOBRAMYCIN, TOBRAMYCIN
 TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIVORA-28, ETHINYL ESTRADIOL

DUCHESNAY

* DUCHESNAY INC
 BONJESTA, DOXYLAMINE SUCCINATE
 DICLEGIS, DOXYLAMINE SUCCINATE
 OSPHENA, OSPEMIFENE

DURAMED PHARMS BARR

* DURAMED PHARMACEUTICALS INC SUB BARR LABORATORIES INC
 AVIANE-28, ETHINYL ESTRADIOL
 CRYSELLE, ETHINYL ESTRADIOL
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 ENPRESSE-28, ETHINYL ESTRADIOL
 VELIVET, DESOGESTREL

DUSA

* DUSA PHARMACEUTICALS INC
 LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE

DUTCH OPHTHALMIC

* DUTCH OPHTHALMIC RESEARCH CENTER INTERNATIONAL BV
 TISSUEBLUE, BRILLIANT BLUE G

REDDYS

* DOCTOR REDDYS LABORATORIES LTD
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

**** E ******E5 PHARMA INC**

* E5 PHARMA INC
 DIAZOXIDE, DIAZOXIDE

EAGLE PHARMS

* EAGLE PHARMACEUTICALS INC
 BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 PEMFEXY, PEMETREXED
 RYANODEX, DANTROLENE SODIUM
 VASOPRESSIN, VASOPRESSIN

ECI PHARMS LLC

* ECI PHARMACEUTICALS LLC
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

ECOLAB

* ECOLAB INC
 CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

EDENBRIDGE PHARMS

* EDENBRIDGE PHARMACEUTICALS LLC
 ALBENDAZOLE, ALBENDAZOLE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 CARBIDOPA, CARBIDOPA
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 IVERMECTIN, IVERMECTIN
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TINIDAZOLE, TINIDAZOLE
 YARGESA, MIGLUSTAT

EDISON THERAPS LLC

* EDISON THERAPEUTICS LLC
 METHERGINE, METHYLERGONOVINE MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******EIGER BIOPHARMS**

* EIGER BIOPHARMACEUTICALS INC
ZOKINVY, LONAFARNIB

EIRGEN

* EIRGEN PHARMA LTD
RAYALDEE, CALCIFEDIOL

EISAI INC

* EISAI INC
ARICEPT, DONEPEZIL HYDROCHLORIDE
BANZEL, RUFINAMIDE
DAYVIGO, LEMBOREXANT
HALAVEN, ERIBULIN MESYLATE
LENVIMA, LENVATINIB MESYLATE

ELI LILLY AND CO

* ELI LILLY AND CO
MOUNJARO, TIRZEPATIDE
OLUMIANT, BARICITINIB
PROZAC, FLUOXETINE HYDROCHLORIDE
REYVOW, LASMIDITAN SUCCINATE
VERZENIO, ABEMACICLIB
ZEPBOUND, TIRZEPATIDE

ELI LILLY CO

* ELI LILLY CO
ADCIRCA, TADALAFIL

ELITE LABS

* ELITE LABORATORIES INC
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

ELITE LABS INC

* ELITE LABORATORIES INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
DANTROLENE SODIUM, DANTROLENE SODIUM
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
ISRADIPINE, ISRADIPINE
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

ELITE PHARM SOLUTION

* ELITE PHARMACEUTICAL SOLUTION INC
NIFEDIPINE, NIFEDIPINE

ELYSIUM

* ELYSIUM PHARMACEUTICALS LTD
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE

EMD SERONO INC

* EMD SERONO INC
CETROTIDE, CETRORELIX ACETATE
MAVENCLAD, CLADRIBINE
TEPMETKO, TEPOTINIB HYDROCHLORIDE

EMED MEDCL

* EMED MEDICAL CO LLC
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

EMERGENT

* EMERGENT OPERATIONS IRELAND LTD
NARCAN, NALOXONE HYDROCHLORIDE (OTC)

EMERGENT BIODEFENSE

* EMERGENT BIODEFENSE OPERATIONS LANSING INC
TEMBEXA, BRINCIDOFVIR
* EMERGENT BIODEFENSE OPERATIONS LANSING LLC
TEMBEXA, BRINCIDOFVIR

EMMAUS MEDCL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EMMAUS MEDICAL INC
ENDARI, L-GLUTAMINE

ENCUBE

* ENCUBE ETHICALS PRIVATE LTD
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
AZELAIC ACID, AZELAIC ACID
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DAPSONE, DAPSONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
DOCOSANOL, DOCOSANOL (OTC)
FLUOROURACIL, FLUOROURACIL
KETOCONAZOLE, KETOCONAZOLE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
MESALAMINE, MESALAMINE
METRONIDAZOLE, METRONIDAZOLE
MUPIROCIN, MUPIROCIN CALCIUM
TACROLIMUS, TACROLIMUS
TAVABOROLE, TAVABOROLE
TESTOSTERONE, TESTOSTERONE
TRETINOIN MICROSPHERE, TRETINOIN
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ENCUBE ETHICALS

* ENCUBE ETHICALS PVT LTD
ADAPALENE, ADAPALENE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
DESONIDE, DESONIDE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
PERMETHRIN, PERMETHRIN

ENDO PHARM

* ENDO PHARMACEUTICAL SOLUTIONS INC
SUPPRELIN LA, HISTRELIN ACETATE
VALSTAR PRESERVATIVE FREE, VALRUBICIN

ENDO PHARMS

* ENDO PHARMACEUTICALS INC
FORTESTA, TESTOSTERONE
FROVA, FROVATRIPTAN SUCCINATE
PERCODAN, ASPIRIN

ENDO PHARMS INC

* ENDO PHARMACEUTICALS INC
AVEED, TESTOSTERONE UNDECANOATE
COLY-MYCIN S, COLISTIN SULFATE
MEGACE ES, MEGESTROL ACETATE
NASCOBAL, CYANOCOBALAMIN

ENTASIS THERAP

* ENTASIS THERAPEUTICS INC
XACDURO (COPACKAGED), DURLOBACTAM SODIUM

EPI HLTH

* EPI HEALTH LLC
CLODERM, CLOCORTOLONE PIVALATE
MINOLIRA, MINOCYCLINE HYDROCHLORIDE

EPIC PHARMA

* EPIC PHARMA LLC
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
SULINDAC, SULINDAC
TRANDOLAPRIL, TRANDOLAPRIL
URSODIOL, URSODIOL

EPIC PHARMA LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******* EPIC PHARMA LLC**

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CILOSTAZOL, CILOSTAZOL
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DESOXIMETASONE, DESOXIMETASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ERYTHROMYCIN, ERYTHROMYCIN
 ESTRADIOL, ESTRADIOL
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 FUROSEMIDE, FUROSEMIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE, GLYBURIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LOVASTATIN, LOVASTATIN
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NIZATIDINE, NIZATIDINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREGABALIN, PREGABALIN
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 RIFAMPIN, RIFAMPIN
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SULFADIAZINE, SULFADIAZINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROPICACYL, TROPICAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EPIC PHARMA LLC
 VORICONAZOLE, VORICONAZOLE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

EPIZYME INC

* EPIZYME INC
 TAZVERIK, TAZEMETOSTAT HYDROBROMIDE

ESKAYEF

* ESKAYEF PHARMACEUTICALS LTD
 PREGABALIN, PREGABALIN

ESPERION THERAPS INC

* ESPERION THERAPEUTICS INC
 NEXLETOL, BEMPEDOIC ACID
 NEXLIZET, BEMPEDOIC ACID

ESSENTIAL ISOTOPES

* ESSENTIAL ISOTOPES LLC
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

ETHYPHARM

* ETHYPHARM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 * ETHYPHARM SA
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM

ETHYPHARM USA CORP

* ETHYPHARM USA CORP
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

ETON

* ETON PHARMACEUTICALS INC
 ALKINDI SPRINKLE, HYDROCORTISONE
 BETAINE, BETAINE
 NITISINONE, NITISINONE

EUGIA PHARMA

* EUGIA PHARMA SPECIALITIES LTD
 ACETAMINOPHEN, ACETAMINOPHEN
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ADENOSINE, ADENOSINE
 AMPHOTERICIN B, AMPHOTERICIN B
 ANASTROZOLE, ANASTROZOLE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZACITIDINE, AZACITIDINE
 AZITHROMYCIN, AZITHROMYCIN
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BIMATOPROST, BIMATOPROST
 BIVALIRUDIN, BIVALIRUDIN
 BORTEZOMIB, BORTEZOMIB
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUDESONIDE, BUDESONIDE
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CAPECITABINE, CAPECITABINE
 CARBOPLATIN, CARBOPLATIN
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DACTINOMYCIN, DACTINOMYCIN
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EUGIA PHARMA SPECIALITIES LTD
 DOCETAXEL, DOCETAXEL
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPTIFIBATIDE, EPTIFIBATIDE
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ETOMIDATE, ETOMIDATE
 EXEMESTANE, EXEMESTANE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 LENALIDOMIDE, LENALIDOMIDE
 LETROZOLE, LETROZOLE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIFITEGRAST, LIFITEGRAST
 LINEZOLID, LINEZOLID
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MEROPENEM, MEROPENEM
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 OXALIPLATIN, OXALIPLATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PLERIXAFOR, PLERIXAFOR
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POSACONAZOLE, POSACONAZOLE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROGESTERONE, PROGESTERONE
 REGADENOSON, REGADENOSON
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EUGIA PHARMA SPECIALITIES LTD
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VASOPRESSIN, VASOPRESSIN
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

EUGIA PHARMA SPECLTS

* EUGIA PHARMA SPECIALITIES LTD
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 OXACILLIN SODIUM, OXACILLIN SODIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

EUROCEPT PHARMS

* EUROCEPT PHARMACEUTICALS
 PASER, AMINOSALICYLIC ACID

EUROHLTH INTL SARL

* EUROHEALTH INTERNATIONAL SARL
 AZACITIDINE, AZACITIDINE

EVEREST LIFE SCI

* EVEREST LIFE SCIENCES LLC
 DAPSONE, DAPSONE

EVOFEM INC

* EVOFEM INC
 PHEXXI, CITRIC ACID

EVOKE PHARMA INC

* EVOKE PHARMA INC
 GIMOTI, METOCLOPRAMIDE HYDROCHLORIDE

EVUS

* EVUS HEALTH SOLUTIONS LLC
 NITROMIST, NITROGLYCERIN

EXELA PHARMA

* EXELA PHARMA SCIENCES LLC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 AKOVAZ, EPHEDRINE SULFATE
 BLOXIVERZ, NEOSTIGMINE METHYLSULFATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CUPRIC CHLORIDE, CUPRIC CHLORIDE
 ELCYS, CYSTEINE HYDROCHLORIDE
 GANZYK-RTU, GANCICLOVIR
 GLYRX-PF, GLYCOPYRROLATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MIDAZOLAM IN 0.8% SODIUM CHLORIDE, MIDAZOLAM
 NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
 POTASSIUM ACETATE, POTASSIUM ACETATE
 SODIUM BICARBONATE, SODIUM BICARBONATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZINC CHLORIDE, ZINC CHLORIDE

EXELA PHARMA SCIENCE

* EXELA PHARMA SCIENCES
 CAFFEINE CITRATE, CAFFEINE CITRATE

EXELIXIS

* EXELIXIS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EXELIXIS INC
COMETRIQ, CABOZANTINIB S-MALATE

EXELIXIS INC

* EXELIXIS INC
CABOMETYX, CABOZANTINIB S-MALATE

EXELTIS USA INC

* EXELTIS USA INC
SLYND, DROSPIRENONE
TYBLUME, ETHINYL ESTRADIOL

EYENOVIA

* EYENOVIA INC
MYDCOMBI, PHENYLEPHRINE HYDROCHLORIDE

EYEPOINT PHARMS

* EYEPOINT PHARMACEUTICALS INC
DEXYCU KIT, DEXAMETHASONE

EYWA

* EYWA PHARMA INC
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
NIACIN, NIACIN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
POTASSIUM CITRATE, POTASSIUM CITRATE
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
VALPROIC ACID, VALPROIC ACID

EYWA PHARMA

* EYWA PHARMA PTE LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
BACLOFEN, BACLOFEN
URSODIOL, URSODIOL

EZRA VENTURES

* EZRA VENTURES LLC
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE

LILLY

* ELI LILLY AND CO
ALIMTA, PEMETREXED DISODIUM
CIALIS, TADALAFIL
CYMBALTA, DULOXETINE HYDROCHLORIDE
EVISTA, RALOXIFENE HYDROCHLORIDE
FORTEO, TERIPARATIDE
STRATTERA, ATOMOXETINE HYDROCHLORIDE
SYMBYAX, FLUOXETINE HYDROCHLORIDE

**** F ******FABRE KRAMER**

* FABRE KRAMER PHARMACEUTICALS INC
EXXUA, GEPIRONE HYDROCHLORIDE

FDC LTD

* FDC LTD
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
LATANOPROST, LATANOPROST
OFLOXACIN, OFLOXACIN
TIMOLOL MALEATE, TIMOLOL MALEATE

FDN CONSUMER

* FOUNDATION CONSUMER BRANDS LLC
ALAVERT, LORATADINE (OTC)
* FOUNDATION CONSUMER HEALTHCARE LLC
PLAN B ONE-STEP, LEVONORGESTREL (OTC)

FEINSTEIN

* FEINSTEIN INSTITUTE MEDICAL RESEARCH
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
FLUORODOPA F18, FLUORODOPA F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ******FENNEC PHARMS INC**

* FENNEC PHARMACEUTICALS INC
PEDMARK, SODIUM THIOSULFATE

FERA PHARMS LLC

* FERA PHARMACEUTICALS LLC
GENTAMICIN SULFATE, GENTAMICIN SULFATE
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE

FERRER INTERNACIONAL

* FERRER INTERNACIONAL SA
XEPI, OZENOXACIN

FERRING

* FERRING PHARMACEUTICALS INC
ENDOMETRIN, PROGESTERONE
FIRMAGON, DEGARELIX ACETATE

FERRING PHARMS INC

* FERRING PHARMACEUTICALS INC
CERVIDIL, DINOPROSTONE
CLENPIQ, CITRIC ACID
DDAVP, DESMOPRESSIN ACETATE

FLAMINGO PHARMS

* FLAMINGO PHARMACEUTICALS LTD
METRONIDAZOLE, METRONIDAZOLE
PIROXICAM, PIROXICAM

FLORIDA

* FLORIDA PHARMACEUTICAL PRODUCTS LLC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
SUMATRIPTAN, SUMATRIPTAN

FOLDRX PHARMS

* FOLDRX PHARMACEUTICALS LLC A WHOLLY OWNED SUB OF PFIZER INC
VYNDAMAX, TAFAMIDIS
VYNDAQEL, TAFAMIDIS MEGLUMINE

FOSUN PHARMA

* FOSUN PHARMA USA INC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

FOUGERA PHARMS

* FOUGERA PHARMACEUTICALS INC
ADAPALENE, ADAPALENE
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CALCIPOTRIENE, CALCIPOTRIENE
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOTRIMAZOLE, CLOTRIMAZOLE
DESONIDE, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HYDROCORTISONE, HYDROCORTISONE
IMIQUIMOD, IMIQUIMOD
KETOCONAZOLE, KETOCONAZOLE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
METRONIDAZOLE, METRONIDAZOLE
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN
NYSTATIN, NYSTATIN
OXISTAT, OXICONAZOLE NITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FOUGERA PHARMACEUTICALS INC
 PREDNICARBATE, PREDNICARBATE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

FOUGERA PHARMS INC

* FOUGERA PHARMACEUTICALS INC
 ACYCLOVIR, ACYCLOVIR
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCORTISONE, HYDROCORTISONE
 LIDOCAINE, LIDOCAINE
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 TACROLIMUS, TACROLIMUS
 TERCONAZOLE, TERCONAZOLE

FRESENIUS

* FRESENIUS KABI DEUTSCHLAND GMBH
 INTRALIPID 20%, SOYBEAN OIL
 INTRALIPID 30%, SOYBEAN OIL
 * FRESENIUS KABI IPSUM SRL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

FRESENIUS KABI

* FRESENIUS KABI ANTI INFECTIVES SRL
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 * FRESENIUS KABI AUSTRIA GMBH
 LACTULOSE, LACTULOSE

FRESENIUS KABI USA

* FRESENIUS KABI USA LLC
 ACETAMINOPHEN, ACETAMINOPHEN
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ADENOSINE, ADENOSINE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 AZTREONAM, AZTREONAM
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BIVALIRUDIN, BIVALIRUDIN
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BORTEZOMIB, BORTEZOMIB
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 CARBOPLATIN, CARBOPLATIN
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CEFOTETAN, CEFOTETAN DISODIUM
 CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CLADRIBINE, CLADRIBINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** F **

* FRESENIUS KABI USA LLC
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAPTOMYCIN, DAPTOMYCIN
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075%, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.225%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DIAZEPAM, DIAZEPAM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILAUDID, HYDROMORPHONE HYDROCHLORIDE
 DIMENHYDRINATE, DIMENHYDRINATE
 DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPRIVAN, PROPOFOL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXY 100, DOXYCYCLINE HYCLATE
 DOXY 200, DOXYCYCLINE HYCLATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FULVESTRANT, FULVESTRANT
 FUROSEMIDE, FUROSEMIDE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GLUCAGON, GLUCAGON HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 IFOSFAMIDE, IFOSFAMIDE
 INDOMETHACIN, INDOMETHACIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MANNITOL 25%, MANNITOL
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MESNA, MESNA
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 5.5, MAGNESIUM CHLORIDE
 MULTIPLE ELECTROLYTES INJECTION TYPE 1 USP PH 7.4, MAGNESIUM CHLORIDE
 NAROPIN, ROPIVACAINE HYDROCHLORIDE
 NEBUPENT, PENTAMIDINE ISETHIONATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
 NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OMEGAVEN, FISH OIL TRIGLYCERIDES
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PENTAM, PENTAMIDINE ISETHIONATE
 PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 POLOCAINE, MEPIVACAINE HYDROCHLORIDE
 POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POSACONAZOLE, POSACONAZOLE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45%, POTASSIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ******* FRESENIUS KABI USA LLC**

POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 10MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9%, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM PHOSPHATES, POTASSIUM PHOSPHATE, DIBASIC
 PROGESTERONE, PROGESTERONE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROMIDEPSIN, ROMIDEPSIN
 SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 SMOFLIPID 20%, FISH OIL
 SODIUM ACETATE, SODIUM ACETATE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM PHOSPHATES, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VALPROATE SODIUM, VALPROATE SODIUM
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VASOPRESSIN, VASOPRESSIN
 VIBISONE, CYANOCOBALAMIN
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 ZINC SULFATE, ZINC SULFATE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

FRESENIUS MEDCL*** FRESENIUS MEDICAL CARE NORTH AMERICA**

DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

- * FRESENIUS MEDICAL CARE NORTH AMERICA
PHOSLO GELCAPS, CALCIUM ACETATE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

**** G ******G1 THERAP**

- * G1 THERAPEUTICS INC
COSELA, TRILACICLIB DIHYDROCHLORIDE

GALDERMA LABS

- * GALDERMA LABORATORIES INC
CLOBEX, CLOBETASOL PROPIONATE
EPIDUO FORTE, ADAPALENE

GALDERMA LABS LP

- * GALDERMA LABORATORIES L P
CLOBEX, CLOBETASOL PROPIONATE
- * GALDERMA LABORATORIES LP
AKLIEF, TRIFAROTENE
CAPEX, FLUOCINOLONE ACETONIDE
CLOBEX, CLOBETASOL PROPIONATE
DESOWEN, DESONIDE
DIFFERIN, ADAPALENE
DIFFERIN, ADAPALENE (OTC)
EPIDUO, ADAPALENE
EPSOLAY, BENZOYL PEROXIDE
METROCREAM, METRONIDAZOLE
METROGEL, METRONIDAZOLE
METROLOTION, METRONIDAZOLE
MIRVASO, BRIMONIDINE TARTRATE
ORACEA, DOXYCYCLINE
SOOLANTRA, IVERMECTIN
TRI-LUMA, FLUOCINOLONE ACETONIDE
TWYNEO, BENZOYL PEROXIDE
VECTICAL, CALCITRIOL

GALT PHARMS

- * GALT PHARMACEUTICALS LLC
DORAL, QUAZEPAM
ORAVIG, MICONAZOLE
ORPHENGESIC FORTE, ASPIRIN
ORPHENGESIC, ASPIRIN

GE HEALTHCARE

- * GE HEALTHCARE
ADREVIEW, IOBENGUANE SULFATE I-123
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CERIANNA, FLUROESTRADIOL F-18
CLARISCAN, GADOTERATE MEGLUMINE
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
MYOVIEW, TECHNETIUM TC-99M TETROFOSMIN KIT
OMNIPAQUE 12, IOHEXOL
OMNIPAQUE 140, IOHEXOL
OMNIPAQUE 180, IOHEXOL
OMNIPAQUE 240, IOHEXOL
OMNIPAQUE 300, IOHEXOL
OMNIPAQUE 350, IOHEXOL
OMNIPAQUE 9, IOHEXOL
OMNISCAN, GADODIAMIDE
OPTISON, ALBUMIN HUMAN
VISIPAQUE 270, IODIXANOL
VISIPAQUE 320, IODIXANOL
VIZAMYL, FLUTEMETAMOL F-18

GE HLTHCARE INC

- * GE HEALTHCARE INC
DATSCAN, IOFLUPANE I-123

GENBIOPRO

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GENBIOPRO INC
MIFEPRISTONE, MIFEPRISTONE

GENENTECH

* GENENTECH INC
ERIVEDGE, VISMODEGIB

GENENTECH INC

* GENENTECH INC
COTELLIC, COBIMETINIB FUMARATE
ESBRIET, PIRFENIDONE
EVRYSDI, RISDIPLAM
GAVRETO, PRALSETINIB
ROZLYTREK, ENTRECTINIB
XOFLUZA, BALOXAVIR MARBOXIL

GENERIC

* GENERICS INTERNATIONAL VENTURES ENTERPRISES LLC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

GENEYORK PHARMS

* GENEYORK PHARMACEUTICALS GROUP LLC
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
PREDNISONE, PREDNISONE

GENSCO

* GENSCO LABORATORIES LLC DBA GENSCO PHARMA
RIZAFILM, RIZATRIPTAN BENZOATE

GENUS

* GENUS LIFESCIENCES INC
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
HYCODAN, HOMATROPINE METHYLBROMIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
NYSTATIN, NYSTATIN

GENUS LIFESCIENCES

* GENUS LIFE SCIENCES INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
AMCINONIDE, AMCINONIDE
GOPRELTO, COCAINE HYDROCHLORIDE
LEVOLET, LEVOTHYROXINE SODIUM **
MONOKET, ISOSORBIDE MONONITRATE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

GENZYME

* GENZYME CORP
CLOLAR, CLOFARABINE
MOZOBIL, PLERIXAFOR
RENAGEL, SEVELAMER HYDROCHLORIDE
RENVELA, SEVELAMER CARBONATE

GENZYME CORP

* GENZYME CORP
CAPRELSA, VANDETANIB
CERDELGA, ELIGLUSTAT TARTRATE

GILEAD

* GILEAD SCIENCES INC
CAYSTON, AZTREONAM
EMTRIVA, EMTRICITABINE
HEPSERA, ADEFOVIR DIPIVOXIL
LETAIRIS, AMBRISENTAN
TRUVADA, EMTRICITABINE

GILEAD SCIENCES

* GILEAD SCIENCES LLC
ATRIPLA, EFAVIRENZA

GILEAD SCIENCES INC

* GILEAD SCIENCES INC
BIKTARVY, BICTEGRAVIR SODIUM
COMPLERA, EMTRICITABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GILEAD SCIENCES INC
 DESCOVY, EMTRICITABINE
 EPCLUSA, SOFOSBUVIR
 GENVOYA, COBICISTAT
 HARVONI, LEDIPASVIR
 ODEFSEY, EMTRICITABINE
 SOVALDI, SOFOSBUVIR
 STRIBILD, COBICISTAT
 SUNLENCA, LENACAPAVIR SODIUM
 TYBOST, COBICISTAT
 VEKLURY, REMDESIVIR
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 VOSEVI, SOFOSBUVIR
 ZYDELIG, IDELALISIB

GISKIT

* GISKIT PHARMA BV
 EXEM FOAM KIT, AIR POLYMER-TYPE A

GLAND

* GLAND PHARMA LTD
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 MEROPENEM, MEROPENEM
 MESNA, MESNA
 OXALIPLATIN, OXALIPLATIN
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

GLAND PHARMA LTD

* GLAND PHARMA LTD
 ADENOSINE, ADENOSINE
 ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARGATROBAN, ARGATROBAN
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BIMATOPROST, BIMATOPROST
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUMETANIDE, BUMETANIDE
 CALCITRIOL, CALCITRIOL
 CARBOPLATIN, CARBOPLATIN
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DOCETAXEL, DOCETAXEL
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLAND PHARMA LTD
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
EPHEDRINE SULFATE, EPHEDRINE SULFATE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
ETOMIDATE, ETOMIDATE
FLUOROURACIL, FLUOROURACIL
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FOMEPIZOLE, FOMEPIZOLE
FOSCARNET SODIUM, FOSCARNET SODIUM
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
FUROSEMIDE, FUROSEMIDE
GANIRELIX ACETATE, GANIRELIX ACETATE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LACOSAMIDE, LACOSAMIDE
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
LEVOFLOXACIN, LEVOFLOXACIN
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MITOMYCIN, MITOMYCIN
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
NELARABINE, NELARABINE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
PACLITAXEL, PACLITAXEL
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
POSACONAZOLE, POSACONAZOLE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
REGADENOSON, REGADENOSON
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TEMSIROLIMUS, TEMSIROLIMUS
THIOTEPA, THIOTEPA
TIMOLOL MALEATE, TIMOLOL MALEATE
TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE
TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TOBRAMYCIN, TOBRAMYCIN
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VORICONAZOLE, VORICONAZOLE
ZINC SULFATE, ZINC SULFATE
ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE
ZOLEDRONIC ACID, ZOLEDRONIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLAND PHARMA LTD
 ZOLEDRONIC, ZOLEDRONIC ACID

GLAUKOS

* GLAUKOS CORP
 IDOSE TR, TRAVOPROST
 PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE
 PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE

GLAXO GRP ENGLAND

* GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
 INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE

GLAXO GRP LTD

* GLAXO GROUP LTD DBA GLAXOSMITHKLINE
 FLOVENT HFA, FLUTICASONE PROPIONATE

* GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
 ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
 ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
 ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
 ADVAIR HFA, FLUTICASONE PROPIONATE
 BREO ELLIPTA, FLUTICASONE FUROATE
 FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 50, FLUTICASONE PROPIONATE

GLAXOSMITHKLINE

* GLAXOSMITHKLINE
 ABREVA, DOCOSANOL (OTC)
 EPIVIR-HBV, LAMIVUDINE
 IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
 IMITREX, SUMATRIPTAN
 IMITREX, SUMATRIPTAN SUCCINATE
 MALARONE PEDIATRIC, ATOVAQUONE
 MALARONE, ATOVAQUONE
 NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
 NICORETTE, NICOTINE POLACRILEX (OTC)
 RELENZA, ZANAMIVIR
 VALTREX, VALACYCLOVIR HYDROCHLORIDE
 VERAMYST, FLUTICASONE FUROATE
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE

* GLAXOSMITHKLINE CONSUMER HEALTHCARE HOLDINGS US LLC
 ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)
 ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
 ADVIL COLD AND SINUS, IBUPROFEN (OTC)
 ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)
 ADVIL DUAL ACTION WITH ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ADVIL LIQUI-GELS, IBUPROFEN (OTC)
 ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)
 ADVIL MULTI-SYMPTOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)
 ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
 ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 ADVIL, IBUPROFEN (OTC)
 ADVIL, IBUPROFEN SODIUM (OTC)
 AXID AR, NIZATIDINE (OTC)
 CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
 CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
 CHILDREN'S ADVIL, IBUPROFEN (OTC)
 CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
 INFANT'S ADVIL, IBUPROFEN (OTC)
 JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
 NICORETTE, NICOTINE POLACRILEX (OTC)

* GLAXOSMITHKLINE INTELLECTUAL PROPERTY DEVELOPMENT LTD ENGLAND
 ANORO ELLIPTA, UMECLIDINIUM BROMIDE
 ARNUITY ELLIPTA, FLUTICASONE FUROATE
 KRINTAFEL, TAFENOQUINE SUCCINATE
 TRELEGY ELLIPTA, FLUTICASONE FUROATE

* GLAXOSMITHKLINE INTELLECTUAL PROPERTY LTD ENGLAND

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLAXOSMITHKLINE INTELLECTUAL PROPERTY LTD ENGLAND
SEREVENT, SALMETEROL XINAFOATE
VENTOLIN HFA, ALBUTEROL SULFATE
- * GLAXOSMITHKLINE INTELLECTUAL PROPERTY NO 2 LTD ENGLAND
JESDUVROQ, DAPRODUSTAT
- * GLAXOSMITHKLINE LLC
OJJAARA, MOMELOTINIB DIHYDROCHLORIDE
ZEJULA, NIRAPARIB TOSYLATE

GLAXOSMITHKLINE CONS

- * GLAXOSMITHKLINE CONSUMER HEALTHCARE
ALLI, ORLISTAT (OTC)
EXCEDRIN (MIGRAINE RELIEF), ACETAMINOPHEN (OTC)
FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)
FLONASE SENSIMIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)
LAMISIL AT, TERBINAFINE (OTC)
LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
VOLTAREN ARTHRITIS PAIN, DICLOFENAC SODIUM (OTC)

GLAXOSMITHKLINE LLC

- * GLAXOSMITHKLINE LLC
FLOLAN, EPOPROSTENOL SODIUM
LAMICTAL CD, LAMOTRIGINE
LAMICTAL ODT, LAMOTRIGINE
LAMICTAL XR, LAMOTRIGINE
LAMICTAL, LAMOTRIGINE
MEPRON, ATOVAQUONE

GLENMARK GENERICS

- * GLENMARK GENERICS INC USA
ADAPALENE, ADAPALENE (OTC)
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
IMIQUIMOD, IMIQUIMOD
MOMETASONE FUROATE, MOMETASONE FUROATE
NIZATIDINE, NIZATIDINE
ZONISAMIDE, ZONISAMIDE
- * GLENMARK GENERICS LIMITED
BRIELLYN, ETHINYL ESTRADIOL
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- * GLENMARK GENERICS LTD
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
ALYACEN 1/35, ETHINYL ESTRADIOL
ALYACEN 7/7/7, ETHINYL ESTRADIOL
ASHLYNA, ETHINYL ESTRADIOL
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
CARVEDILOL, CARVEDILOL
CICLOPIROX, CICLOPIROX
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
DESOXIMETASONE, DESOXIMETASONE
ESZOPICLONE, ESZOPICLONE
FELODIPINE, FELODIPINE
FLUCONAZOLE, FLUCONAZOLE
FLUOCINONIDE, FLUOCINONIDE
HEATHER, NORETHINDRONE
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
LAMOTRIGINE, LAMOTRIGINE
LEVOFLOXACIN, LEVOFLOXACIN
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LITHIUM CARBONATE, LITHIUM CARBONATE
MARLISSA, ETHINYL ESTRADIOL
MELOXICAM, MELOXICAM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE
MONTELUKAST SODIUM, MONTELUKAST SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLENMARK GENERICS LTD
 - NAPROXEN, NAPROXEN
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - NORETHINDRONE, NORETHINDRONE
 - NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - OMEPRAZOLE, OMEPRAZOLE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON, ONDANSETRON
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 - THEOPHYLLINE, THEOPHYLLINE
 - TOPIRAMATE, TOPIRAMATE
 - TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
 - TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 - URSODIOL, URSODIOL
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 - VIORELE, DESOGESTREL
 - ZOLMITRIPTAN, ZOLMITRIPTAN
- * GLENMARK GENERICS LTD INDIA
 - INDOMETHACIN, INDOMETHACIN
 - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
- GLENMARK PHARMS**
- * GLENMARK PHARMACEUTICALS INC
 - TERIFLUNOMIDE, TERIFLUNOMIDE
- * GLENMARK PHARMACEUTICALS INC USA
 - CICLOPIROX, CICLOPIROX
 - CLOTRIMAZOLE, CLOTRIMAZOLE
 - MUPIROCIN, MUPIROCIN
- * GLENMARK PHARMACEUTICALS LTD
 - MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
- * GLENMARK PHARMACEUTICALS SA
 - ABIRATERONE ACETATE, ABIRATERONE ACETATE
 - ATOVAQUONE, ATOVAQUONE
 - AZELAIC ACID, AZELAIC ACID
 - CALCIPOTRIENE, CALCIPOTRIENE
 - CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - DESONIDE, DESONIDE
 - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 - HAILEY 1.5/30, ETHINYL ESTRADIOL
 - HAILEY FE 1.5/30, ETHINYL ESTRADIOL
 - LINEZOLID, LINEZOLID
 - NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
- GLENMARK PHARMS INC**
- * GLENMARK PHARMACEUTICALS INC USA
 - CALCIPOTRIENE, CALCIPOTRIENE
 - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - FULVESTRANT, FULVESTRANT
 - LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 - LITHIUM CARBONATE, LITHIUM CARBONATE
 - MUPIROCIN, MUPIROCIN CALCIUM
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
- GLENMARK PHARMS LTD**
- * GLENMARK PHARMACEUTICALS LTD
 - ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 - AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLENMARK PHARMACEUTICALS LTD
 APREMILAST, APREMILAST
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESONIDE, DESONIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ESTRADIOL, ESTRADIOL
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUCINOLONE ACETONIDE, FLUCINOLONE ACETONIDE
 FLUCINONIDE ACETONIDE, FLUCINOLONE ACETONIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 GABAPENTIN, GABAPENTIN
 HAILEY 24 FE, ETHINYL ESTRADIOL
 HAILEY FE 1/20, ETHINYL ESTRADIOL
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 INDOMETHACIN, INDOMETHACIN
 LACOSAMIDE, LACOSAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LIDOCAINE, LIDOCAINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXCARBAZEPINE, OXCARBAZEPINE
 PIMECROLIMUS, PIMECROLIMUS
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 RUFINAMIDE, RUFINAMIDE
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SIROLIMUS, SIROLIMUS
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 TACROLIMUS, TACROLIMUS
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 THEOPHYLLINE, THEOPHYLLINE
 TOPIRAMATE, TOPIRAMATE
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLENMARK PHARMACEUTICALS LTD
VORICONAZOLE, VORICONAZOLE

GLENMARK PHARMS SA

* GLENMARK PHARMACEUTICALS SA SWITZERLAND
ACYCLOVIR, ACYCLOVIR
APREPITANT, APREPITANT
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
NITROGLYCERIN, NITROGLYCERIN

GLENMARK SPECIALTY

* GLENMARK SPECIALTY SA
RYALTRIS, MOMETASONE FUROATE

GLOBAL BLOOD THERAPIS

* GLOBAL BLOOD THERAPEUTICS INC
OXBRYTA, VOXELOTOR

GLW

* GLW PHARMA GMBH
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS

GRANULATION TECH

* GRANULATION TECHNOLOGY INC
PREDNISONE, PREDNISONE

GRANULES

* GRANULES INDIA LTD
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
GABAPENTIN, GABAPENTIN
GUAIFENESIN, GUAIFENESIN (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
LEVETIRACETAM, LEVETIRACETAM
LORATADINE, LORATADINE (OTC)
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
METOPROLOL SUCCINATE
NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
NAPROXEN, NAPROXEN
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
ZONISAMIDE, ZONISAMIDE

* GRANULES PHARMACEUTICALS INC
ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
COLCHICINE, COLCHICINE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DOFETILIDE, DOFETILIDE
METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
PENICILLAMINE, PENICILLAMINE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
RAMELTEON, RAMELTEON
SILDENAFIL CITRATE, SILDENAFIL CITRATE
TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

GRANULES INDIA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GRANULES INDIA LTD
IBUPROFEN, IBUPROFEN (OTC)
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

GRAVITI PHARMS

- * GRAVITI PHARMACEUTICALS INC
GABAPENTIN, GABAPENTIN
MONTELUKAST SODIUM, MONTELUKAST SODIUM
- * GRAVITI PHARMACEUTICALS PRIVATE LTD
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BACLOFEN, BACLOFEN
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CHLORZOXAZONE, CHLORZOXAZONE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
FAMOTIDINE, FAMOTIDINE
FENOFIBRATE, FENOFIBRATE
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
FUROSEMIDE, FUROSEMIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE

GUANGZHOU NOVAKEN

- * GUANGZHOU NOVAKEN PHARMACEUTICAL CO LTD
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

GUARDIAN DRUG

- * GUARDIAN DRUG CO
GUAIFENESIN, GUAIFENESIN (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
LORATADINE, LORATADINE (OTC)

GUERBET

- * GUERBET
ELUCIREM, GADOPICLENOL
- * GUERBET LLC
DOTAREM, GADOTERATE MEGLUMINE
LIPIODOL, ETHIODIZED OIL

**** H ******HAEMONETICS**

- * HAEMONETICS CORP
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

HAINAN POLY

- * HAINAN POLY PHARM CO LTD
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
AZITHROMYCIN, AZITHROMYCIN
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
FLUCONAZOLE, FLUCONAZOLE
IOPAMIDOL, IOPAMIDOL
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VORICONAZOLE, VORICONAZOLE

HAINAN POLY PHARM

- * HAINAN POLY PHARMACEUTICAL CO LTD
DAPTOMYCIN, DAPTOMYCIN
GADOBUTROL, GADOBUTROL
LEVETIRACETAM, LEVETIRACETAM
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

HALOCARBON PRODS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HALOCARBON PRODUCTS CORP
 ISOFLURANE, ISOFLURANE
 SEVOFLURANE, SEVOFLURANE

HANGZHOU BINJIANG

* HANGZHOU MINSHENG BINJIANG PHARMACEUTICAL CO LTD
 ALENDRONATE SODIUM, ALENDRONATE SODIUM

HANGZHOU ZHONGMEI

* HANGZHOU ZHONGMEI HUADONG PHARMACEUTICAL CO LTD
 DAPTOMYCIN, DAPTOMYCIN
 TACROLIMUS, TACROLIMUS

HARM REDUCTION THERP

* HARM REDUCTION THERAPEUTICS INC
 RIVIVE, NALOXONE HYDROCHLORIDE (OTC)

HARMAN FINOCHEM

* HARMAN FINOCHEM LTD
 ALLOPURINOL, ALLOPURINOL
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE

HARMONY

* HARMONY BIOSCIENCES LLC
 WAKIX, PITOLISANT HYDROCHLORIDE

HARROW EYE

* HARROW EYE LLC
 FLAREX, FLUOROMETHOLONE ACETATE
 IHEEZO, CHLOROPROCAINE HYDROCHLORIDE
 ILEVRO, NEPAFENAC
 IOPIDINE, APRACLOPIDINE HYDROCHLORIDE
 MAXIDEX, DEXAMETHASONE
 MAXITROL, DEXAMETHASONE
 NATAACYN, NATAMYCIN
 NEVANAC, NEPAFENAC
 TOBRADEX ST, DEXAMETHASONE
 TRIESENCE, TRIAMCINOLONE ACETONIDE
 VERKAZIA, CYCLOSPORINE
 VEVEYE, CYCLOSPORINE
 ZERVIAE, CETIRIZINE HYDROCHLORIDE

HBT LABS INC

* HBT LABS INC
 FULVESTRANT, FULVESTRANT

HEC PHARM

* HEC PHARM USA INC
 CLARITHROMYCIN, CLARITHROMYCIN
 LEVOFLOXACIN, LEVOFLOXACIN
 OLANZAPINE, OLANZAPINE
 PRASUGREL, PRASUGREL HYDROCHLORIDE

HEC PHARM CO LTD

* HEC PHARM CO LTD
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE

HELSINN

* HELSINN BIREX PHARMACEUTICALS LTD
 VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE

HELSINN HLTHCARE

* HELSINN HEALTHCARE SA
 AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
 AKYNZEO, NETUPITANT

HERITAGE LIFE

* HERITAGE LIFE SCIENCES BARBADOS INC
 CLOZARIL, CLOZAPINE

HERITAGE PHARMA

* HERITAGE PHARMA LABS INC
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HERITAGE PHARMA LABS INC
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIFLUNISAL, DIFLUNISAL
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METHIMAZOLE, METHIMAZOLE
 NIFEDIPINE, NIFEDIPINE

HERITAGE PHARMA AVET

* HERITAGE PHARMA LABS INC DBA AVET PHARMACEUTICALS LABS INC
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE

HERITAGE PHARMS

* HERITAGE PHARMACEUTICALS INC DBA AVET PHARMACEUTICALS INC
 ACYCLOVIR, ACYCLOVIR
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 FELODIPINE, FELODIPINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE, GLYBURIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LEFLUNOMIDE, LEFLUNOMIDE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 NYSTATIN, NYSTATIN
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

HERITAGE PHARMS INC

* HERITAGE PHARMACEUTICALS INC
 CALCIUM ACETATE, CALCIUM ACETATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE

HERITAGE PHARMS LABS

* HERITAGE PHARMACEUTICALS LABS INC DBA AVET PHARMACEUTICALS LABS INC
 BENZONATATE, BENZONATATE

HERON THERAPS INC

* HERON THERAPEUTICS INC
 APONVIE, APREPITANT
 CINVANTI, APREPITANT
 SUSTOL, GRANISETRON
 ZYNRELEF KIT, BUPIVACAINE

HETERO LABS

* HETERO LABS LTD
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LINEZOLID, LINEZOLID

HETERO LABS LTD III

* HETERO LABS LTD UNIT III

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HETERO LABS LTD UNIT III
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACYCLOVIR, ACYCLOVIR
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATOVAQUONE, ATOVAQUONE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOBAZAM, CLOBAZAM
 DABIGATRAN ETEXILATE MESYLATE, DABIGATRAN ETEXILATE MESYLATE
 DARUNAVIR, DARUNAVIR
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 EZETIMIBE, EZETIMIBE
 FENOFIBRATE, FENOFIBRATE
 FINASTERIDE, FINASTERIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LACOSAMIDE, LACOSAMIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVOCETIRIZINE HYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LORATADINE, LORATADINE (OTC)
 MARAVIROC, MARAVIROC
 METHOCARBAMOL, METHOCARBAMOL
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NAPROXEN, NAPROXEN
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NEVIRAPINE, NEVIRAPINE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXCARBAZEPINE, OXCARBAZEPINE
 POSACONAZOLE, POSACONAZOLE
 PREGABALIN, PREGABALIN
 RANOLAZINE, RANOLAZINE
 RITONAVIR, RITONAVIR
 ROFLUMILAST, ROFLUMILAST
 RUFINAMIDE, RUFINAMIDE
 SIMVASTATIN, SIMVASTATIN
 TADALAFIL, TADALAFIL
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TORSEMIDE, TORSEMIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 ZIDOVUDINE, ZIDOVUDINE

HETERO LABS LTD V

* HETERO LABS LTD UNIT V
 ACYCLOVIR, ACYCLOVIR
 ALLOPURINOL, ALLOPURINOL
 APIXABAN, APIXABAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HETERO LABS LTD UNIT V
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BREXPIPIRAZOLE, BREXPIPIRAZOLE
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLCHICINE, COLCHICINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ENTECAVIR, ENTECAVIR
 ESLICARBAZEPINE ACETATE, ESLICARBAZEPINE ACETATE
 ESZOPICLONE, ESZOPICLONE
 FAMCICLOVIR, FAMCICLOVIR
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 LACOSAMIDE, LACOSAMIDE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LENALIDOMIDE, LENALIDOMIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LINEZOLID, LINEZOLID
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIRFENIDONE, PIRFENIDONE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SODIUM PICOSULFATE, MAGNESIUM OXIDE AND ANHYDROUS CITRIC ACID, CITRIC ACID
 TELMISARTAN, TELMISARTAN
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TETRABENAZINE, TETRABENAZINE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TOLVAPTAN, TOLVAPTAN
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN

HEYL CHEMISCH

* HEYL CHEMISCH PHARMAZEUTISCHE FABRIK
 RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)

HIKMA

* HIKMA FARMACEUTICA PORTUGAL SA
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
 CEFTRIAXONE SODIUM, CEFTRIAXONE SODIUM
 CEFUROXIME SODIUM, CEFUROXIME SODIUM
 DOCETAXEL, DOCETAXEL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 ETOMIDATE, ETOMIDATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA FARMACEUTICA PORTUGAL SA
 VECURONIUM BROMIDE, VECURONIUM BROMIDE

* HIKMA PHARMACEUTICALS
 AMOXICILLIN, AMOXICILLIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE

* HIKMA PHARMACEUTICALS INTERNATIONAL LTD
 CODEINE SULFATE, CODEINE SULFATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 FUROSEMIDE, FUROSEMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 RUFINAMIDE, RUFINAMIDE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

* HIKMA PHARMACEUTICALS LLC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DOXERCALCIFEROL, DOXERCALCIFEROL
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

* HIKMA PHARMACEUTICALS USA INC
 ACARBOSE, ACARBOSE
 ACETAMINOPHEN, ACETAMINOPHEN
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ACYCLOVIR, ACYCLOVIR
 ADENOSINE, ADENOSINE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
 ALPRAZOLAM, ALPRAZOLAM
 ALPROSTADIL, ALPROSTADIL
 ALVIMOPAN, ALVIMOPAN
 AMICAR, AMINOCAPROIC ACID
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMRINONE LACTATE, INAMRINONE LACTATE
 ATIVAN, LORAZEPAM
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BEXAROTENE, BEXAROTENE
 BIMATOPROST, BIMATOPROST
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CAFKIT, CAFFEINE CITRATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CAPECITABINE, CAPECITABINE
 CARBOPLATIN, CARBOPLATIN
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFOXITIN, CEFOXITIN SODIUM
 CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CICLOPIROX, CICLOPIROX
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLADRIBINE, CLADRIBINE
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COMBOGESIC IV, ACETAMINOPHEN
 CORMAX, CLOBETASOL PROPIONATE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYCLOSPORINE, CYCLOSPORINE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEFERIPRONE, DEFERIPRONE
 DESONIDE, DESONIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXAMETHASONE INTENSOL, DEXAMETHASONE
 DEXAMETHASONE, DEXAMETHASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DIAZEPAM INTENSOL, DIAZEPAM
 DIAZEPAM, DIAZEPAM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOPRAM, DOXAPRAM HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DRONABINOL, DRONABINOL
 DURAMORPH PF, MORPHINE SULFATE
 EMBELINE, CLOBETASOL PROPIONATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETOMIDATE, ETOMIDATE
 ETOPOSIDE, ETOPOSIDE
 EVEROLIMUS, EVEROLIMUS
 EXEMESTANE, EXEMESTANE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FEBUXOSTAT, FEBUXOSTAT
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 GATIFLOXACIN, GATIFLOXACIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IMMPHENTIV, PHENYLEPHRINE HYDROCHLORIDE
 INFUMORPH, MORPHINE SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KLOXXADO, NALOXONE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN DEXTROSE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE IN SODIUM CHLORIDE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCARNITINE, LEVOCARNITINE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LORAZEPAM INTENSOL, LORAZEPAM
 LORAZEPAM, LORAZEPAM
 MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MESNA, MESNA
 METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALOXONE, NALOXONE HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA PHARMACEUTICALS USA INC
 OFLOXACIN, OFLOXACIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
 PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 POSACONAZOLE, POSACONAZOLE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISOLONE, PREDNISOLONE
 PREDNISON INTENSOL, PREDNISON
 PREDNISON, PREDNISON
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 PYRAZINAMIDE, PYRAZINAMIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 REGADENOSON, REGADENOSON
 REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 ROBAXIN, METHOCARBAMOL
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 RUFINAMIDE, RUFINAMIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM TETRADECYL SULFATE, SODIUM TETRADECYL SULFATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 THIOTEPA, THIOTEPA
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN
 TORSEMIDE, TORSEMIDE
 TRIAZOLAM, TRIAZOLAM
 VALPROIC ACID, VALPROIC ACID
 VALRUBICIN, VALRUBICIN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VIGABATRIN, VIGABATRIN
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINOELBINE TARTRATE, VINOELBINE TARTRATE
 VOSOL, ACETIC ACID, GLACIAL
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 ZALEPLON, ZALEPLON

HIKMA FARMACEUTICA

* HIKMA FARMACEUTICA (PORTUGAL) SA
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CEFOXITIN, CEFOXITIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HIKMA FARMACEUTICA (PORTUGAL) SA
 - CEFTRIAZONE, CEFTRIAZONE SODIUM
 - CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 - CIPROFLOXACIN, CIPROFLOXACIN
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - ENALAPRILAT, ENALAPRILAT
 - FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 - FLUMAZENIL, FLUMAZENIL
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE
 - MILRINONE LACTATE, MILRINONE LACTATE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PROGESTERONE, PROGESTERONE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - TERBUTALINE SULFATE, TERBUTALINE SULFATE
 - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 - VALPROATE SODIUM, VALPROATE SODIUM
- * HIKMA FARMACEUTICA PORTUGAL LDA
 - CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 - FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
- * HIKMA FARMACEUTICA PORTUGAL SA
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - OXYTOCIN, OXYTOCIN
 - TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
- * HIKMA FARMACEUTICA SA
 - ZOLEDRONIC ACID, ZOLEDRONIC ACID

HIKMA INTL PHARMS

- * HIKMA INTERNATIONAL PHARMACEUTICALS LLC
 - CAPTOPRIL, CAPTOPRIL
 - DIGOXIN, DIGOXIN
 - DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 - MITIGARE, COLCHICINE
 - NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE
 - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

HIKMA PHARM CO LTD

- * HIKMA PHARM CO LTD
 - ARGATROBAN, ARGATROBAN

HIKMA PHARMS

- * HIKMA PHARMACEUTICALS
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - AMOXICILLIN, AMOXICILLIN
 - BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 - BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 - DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 - PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 - RIFAMPIN, RIFAMPIN
 - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
- * HIKMA PHARMACEUTICALS CO LTD
 - PARICALCITOL, PARICALCITOL

HILL DERMAC

- * HILL DERMACEUTICALS INC
 - DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
 - DERMOTIC, FLUOCINOLONE ACETONIDE

HILL DERMACEUTICALS

- * HILL DERMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HILL DERMACEUTICALS INC
TOLAK, FLUOROURACIL

HISAMITSU

* HISAMITSU PHARMACEUTICAL CO INC
SECUADO, ASENAPINE

HISAMITSU PHARM CO

* HISAMITSU PHARMACEUTICAL CO INC
SALONPAS, MENTHOL (OTC)

HISUN PHARM HANGZHOU

* HISUN PHARMACEUTICAL (HANGZHOU) CO LTD
CLADRIBINE, CLADRIBINE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
OLANZAPINE, OLANZAPINE

* HISUN PHARMACEUTICAL HANGZHOU CO LTD
DACTINOMYCIN, DACTINOMYCIN
DAPTOMYCIN, DAPTOMYCIN
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
TICAGRELOR, TICAGRELOR

HLTHCARE

* HEALTHCARE PHARMACEUTICALS LTD
ATENOLOL, ATENOLOL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE

HOFFMANN LA ROCHE

* HOFFMANN LA ROCHE INC
ZELBORAF, VEMURAFENIB

HOFFMANN-LA ROCHE

* HOFFMANN-LA ROCHE INC
ALECENSA, ALECTINIB HYDROCHLORIDE

HONG KONG

* HONG KONG KING FRIEND INDUSTRIAL CO LTD
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM

HOPE PHARMS

* HOPE PHARMACEUTICALS
NITHIODE, SODIUM NITRITE
SODIUM NITRITE, SODIUM NITRITE
SODIUM THIOSULFATE, SODIUM THIOSULFATE

HORIZON

* HORIZON MEDICINES LLC
DUEXIS, FAMOTIDINE

* HORIZON THERAPEUTICS IRELAND DAC
PENNSAID, DICLOFENAC SODIUM

* HORIZON THERAPEUTICS USA INC
PROCYSBI, CYSTEAMINE BITARTRATE
RAYOS, PREDNISONE

HORIZON THERAP US

* HORIZON THERAPEUTICS US HOLDING LLC
BUPHENYL, SODIUM PHENYL BUTYRATE
RAVICTI, GLYCEROL PHENYL BUTYRATE

HOSPIRA

* HOSPIRA INC
ACETYLCYSTEINE, ACETYLCYSTEINE
ADENOSINE, ADENOSINE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
AMIDATE, ETOMIDATE
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOPHYLLINE, AMINOPHYLLINE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
ARGATROBAN, ARGATROBAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

ATROPINE SULFATE, ATROPINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 AZTREONAM, AZTREONAM
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 BIVALIRUDIN, BIVALIRUDIN
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BORTEZOMIB, BORTEZOMIB
 BUMETANIDE, BUMETANIDE
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CORLOPAM, FENOLDOPAM MESYLATE
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAPTOMYCIN, DAPTOMYCIN
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DEXTROSE 25%, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50%, DEXTROSE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 ENALAPRILAT, ENALAPRILAT
 EPINEPHRINE, EPINEPHRINE
 ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FUROSEMIDE, FUROSEMIDE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOPHED, NOREPINEPHRINE BITARTRATE
 LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
 LORAZEPAM, LORAZEPAM
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
 MANNITOL 25%, MANNITOL
 MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE, BUPIVACAINE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PACLITAXEL, PACLITAXEL
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PEMETREXED DITROMETHAMINE, PEMETREXED DITROMETHAMINE
 PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 POTASSIUM ACETATE, POTASSIUM ACETATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 QUELICIN, SUCCINYLCHOLINE CHLORIDE
 REGADENOSON, REGADENOSON
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM ACETATE, SODIUM ACETATE
 SODIUM BICARBONATE, SODIUM BICARBONATE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 14.6%, SODIUM CHLORIDE
 SODIUM CHLORIDE 23.4%, SODIUM CHLORIDE
 SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 TAZICEF, CEFTAZIDIME
 THAM, TROMETHAMINE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VITAMIN K1, PHYTONADIONE
 ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HOSPIRA INC
ZOLEDRONIC ACID, ZOLEDRONIC ACID
- * HOSPIRA WORLDWIDE, INC
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
VINCRIStINE SULFATE PFS, VINCRIStINE SULFATE

HOSPIRA INC

- * HOSPIRA INC
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
DOCETAXEL, DOCETAXEL
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
HEPARIN SODIUM, HEPARIN SODIUM
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
MORPHINE SULFATE, MORPHINE SULFATE
NIPENT, PENTOSTATIN
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
ZOLEDRONIC ACID, ZOLEDRONIC ACID

HOSPIRA WORLDWIDE

- * HOSPIRA WORLDWIDE PTY
OXALIPLATIN, OXALIPLATIN

HOT SHOTS NM LLC

- * HOT SHOTS NUCLEAR MEDICINE LLC
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

HQ SPCLT PHARMA

- * HQ SPECIALTY PHARMA CORP
CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CISPLATIN, CISPLATIN
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
LINEZOLID, LINEZOLID
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MEROPENEM, MEROPENEM

HQ SPECLT PHARMA

- * HQ SPECIALTY PHARMA CORP
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

HRA PHARMA

- * HRA PHARMA RARE DISEASES
LYSODREN, MITOTANE
METOPIRONE, METYRAPONE

HUMANWELL PURACAP

- * HUMANWELL PURACAP PHARMACEUTICAL WUHAN CO LTD
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
DUTASTERIDE, DUTASTERIDE
IBUPROFEN, IBUPROFEN (OTC)
ICOSAPENT ETHYL, ICOSAPENT ETHYL

HUONS

- * HUONS CO LTD
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

ROCHE

- * HOFFMANN LA ROCHE INC
FUZEON, ENFUVIRTIDE
TAMIFLU, OSELTAMIVIR PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******SHUANGCHENG**

* HAINAN SHUANGCHENG PHARMACEUTICALS CO LTD
 BIVALIRUDIN, BIVALIRUDIN
 EPTIFIBATIDE, EPTIFIBATIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

**** I ******I 3 PHARMS**

* I 3 PHARMACEUTICALS LLC
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 MARAVIROC, MARAVIROC
 POSACONAZOLE, POSACONAZOLE

I3 PHARMS

* I3 PHARMACEUTICALS LLC
 RAMELTEON, RAMELTEON
 RANOLAZINE, RANOLAZINE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

IBSA

* IBSA INSTITUT BIOCHIMIQUE SA
 FLECTOR, DICLOFENAC EPOLAMINE
 LIDOCAINE, LIDOCAINE
 TIROSINT, LEVOTHYROXINE SODIUM
 TIROSINT-SOL, LEVOTHYROXINE SODIUM

IBSA INST BIO

* IBSA INSTITUT BIOCHIMIQUE SA
 LICART, DICLOFENAC EPOLAMINE

ICHNOS

* ICHNOS SCIENCES SA
 DEFERASIROX, DEFERASIROX

ICU MEDICAL INC

* ICU MEDICAL INC
 ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
 AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
 AMINOSYN-PF 10%, AMINO ACIDS
 AMINOSYN-PF 7%, AMINO ACIDS
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
 NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* ICU MEDICAL INC**

POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION

IDORSIA*** IDORSIA PHARMACEUTICALS LTD**

QUVIVIQ, DARIDOREXANT HYDROCHLORIDE

IMPACT*** IMPACT BIOMEDICINES INC A WHOLLY OWNED SUB OF CELGENE CORP**

INREBIC, FEDRATINIB HYDROCHLORIDE

IMPAX*** IMPAX LABORATORIES LLC**

ADRENACLICK, EPINEPHRINE
 BACLOFEN, BACLOFEN
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE

IMPAX LABS*** IMPAX LABORATORIES INC**

ACARBOSE, ACARBOSE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIGOXIN, DIGOXIN
 DIPYRIDAMOLE, DIPYRIDAMOLE
 FENOFIBRATE, FENOFIBRATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RILUZOLE, RILUZOLE
 RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE

IMPAX LABS INC*** IMPAX LABORATORIES INC**

ACITRETIN, ACITRETIN
 BUDESONIDE, BUDESONIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 EMVERM, MEBENDAZOLE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HYDROCORTISONE, HYDROCORTISONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* IMPAX LABORATORIES INC**

HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RYTARY, CARBIDOPA

IMPAX PHARMS*** IMPAX PHARMACEUTICALS**

GEMFIBROZIL, GEMFIBROZIL
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

IMPEL PHARMS*** IMPEL PHARMACEUTICALS INC**

TRUDHESA, DIHYDROERGOTAMINE MESYLATE

IMS LTD*** INTERNATIONAL MEDICATION SYSTEMS LTD**

REGADENOSON, REGADENOSON

INCYTE CORP*** INCYTE CORP**

JAKAFI, RUXOLITINIB PHOSPHATE
 OPZELURA, RUXOLITINIB PHOSPHATE
 PEMAZYRE, PEMIGATINIB

INDCHEMIE HEALTH*** INDCHEMIE HEALTH SPECIALTIES PVT LTD**

ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE

INDICUS PHARMA*** INDICUS PHARMA LLC**

ACETAZOLAMIDE, ACETAZOLAMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE

INDIVIOR*** INDIVIOR INC**

OPVEE, NALMEFENE HYDROCHLORIDE
 PERSERIS KIT, RISPERIDONE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE

INDOCO*** INDOCO REMEDIES LTD**

ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR, ACYCLOVIR
 ALLOPURINOL, ALLOPURINOL
 APIXABAN, APIXABAN
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 FEBUXOSTAT, FEBUXOSTAT
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LACOSAMIDE, LACOSAMIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 OLANZAPINE, OLANZAPINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

INDOCO REMEDIES*** INDOCO REMEDIES LTD**

GLIMEPIRIDE, GLIMEPIRIDE

INFORLIFE*** INFORLIFE SA**

ACETAMINOPHEN, ACETAMINOPHEN
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* INFORLIFE SA**

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MIDAZOLAM IN 0.9% SODIUM CHLORIDE, MIDAZOLAM
 NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE, NOREPINEPHRINE BITARTRATE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

INGENUS PHARMS LLC*** INGENUS PHARMACEUTICALS LLC**

ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CABERGOLINE, CABERGOLINE
 CARMUSTINE, CARMUSTINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 TIMOLOL MALEATE, TIMOLOL MALEATE

INNOCOLL*** INNOCOLL PHARMACEUTICALS LTD**

POSIMIR, BUPIVACAINE

INNOCOLL PHARMS*** INNOCOLL PHARMACEUTICALS**

XARACOLL, BUPIVACAINE HYDROCHLORIDE

INNOGENIX*** INNOGENIX LLC**

BACLOFEN, BACLOFEN
 HALOPERIDOL, HALOPERIDOL
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 METOLAZONE, METOLAZONE
 METRONIDAZOLE, METRONIDAZOLE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 RAMELTEON, RAMELTEON

INNOPHARMA*** INNOPHARMA LICENSING LLC A SUB OF PFIZER INC**

PROPOFOL, PROPOFOL

INSMED INC*** INSMED INC**

ARIKAYCE KIT, AMIKACIN SULFATE

INTAS PHARMS USA*** INTAS PHARMACEUTICALS USA**

IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

INTELLIPHARMACEUTICS*** INTELLIPHARMACEUTICS CORP**

DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

INTERCEPT PHARMS INC*** INTERCEPT PHARMACEUTICALS INC**

OALIVA, OBETICHOLIC ACID

INTERSECT ENT INC*** INTERSECT ENT INC**

SINUVA, MOMETASONE FUROATE

INTL ISOTOPES*** INTERNATIONAL ISOTOPES INC**

SODIUM IODIDE I 131, SODIUM IODIDE I-131

INTL MEDICATION*** INTERNATIONAL MEDICATION SYSTEM**

LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 PHYTONADIONE, PHYTONADIONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* INTERNATIONAL MEDICATION SYSTEM
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE

INTL MEDICATION SYS

* INTERNATIONAL MEDICATION SYSTEMS LTD
 ATROPINE SULFATE, ATROPINE SULFATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 DEXTROSE 50%, DEXTROSE
 EPINEPHRINE, EPINEPHRINE
 LORAZEPAM, LORAZEPAM
 MORPHINE SULFATE, MORPHINE SULFATE
 SODIUM BICARBONATE, SODIUM BICARBONATE

INTRA-CELLULAR

* INTRA-CELLULAR THERAPIES INC
 CAPLYTA, LUMATEPERONE TOSYLATE

INVAGEN PHARMS

* INVAGEN PHARMACEUTICALS INC
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CALCIUM ACETATE, CALCIUM ACETATE
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 GEMFIBROZIL, GEMFIBROZIL
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LANREOTIDE ACETATE, LANREOTIDE ACETATE
 LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LEUPROLIDE ACETATE FOR DEPOT SUSPENSION, LEUPROLIDE ACETATE
 LEVETIRACETAM, LEVETIRACETAM
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEPROBAMATE, MEPROBAMATE
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NAPROXEN, NAPROXEN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PENICILLAMINE, PENICILLAMINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIMVASTATIN, SIMVASTATIN
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 VIGABATRIN, VIGABATRIN
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 ZONISAMIDE, ZONISAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******INVATECH**

* INVATECH PHARMA SOLUTIONS LLC
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

INVENTIA

* INVENTIA HEALTHCARE LTD
 CHLORTHALIDONE, CHLORTHALIDONE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 PALIPERIDONE, PALIPERIDONE
 TELMISARTAN, TELMISARTAN

INVENTIA HLTHCARE

* INVENTIA HEALTHCARE PRIVATE LTD
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

IONETIX

* IONETIX CORP
 AMMONIA N 13, AMMONIA N-13

IONIS PHARMS INC

* IONIS PHARMACEUTICALS INC
 WAINUA, EPLONTERSEN SODIUM

IPCA LABS LTD

* IPCA LABORATORIES LTD
 ALLOPURINOL, ALLOPURINOL
 ATENOLOL, ATENOLOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 ESZOPICLONE, ESZOPICLONE
 FUROSEMIDE, FUROSEMIDE
 GABAPENTIN, GABAPENTIN
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

IPR

* IPR PHARMACEUTICALS INC
 CRESTOR, ROSUVASTATIN CALCIUM

IPSEN

* IPSEN BIOPHARMACEUTICALS INC
 ONIVYDE, IRINOTECAN HYDROCHLORIDE
 SOHONOS, PALOVAROTENE

IPSEN PHARMA

* IPSEN PHARMA BIOTECH SAS
 SOMATULINE DEPOT, LANREOTIDE ACETATE

IRONSHORE PHARMS

* IRONSHORE PHARMACEUTICALS AND DEVELOPMENT INC
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE

ISOLOGIC INNOVATIVE

* ISOLOGIC INNOVATIVE RADIOPHARMACEUTICALS LTD
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

ISOTEX

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* ISOTEX DIAGNOSTICS**

GLOFIL-125, IOTHALAMATE SODIUM I-125

ISTITUTO BIO ITA SPA*** ISTITUTO BIOCHIMICO ITALIANO SPA**AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
NAFCILLIN SODIUM, NAFCILLIN SODIUM
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PIPERACILLIN, PIPERACILLIN SODIUM**ITALFARMACO SA***** ITALFARMACO SA**

TIGLUTIK KIT, RILUZOLE

IVAX SUB TEVA PHARMS*** IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA**ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
BACLOFEN, BACLOFEN
CABERGOLINE, CABERGOLINE
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CLOZAPINE, CLOZAPINE
CYCLOSPORINE, CYCLOSPORINE
DIAZEPAM, DIAZEPAM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
FAMOTIDINE, FAMOTIDINE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE**IVERIC BIO***** IVERIC BIO INC**

IZERVAY, AVACINCAPTAD PEGOL SODIUM

**** J ******HENGRUI PHARMA***** JIANGSU HENGRUI PHARMACEUTICALS CO LTD**CARMUSTINE, CARMUSTINE
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DAPTOMYCIN, DAPTOMYCIN
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
GADOBUTROL, GADOBUTROL
GADOTERATE MEGLUMINE, GADOTERATE MEGLUMINE
IODIXANOL, IODIXANOL
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
THIOTEPA, THIOTEPA**J AND J CONSUMER INC***** JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIV**CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
CHILDREN'S MOTRIN, IBUPROFEN (OTC)
CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
MOTRIN IB, IBUPROFEN (OTC)
PEPCID AC, FAMOTIDINE (OTC)
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
RHINOCORT ALLERGY, BUDESONIDE (OTC)
SINE-AID IB, IBUPROFEN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIV
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 TYLENOL, ACETAMINOPHEN (OTC)
 ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

JANSSEN BIOTECH

* JANSSEN BIOTECH INC
 AKEEGA, ABIRATERONE ACETATE
 BALVERSA, ERDAFITINIB
 ERLEADA, APALUTAMIDE
 ZYTIGA, ABIRATERONE ACETATE

JANSSEN PHARMS

* JANSSEN PHARMACEUTICALS INC
 CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 HALDOL, HALOPERIDOL DECANOATE
 INVEGA HAFYERA, PALIPERIDONE PALMITATE
 INVEGA SUSTENNA, PALIPERIDONE PALMITATE
 INVEGA TRINZA, PALIPERIDONE PALMITATE
 INVEGA, PALIPERIDONE
 INVOKAMET XR, CANAGLIFLOZIN
 INVOKAMET, CANAGLIFLOZIN
 INVOKANA, CANAGLIFLOZIN
 PONVORY, PONESIMOD
 RISPERDAL CONSTA, RISPERIDONE
 RISPERDAL, RISPERIDONE
 SPORANOX, ITRACONAZOLE
 SPRAVATO, ESKETAMINE HYDROCHLORIDE
 TOPAMAX, TOPIRAMATE
 XARELTO, RIVAROXABAN

JANSSEN PRODS

* JANSSEN PRODUCTS LP
 EDURANT, RILPIVIRINE HYDROCHLORIDE
 PREZCOBIX, COBICISTAT
 PREZISTA, DARUNAVIR
 SYMTUZA, COBICISTAT
 YONDELIS, TRABECTEDIN

JANSSEN R AND D

* JANSSEN RESEARCH AND DEVELOPMENT LLC
 INTELENCE, ETRAVIRINE

JANSSEN THERAP

* JANSSEN THERAPEUTICS DIV JANSSEN PRODUCTS LP
 SIRTURO, BEDAQUILINE FUMARATE

JAZZ

* JAZZ PHARMACEUTICALS IRELAND LTD
 XYWAV, CALCIUM OXYBATE
 ZEPZELCA, LURBINECTEDIN

JAZZ PHARMS

* JAZZ PHARMACEUTICALS INC
 XYREM, SODIUM OXYBATE

JAZZ PHARMS INC

* JAZZ PHARMACEUTICALS INC
 DEFITELIO, DEFIBROTIDE SODIUM

JAZZ PHARMS RES

* JAZZ PHARMACEUTICALS RESEARCH UK LTD
 EPIDIOLEX, CANNABIDIOL

JDP

* JDP THERAPEUTICS LLC
 QUZYTIR, CETIRIZINE HYDROCHLORIDE

JIANGSU HANSOH PHARM

* JIANGSU HANSOH PHARMACEUTICAL GROUP CO LTD
 AZACITIDINE, AZACITIDINE
 BORTEZOMIB, BORTEZOMIB
 DECITABINE, DECITABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JIANGSU HANSOH PHARMACEUTICAL GROUP CO LTD
 FULVESTRANT, FULVESTRANT
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 MICAFFUNGIN SODIUM, MICAFFUNGIN SODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 VINOELBINE TARTRATE, VINOELBINE TARTRATE

JOHNS HOPKINS UNIV

* JOHNS HOPKINS UNIV
 AMMONIA N 13, AMMONIA N-13

JOHNSON AND JOHNSON

* JOHNSON AND JOHNSON CONSUMER INC
 VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
 VISINE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 * JOHNSON AND JOHNSON GROUP CONSUMER COMPANIES
 MEN'S ROGAINE, MINOXIDIL (OTC)
 ROGAINE (FOR MEN), MINOXIDIL (OTC)
 ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
 ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 WOMEN'S ROGAINE, MINOXIDIL (OTC)

JOURNEY

* JOURNEY MEDICAL CORP
 AMZEEQ, MINOCYCLINE HYDROCHLORIDE
 EURAX, CROTAMITON
 EXELDERM, SULCONAZOLE NITRATE
 QBREXZA, GLYCOPYRRONIUM TOSYLATE
 ZILXI, MINOCYCLINE HYDROCHLORIDE

JUBILANT

* JUBILANT DRAXIMAGE INC DBA JUBILANT RADIOPHARMA
 DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT
 DRAXIMAGE DTPA, TECHNETIUM TC-99M PENTETATE KIT
 DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
 HICON, SODIUM IODIDE I-131
 RUBY-FILL, RUBIDIUM CHLORIDE RB-82
 SODIUM IODIDE I 131, SODIUM IODIDE I-131
 TECHNETIUM TC-99M SULFUR COLLOID KIT, TECHNETIUM TC-99M SULFUR COLLOID KIT

JUBILANT CADISTA

* JUBILANT CADISTA PHARMACEUTICALS INC
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 PREDNISONE, PREDNISONE
 PROCOMP, PROCHLORPERAZINE MALEATE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

JUBILANT DRAXIMAGE

* JUBILANT DRAXIMAGE INC
 TECHNETIUM TC 99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
 * JUBILANT DRAXIMAGE RADIOPHARMACIES INC
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
 * JUBILANT DRAXIMAGE USA INC
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT

JUBILANT GENERICS

* JUBILANT GENERICS LTD
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JUBILANT GENERICS LTD
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 IRBESARTAN, IRBESARTAN
 OLANZAPINE, OLANZAPINE
 RISPERIDONE, RISPERIDONE
 SPIRONOLACTONE, SPIRONOLACTONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 ZOLMITRIPTAN, ZOLMITRIPTAN

STEVENS J

* JEROME STEVENS PHARMACEUTICALS INC
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 DIGOXIN, DIGOXIN
 UNITHROID, LEVOTHYROXINE SODIUM **

**** K ******GRIFFEN**

* KW GRIFFEN CO
 BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

KADMON PHARMS LLC

* KADMON PHARMACEUTICALS LLC
 REZUROCK, BELUMOSUDIL MESYLATE

KAI PHARMS INC

* KAI PHARMACEUTICALS INC A WHOLLY OWNED SUBSIDIARY OF AMGEN INC
 PARSABIV, ETELCALCETIDE

KALEO INC

* KALEO INC
 AUVI-Q, EPINEPHRINE

KARYOPHARM THERAPS

* KARYOPHARM THERAPEUTICS INC
 XPOVIO, SELINEXOR

KENTON

* KENTON CHEMICALS AND PHARMACEUTICALS CORP
 ANASTROZOLE, ANASTROZOLE
 CAPTOPRIL, CAPTOPRIL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 RILUZOLE, RILUZOLE

KERYX BIOPHARMS

* KERYX BIOPHARMACEUTICALS INC
 AURYXIA, FERRIC CITRATE

KETTERING MEDCTR

* KETTERING MEDCTR
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

KINDEVA

* KINDEVA DRUG DELIVERY LP
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL
 PROVENTIL-HFA, ALBUTEROL SULFATE

KINDOS

* KINDOS PHARMACEUTICALS CO LTD
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 NELARABINE, NELARABINE
 PLERIXAFOR, PLERIXAFOR
 THIOTEPA, THIOTEPA

KING PHARMS

* KING PHARMACEUTICALS INC
 SYNERCID, DALFOPRISTIN
 * KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ****

* KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC
 CYTOMEL, LIOTHYRONINE SODIUM
 LEVOXYL, LEVOTHYROXINE SODIUM **

KING PHARMS LLC

* KING PHARMACEUTICALS LLC
 ALTACE, RAMIPRIL
 BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
 BICILLIN C-R, PENICILLIN G BENZATHINE
 BICILLIN L-A, PENICILLIN G BENZATHINE
 PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
 SILVADENE, SILVER SULFADIAZINE

KNACK

* KNACK PHARMACEUTICALS INC
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

KNIGHT THERAPS

* KNIGHT THERAPEUTICS USA INC
 IMPAVIDO, MILTEFOSINE

KOWA CO

* KOWA CO LTD
 LIVALO, PITAVASTATIN CALCIUM

KOWA PHARMS

* KOWA PHARMACEUTICALS AMERICA INC
 SEGLENTIS, CELECOXIB

KRAMER

* KRAMER LABORATORIES INC
 NIZORAL ANTI-DANDRUFF, KETOCONAZOLE (OTC)

KREITCHMAN PET CTR

* KREITCHMAN PET CENTER
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

KVK TECH

* KVK TECH INC
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 KALEXATE, SODIUM POLYSTYRENE SULFONATE
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

KVK TECH INC

* KVK TECH INC
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

KYOWA KIRIN

* KYOWA KIRIN INC
 FARESTON, TOREMIFENE CITRATE
 NOURIANZ, ISTRADÉFYLLINE

KYTHERA BIOPHARMS

* KYTHERA BIOPHARMACEUTICALS INC
 KYBELLA, DEOXYCHOLIC ACID

**** L ******L PERRIGO CO**

* L PERRIGO CO
 ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
 CIMETIDINE, CIMETIDINE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* L PERRIGO CO**

NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

LA JOLLA PHARMA

* LA JOLLA PHARMA LLC
 GIAPREZA, ANGIOTENSIN II ACETATE

LAB HRA PHARMA

* LABORATOIRE HRA PHARMA
 ELLA, ULIPRISTAL ACETATE

LABORATOIRE HRA

* LABORATOIRE HRA PHARMA
 LEVONORGESTREL, LEVONORGESTREL (OTC)

LABORATORIOS GRIFOLS

* LABORATORIOS GRIFOLS SA
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LABORATORIOS SALVAT

* LABORATORIOS SALVAT SA
 OTOVEL, CIPROFLOXACIN HYDROCHLORIDE

LABS JUVISE

* LABORATOIRES JUVISE PHARMACEUTICALS
 PYLERA, BISMUTH SUBCITRATE POTASSIUM

LACER PHARMA

* LACER PHARMA LLC
 ERTACZO, SERTACONAZOLE NITRATE

LANDELA PHARM

* LANDELA PHARMACEUTICAL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

LANNETT

* LANNETT CO INC
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 LANORINAL, ASPIRIN
 PRIMIDONE, PRIMIDONE
 PROBALAN, PROBENECID

LANNETT CO INC

* LANNETT CO INC
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BACLOFEN, BACLOFEN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CODEINE SULFATE, CODEINE SULFATE
 DANAZOL, DANAZOL
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LORATADINE, LORATADINE (OTC)
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 METAXALONE, METAXALONE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LANNETT CO INC**

MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NIACIN, NIACIN
 OMEPRAZOLE, OMEPRAZOLE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SUMATRIPTAN, SUMATRIPTAN
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 URSODIOL, URSODIOL

LANTHEUS MEDCL*** LANTHEUS MEDICAL IMAGING INC**

CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
 DEFINITY RT, PERFLUTREN
 DEFINITY, PERFLUTREN
 NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
 TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 XENON XE 133, XENON XE-133

LARKEN LABS*** LARKEN LABORATORIES INC**

DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 OFLOXACIN, OFLOXACIN

LARKEN LABS INC*** LARKEN LABORATORIES INC**

ALLZITAL, ACETAMINOPHEN
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 DEXAMETHASONE, DEXAMETHASONE

LAURUS*** LAURUS LABS LTD**

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 GABAPENTIN, GABAPENTIN
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIRFENIDONE, PIRFENIDONE

LAVIPHARM*** LAVIPHARM SA**

CATAPRES-TTS-1, CLONIDINE
 CATAPRES-TTS-2, CLONIDINE
 CATAPRES-TTS-3, CLONIDINE

LEADIANT BIOSCI INC*** LEADIANT BIOSCIENCES INC**

ABELCET, AMPHOTERICIN B
 CARNITOR SF, LEVOCARNITINE
 CARNITOR, LEVOCARNITINE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 MATULANE, PROCARBAZINE HYDROCHLORIDE

LEADING*** LEADING PHARMA LLC**

BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LEADING PHARMA LLC
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYUREA, HYDROXYUREA
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LORAZEPAM, LORAZEPAM
 NIFEDIPINE, NIFEDIPINE
 NYSTATIN, NYSTATIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 THEOPHYLLINE, THEOPHYLLINE

LEADING PHARMA

* LEADING PHARMA LLC
 LACOSAMIDE, LACOSAMIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE

LEGACY PHARMA

* LEGACY PHARMA INC
 ALA-SCALP, HYDROCORTISONE
 BETAPACE AF, SOTALOL HYDROCHLORIDE
 BETAPACE, SOTALOL HYDROCHLORIDE
 CROTAN, CROTAMITON

LEGACY PHARMA USA

* LEGACY PHARMA USA INC
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE

LEO PHARMA AS

* LEO PHARMA AS
 DOVONEX, CALCIPOTRIENE
 ENSTILAR, BETAMETHASONE DIPROPIONATE
 FINACEA, AZELAIC ACID
 PROTOPIC, TACROLIMUS
 TACLONEX, BETAMETHASONE DIPROPIONATE

LEXICON PHARMS INC

* LEXICON PHARMACEUTICALS INC
 INPEFA, SOTAGLIFLOZIN

LGM PHARMA

* LGM PHARMA SOLUTIONS LLC
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 CHENODIOL, CHENODIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCODONE BITARTRATE AND ASPIRIN, ASPIRIN
 MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 URSODIOL, URSODIOL

LIEBEL-FLARSHEIM

* LIEBEL-FLARSHEIM CO LLC
 CONRAY, IOTHALAMATE MEGLUMINE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
 OPTIRAY 300, IOVERSOL
 OPTIRAY 320, IOVERSOL
 OPTIRAY 350, IOVERSOL
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LIFE MOLECULAR

* LIFE MOLECULAR IMAGING LTD
 NEURACEQ, FLORBETABEN F-18

LINDE GAS EQUIP

* LINDE GAS AND EQUIPMENT INC
 NOXIVENT, NITRIC OXIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******LNHC**

* LNHC INC
SITAVIG, ACYCLOVIR

LNK

* LNK INTERNATIONAL INC
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

LNK INTL INC

* LNK INTERNATIONAL INC
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

LONG GROVE PHARMS

* LONG GROVE PHARMACEUTICALS LLC
AK-FLUOR 10%, FLUORESCEIN SODIUM
AK-FLUOR 25%, FLUORESCEIN SODIUM
CALCITRIOL, CALCITRIOL
CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

LOREAL USA

* LOREAL USA PRODUCTS INC
ANTHELIOS 20, AVOBENZONE (OTC)
ANTHELIOS 40, AVOBENZONE (OTC)
ANTHELIOS SX, AVOBENZONE (OTC)
CAPITAL SOLEIL 15, AVOBENZONE (OTC)

LOTUS PHARM CO LTD

* LOTUS PHARMACEUTICAL CO LTD NANTOU PLANT
LENALIDOMIDE, LENALIDOMIDE
METHOTREXATE SODIUM, METHOTREXATE SODIUM

LOXO ONCOL

* LOXO ONCOLOGY INC
JAYPIRCA, PIRTOBRUTINIB

LOXO ONCOL ELI LILLY

* LOXO ONCOLOGY INC A WHOLLY OWNED SUB OF ELI LILLY AND CO
RETEVMO, SELPERCATINIB

LUITPOLD

* LUITPOLD PHARMACEUTICALS INC
AMINOCAPROIC ACID, AMINOCAPROIC ACID

LUKARE MEDICAL LLC

* LUKARE MEDICAL LLC
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE

LUNDBECK NA LTD

* LUNDBECK NA LTD
NORTHERA, DROXIDOPA

LUNDBECK PHARMS LLC

* LUNDBECK PHARMACEUTICALS LLC
ONFI, CLOBAZAM
SABRIL, VIGABATRIN

LUOXIN AUROVITAS

* LUOXIN AUROVITAS PHARMA CHENGDU CO LTD
ALBUTEROL SULFATE, ALBUTEROL SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

LUPIN

* LUPIN INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ANTARA (MICRONIZED), FENOFIBRATE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
BROVANA, ARFORMOTEROL TARTRATE
BUDESONIDE, BUDESONIDE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CYANOCOBALAMIN, CYANOCOBALAMIN
DESOXIMETASONE, DESOXIMETASONE
DIMETHYL FUMARATE, DIMETHYL FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LUPIN INC**

FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SOLOSEC, SECNIDAZOLE
 TESTOSTERONE, TESTOSTERONE
 TIOTROPIUM BROMIDE, TIOTROPIUM BROMIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 XOPENEX HFA, LEVALBUTEROL TARTRATE

*** LUPIN LTD**

AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 CARVEDILOL, CARVEDILOL
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 LOVASTATIN, LOVASTATIN
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 TRANDOLAPRIL, TRANDOLAPRIL

LUPIN LTD*** LUPIN LIMITED**

LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

*** LUPIN LTD**

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE
 ALLOPURINOL, ALLOPURINOL
 AMABELZ, ESTRADIOL
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 ARMODAFINIL, ARMODAFINIL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATOVAQUONE, ATOVAQUONE
 AZITHROMYCIN, AZITHROMYCIN
 BEKYREE, DESOGESTREL
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BIMATOPROST, BIMATOPROST
 BLISOVI 24 FE, ETHINYL ESTRADIOL
 BLISOVI FE 1.5/30, ETHINYL ESTRADIOL
 BLISOVI FE 1/20, ETHINYL ESTRADIOL
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CALCIUM ACETATE, CALCIUM ACETATE
 CELECOXIB, CELECOXIB
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LUPIN LTD
 DARUNAVIR, DARUNAVIR
 DAYSEE, ETHINYL ESTRADIOL
 DECITABINE, DECITABINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENSKYCE, DESOGESTREL
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FALLBACK SOLO, LEVONORGESTREL (OTC)
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FYAVOLV, ETHINYL ESTRADIOL
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GATIFLOXACIN, GATIFLOXACIN
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 JENCYCLA, NORETHINDRONE
 KAITLIB FE, ETHINYL ESTRADIOL
 KETOROLAC TROMETHAMINE AND PHENYLEPHRINE HYDROCHLORIDE, KETOROLAC TROMETHAMINE
 KURVELO, ETHINYL ESTRADIOL
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOTE Prednol ETABONATE, LOTE Prednol ETABONATE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEFENAMIC ACID, MEFENAMIC ACID
 MELOXICAM, MELOXICAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRABEGRON, MIRABEGRON
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NIACIN, NIACIN
 NIKKI, DROSPIRENONE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LUPIN LTD**

PENICILLAMINE, PENICILLAMINE
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANOLAZINE, RANOLAZINE
 RIFABUTIN, RIFABUTIN
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SUPRAX, CEFIXIME
 TADALAFIL, TADALAFIL
 TAVABOROLE, TAVABOROLE
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TENOFOVIR ALAFENAMIDE, TENOFOVIR ALAFENAMIDE FUMARATE
 TESTOSTERONE, TESTOSTERONE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TURQOZ, ETHINYL ESTRADIOL
 TYDEMY, DROSPIRENONE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VYFEMLA, ETHINYL ESTRADIOL
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

LUPIN PHARMS*** LUPIN PHARMACEUTICALS INC**

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 DESLORATADINE, DESLORATADINE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DROXIDOPA, DROXIDOPA
 MELOXICAM, MELOXICAM
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 RIFAMPIN, RIFAMPIN
 SUPRAX, CEFIXIME
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

LYNE*** LYNE LABORATORIES INC**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXAMETHASONE, DEXAMETHASONE
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 LEVOCARNITINE, LEVOCARNITINE
 NYSTATIN, NYSTATIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

LYRUS LIFE SCIENCES*** LYRUS LIFE SCIENCES PRIVATE LTD**

METHENAMINE HIPPURATE, METHENAMINE HIPPURATE

PERRIGO*** L PERRIGO CO**

ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
 DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* L PERRIGO CO
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 TIOCONAZOLE, TIOCONAZOLE (OTC)

**** M ******MA GENERAL HOSP**

* MASSACHUSETTS GENERAL HOSP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MACLEODS PHARMS LTD

* MACLEODS PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CELECOXIB, CELECOXIB
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENTACAPONE, ENTACAPONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESZOPICLONE, ESZOPICLONE
 FAMCICLOVIR, FAMCICLOVIR
 FEBUXOSTAT, FEBUXOSTAT
 FLUOCINONIDE, FLUOCINONIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LIDOCAINE, LIDOCAINE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 NIACIN, NIACIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 PYRAZINAMIDE, PYRAZINAMIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 REPAGLINIDE, REPAGLINIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MACLEODS PHARMACEUTICALS LTD**

SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SEVELAMER HYDROCHLORIDE, SEVELAMER HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

MAIA PHARMS INC*** MAIA PHARMACEUTICALS INC**

ANGIOMAX RTU, BIVALIRUDIN
 BACLOFEN, BACLOFEN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 SINCALIDE, SINCALIDE
 SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

MAINPOINTE*** MAINPOINTE PHARMACEUTICALS LLC**

TUXARIN ER, CHLORPHENIRAMINE MALEATE

MALLINCKRODT ARD*** MALLINCKRODT ARD INC**

ACTHAR GEL, CORTICOTROPIN

MALLINCKRODT HOSP*** MALLINCKRODT HOSP PRODUCTS IP LTD**

INOMAX, NITRIC OXIDE
 UVADEX, METHOXSALEN

MALLINCKRODT IRELAND*** MALLINCKRODT PHARMACEUTICALS IRELAND LTD**

TERLIVAZ, TERLIPRESSIN ACETATE

MANKIND PHARMA*** MANKIND PHARMA LTD**

ACETAZOLAMIDE, ACETAZOLAMIDE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATROPINE SULFATE, ATROPINE SULFATE
 BACLOFEN, BACLOFEN
 CHLORTHALIDONE, CHLORTHALIDONE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRATE, FENOFIBRATE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 FUROSEMIDE, FUROSEMIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NITROGLYCERIN, NITROGLYCERIN
 OFLOXACIN, OFLOXACIN
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MANKIND PHARMA LTD
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBRAMYCIN, TOBRAMYCIN
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

MARINUS

* MARINUS PHARMACEUTICALS INC
 ZTALMY, GANAXOLONE

MARIUS PHARMS LLC

* MARIUS PHARMACEUTICALS LLC
 KYZATREX, TESTOSTERONE UNDECANOATE

MARKSANS PHARMA

* MARKSANS PHARMA LTD
 ACETAMINOPHEN AND IBUPROFEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 PARICALCITOL, PARICALCITOL

MAYNE PHARMA

* MAYNE PHARMA INTERNATIONAL PTY LTD
 DORYX MPC, DOXYCYCLINE HYCLATE
 DORYX, DOXYCYCLINE HYCLATE
 TOLSURA, ITRACONAZOLE

* MAYNE PHARMA LLC
 ANNOVERA, ETHINYL ESTRADIOL
 BIJUVA, ESTRADIOL
 FABIOR, TAZAROTENE
 IMVEXXY, ESTRADIOL
 LEXETTE, HALOBETASOL PROPIONATE
 NEXTSTELLIS, DROSPIRENONE
 RHOFAD, OXYMETAZOLINE HYDROCHLORIDE
 SORILUX, CALCIPOTRIENE

MAYNE PHARMA COMMRCCL

* MAYNE PHARMA COMMERCIAL LLC
 SOLTAMOX, TAMOXIFEN CITRATE

MC2

* MC2 THERAPEUTICS LTD
 WYNZORA, BETAMETHASONE DIPROPIONATE

MCGUFF

* MCGUFF PHARMACEUTICALS INC
 ASCOR, ASCORBIC ACID

MCNEIL

* MCNEIL CONSUMER PRODUCTS CO DIV MCNEILAB INC
 IBUPROFEN, IBUPROFEN (OTC)

MCPRF

* MAYO CLINIC PET RADIOCHEMISTRY FACILITY
 AMMONIA N 13, AMMONIA N-13
 CHOLINE C-11, CHOLINE C-11
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MDD US

* MDD US OPERATIONS LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MDD US OPERATIONS LLC
 APOKYN, APOMORPHINE HYDROCHLORIDE
 XADAGO, SAFINAMIDE MESYLATE

MDGH

* MEDICINES DEVELOPMENT FOR GLOBAL HEALTH
 MOXIDECTIN, MOXIDECTIN

MEDEFIL INC

* MEDEFIL INC
 ATROPINE SULFATE, ATROPINE SULFATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION

MEDEXUS

* MEDEXUS PHARMA INC
 RASUVO, METHOTREXATE

MEDI-RADIOPHARMA

* MEDI-RADIOPHARMA LTD
 TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT

MEDICINES360

* MEDICINES360
 LILETTA, LEVONORGESTREL

MEDICURE

* MEDICURE INTERNATIONAL INC
 AGGRASTAT, TIROFIBAN HYDROCHLORIDE
 ZYPITAMAG, PITAVASTATIN MAGNESIUM

MEDIMETRIKS PHARMS

* MEDIMETRIKS PHARMACEUTICALS INC
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 LOPROX, CICLOPIROX
 NEO-SYNALAR, FLUOCINOLONE ACETONIDE
 SYNALAR, FLUOCINOLONE ACETONIDE

MEDLINE INDUSTRIES

* MEDLINE INDUSTRIES INC
 READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)

MEDTECH PRODUCTS

* MEDTECH PRODUCTS INC
 MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
 MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 3, MICONAZOLE NITRATE
 MONISTAT 3, MICONAZOLE NITRATE (OTC)
 MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 7, MICONAZOLE NITRATE (OTC)
 NIX, PERMETHRIN (OTC)
 TAGAMET HB, CIMETIDINE (OTC)

MEDUNIK

* MEDUNIK CANADA INC
 NITISINONE, NITISINONE
 PHEBURANE, SODIUM PHENYL BUTYRATE

MEITHEAL

* MEITHEAL PHARMACEUTICALS INC
 ADENOSINE, ADENOSINE
 ALPROSTADIL, ALPROSTADIL
 AMIKACIN SULFATE, AMIKACIN SULFATE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZACITIDINE, AZACITIDINE
 BIVALIRUDIN, BIVALIRUDIN
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BORTEZOMIB, BORTEZOMIB
 BUSULFAN, BUSULFAN
 CARMUSTINE, CARMUSTINE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLOFARABINE, CLOFARABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MEITHEAL PHARMACEUTICALS INC
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DACTINOMYCIN, DACTINOMYCIN
 DAPTOMYCIN, DAPTOMYCIN
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DOXERCALCIFEROL, DOXERCALCIFEROL
 ETOPOSIDE, ETOPOSIDE
 FUROSEMIDE, FUROSEMIDE
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 REGADENOSON, REGADENOSON
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

MELINTA

* MELINTA SUBSIDIARY CORP
 BAXDELA, DELAFLOXACIN MEGLUMINE

MELINTA THERAP

* MELINTA THERAPEUTICS LLC
 KIMYRSA, ORITAVANCIN DIPHOSPHATE
 ORBACTIV, ORITAVANCIN DIPHOSPHATE

MEM SLOAN-KETTERING

* MEMORIAL SLOAN-KETTERING CANCER CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MERCK

* MERCK AND CO INC
 CANCIDAS, CASPOFUNGIN ACETATE
 EMEND, APREPITANT
 PRIMAXIN, CILASTATIN SODIUM

MERCK AND CO INC

* MERCK AND CO INC
 EMEND, FOSAPREPITANT DIMEGLUMINE

MERCK SHARP DOHME

* MERCK SHARP AND DOHME CORP

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MERCK SHARP AND DOHME CORP
BELSOMRA, SUVOREXANT
JANUVIA, SITAGLIPTIN PHOSPHATE
NOXAFIL, POSACONAZOLE
PREVYMIS, LETERMOVIR
STROMECTOL, IVERMECTIN
TEMODAR, TEMOZOLOMIDE
VERQUVO, VERICIGUAT
- * MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
WELIREG, BELZUTIFAN

MERIDIAN BIOSCIENCE

- * MERIDIAN BIOSCIENCE ISRAEL LTD
IDKIT:HP, CITRIC ACID

MERRO PHARM USA

- * MERRO PHARMACEUTICAL USA INC
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

MERZ PHARMS

- * MERZ PHARMACEUTICALS LLC
CUVPOSA, GLYCOPYRROLATE

METACEL PHARMS LLC

- * METACEL PHARMACEUTICALS LLC
OZOBAX DS, BACLOFEN

METHAPHARM

- * METHAPHARM INC
PROVOCHOLINE, METHACHOLINE CHLORIDE

METUCHEN PHARMS

- * METUCHEN PHARMACEUTICALS LLC
STENDRA, AVANAFIL

MICRO LABS

- * MICRO LABS LTD
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
BACLOFEN, BACLOFEN
BIMATOPROST, BIMATOPROST
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CAFFEINE CITRATE, CAFFEINE CITRATE
CELECOXIB, CELECOXIB
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLOBAZAM, CLOBAZAM
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CROMOLYN SODIUM, CROMOLYN SODIUM
DALFAMPRIDINE, DALFAMPRIDINE
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ERYTHRA-DERM, ERYTHROMYCIN
FAMOTIDINE, FAMOTIDINE
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
GLIMEPIRIDE, GLIMEPIRIDE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
LIFITEGRAST, LIFITEGRAST
LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MEFENAMIC ACID, MEFENAMIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MICRO LABS LTD**

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MISOPROSTOL, MISOPROSTOL
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 PIRFENIDONE, PIRFENIDONE
 PIROXICAM, PIROXICAM
 RAMELTEON, RAMELTEON
 RANOLAZINE, RANOLAZINE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 ROFLUMILAST, ROFLUMILAST
 RUFINAMIDE, RUFINAMIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIMVASTATIN, SIMVASTATIN
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TAFLUPROST, TAFLUPROST
 TELMISARTAN, TELMISARTAN
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBRAMYCIN, TOBRAMYCIN
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAVOPROST, TRAVOPROST
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

MICRO LABS LTD*** MICRO LABS LTD**

NEVIRAPINE, NEVIRAPINE

MICRO LABS LTD INDIA*** MICRO LABS LTD INDIA**

ACETAZOLAMIDE, ACETAZOLAMIDE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CROMOLYN SODIUM, CROMOLYN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN

MIDWEST MEDCL*** MIDWEST MEDICAL ISOTOPES LLC CYCLOTRON DIV**

AMMONIA N 13, AMMONIA N-13
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MIKART*** MIKART LLC**

BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTAPAP, ACETAMINOPHEN
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 CHLORZOAZONE, CHLORZOAZONE
 ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METHAZOLAMIDE, METHAZOLAMIDE
 METHOCARBAMOL, METHOCARBAMOL
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

MILLA PHARMS*** MILLA PHARMACEUTICALS INC**

DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 SODIUM ACETATE, SODIUM ACETATE

MILLICENT*** MILLICENT HOLDINGS LTD**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MILLICENT HOLDINGS LTD
FEMRING, ESTRADIOL ACETATE

* MILLICENT PHARMA LTD
INTRAROSA, PRASTERONE

MIPS CRF

* MIPS CYCLOTRON AND RADIOCHEMISTRY FACILITY
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MIRATI THERAPS

* MIRATI THERAPEUTICS INC
KRAZATI, ADAGRASIB

MIRUM

* MIRUM PHARMACEUTICALS INC
CHOLBAM, CHOLIC ACID
LIVMARLI, MARALIXIBAT CHLORIDE

MISEMER

* MISEMER PHARMACEUTICALS INC
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
KETOPROFEN, KETOPROFEN

MISSION PHARMA

* MISSION PHARMACAL CO
LITHOSTAT, ACETOHYDROXAMIC ACID
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
TEXACORT, HYDROCORTISONE
THIOLA, TIOPRONIN
TINDAMAX, TINIDAZOLE
UROCIK-K, POTASSIUM CITRATE

MISSION PHARMACAL

* MISSION PHARMACAL CO
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
GABAPENTIN, GABAPENTIN
POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)
THIOLA EC, TIOPRONIN

mitsubishi tanabe

* MITSUBISHI TANABE PHARMA CORP
RADICAVA ORS, EDARAVONE
RADICAVA, EDARAVONE

MLV

* MLV PHARMA LLC
CARISOPRODOL, CARISOPRODOL
METHOCARBAMOL, METHOCARBAMOL
NYSTATIN, NYSTATIN
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

MMT

* MERIDIAN MEDICAL TECHNOLOGIES LLC
DUODOTE, ATROPINE
SEIZALAM, MIDAZOLAM HYDROCHLORIDE

MOBIUS THERAP

* MOBIUS THERAPEUTICS LLC
MITOSOL, MITOMYCIN

MOLNLYCKE HLTH

* MOLNLYCKE HEALTH CARE
HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)

MONARCH PHARMS

* MONARCH PHARMACEUTICALS LLC
MENESE, ESTROGENS, ESTERIFIED
NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
NEOSPORIN, GRAMICIDIN
SEPTRA DS, SULFAMETHOXAZOLE
SEPTRA, SULFAMETHOXAZOLE
VIROPTIC, TRIFLURIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MONTEREY PHARMS LLC**

* MONTEREY PHARMACEUTICALS LLC
METHOCARBAMOL, METHOCARBAMOL

MOUNTAIN

* MOUNTAIN LLC
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
METAXALONE, METAXALONE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

MSD MERCK CO

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
DELSTRIGO, DORAVIRINE
EMEND, APREPITANT
NOXAFIL POWDERMIX KIT, POSACONAZOLE
PIFELTRO, DORAVIRINE
RECARBRIO, CILASTATIN SODIUM

MSD SUB MERCK

* MERCK SHARP AND DOHME LLC A SUB OF MERCK AND CO INC
BRIDION, SUGAMMADEX SODIUM
INVANZ, ERTAPENEM SODIUM
ISENTRESS HD, RALTEGRAVIR POTASSIUM
ISENTRESS, RALTEGRAVIR POTASSIUM
JANUMET XR, METFORMIN HYDROCHLORIDE
JANUMET, METFORMIN HYDROCHLORIDE
SEGLUROMET, ERTUGLIFLOZIN
STEGLATRO, ERTUGLIFLOZIN
STEGLUJAN, ERTUGLIFLOZIN
ZEPATIER, ELBASVIR
ZOLINZA, VORINOSTAT

MSN

* MSN LABORATORIES PRIVATE LTD
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ACYCLOVIR, ACYCLOVIR
ALBENDAZOLE, ALBENDAZOLE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BORTEZOMIB, BORTEZOMIB
BUMETANIDE, BUMETANIDE
CAPECITABINE, CAPECITABINE
CARMUSTINE, CARMUSTINE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CLOBAZAM, CLOBAZAM
CLOFARABINE, CLOFARABINE
DARUNAVIR, DARUNAVIR
DECITABINE, DECITABINE
DEFERASIROX, DEFERASIROX
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DOFETILIDE, DOFETILIDE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
FEBUXOSTAT, FEBUXOSTAT
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
HALOPERIDOL, HALOPERIDOL
ISOSULFAN BLUE, ISOSULFAN BLUE
LACOSAMIDE, LACOSAMIDE
LEVETIRACETAM, LEVETIRACETAM
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
PACLITAXEL, PACLITAXEL
PIRFENIDONE, PIRFENIDONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MSN LABORATORIES PRIVATE LTD**

PLERIXAFOR, PLERIXAFOR
 PREGABALIN, PREGABALIN
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RIOCIQUAT, RIOCIQUAT
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SIROLIMUS, SIROLIMUS
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TASIMELTEON, TASIMELTEON
 TERIFLUNOMIDE, TERIFLUNOMIDE
 THIOTEPA, THIOTEPA
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 TOLVAPTAN, TOLVAPTAN
 TOREMIFENE CITRATE, TOREMIFENE CITRATE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZIPRASIDONE MESYLATE, ZIPRASIDONE MESYLATE

MSN LABS PVT LTD*** MSN LABORATORIES PRIVATE LTD**

LACOSAMIDE, LACOSAMIDE

MSN PHARMS INC*** MSN PHARMACEUTICALS INC**

DROXIDOPA, DROXIDOPA

MYCOVIA PHARMS*** MYCOVIA PHARMACEUTICALS INC**

VIVJOA, OTESECONAZOLE

MYLAN*** MYLAN INSTITUTIONAL LLC A VIATRIS CO**

ALOPRIM, ALLOPURINOL SODIUM

*** MYLAN LABORATORIES LTD A VIATRIS CO**

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

*** MYLAN PHARMACEUTICALS**

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

*** MYLAN PHARMACEUTICALS INC**

ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ACETAMINOPHEN, ACETAMINOPHEN
 ACITRETIN, ACITRETIN
 ALLOPURINOL, ALLOPURINOL
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ATENOLOL, ATENOLOL
 ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 BEPOTASTINE BESILATE, BEPOTASTINE BESILATE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BREYNA, BUDESONIDE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BUDESONIDE, BUDESONIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLORTHALIDONE, CHLORTHALIDONE
 CIMETIDINE, CIMETIDINE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
 CLOZAPINE, CLOZAPINE
 COLCHICINE, COLCHICINE
 CYCLOSPORINE, CYCLOSPORINE
 CYSTAGON, CYSTEAMINE BITARTRATE
 DAPSONE, DAPSONE
 DENAVIR, PENCICLOVIR
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ERMEZA, LEVOTHYROXINE SODIUM
 ERYGEL, ERYTHROMYCIN
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESTRADIOL, ESTRADIOL
 ETOPOSIDE, ETOPOSIDE
 EVEROLIMUS, EVEROLIMUS
 EXTINA, KETOCONAZOLE
 FENOFIBRATE, FENOFIBRATE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FUROSEMIDE, FUROSEMIDE
 GLATIRAMER ACETATE, GLATIRAMER ACETATE
 HALCINONIDE, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 IMATINIB MESYLATE, IMATINIB MESYLATE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LENALIDOMIDE, LENALIDOMIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIRTAZAPINE, MIRTAZAPINE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NEVIRAPINE, NEVIRAPINE
 NISOLDIPINE, NISOLDIPINE
 OLANZAPINE, OLANZAPINE
 ONDANSETRON, ONDANSETRON
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 PHENYTEK, PHENYTOIN SODIUM
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MYLAN PHARMACEUTICALS INC
 - REGADENOSON, REGADENOSON
 - ROFLUMILAST, ROFLUMILAST
 - RUFINAMIDE, RUFINAMIDE
 - SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 - SILDENAFIL CITRATE, SILDENAFIL CITRATE
 - SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 - SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 - SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - SPIRONOLACTONE, SPIRONOLACTONE
 - SUCRALFATE, SUCRALFATE
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - SUNITINIB MALATE, SUNITINIB MALATE
 - SYMFI LO, EFAVIRENZ
 - TACROLIMUS, TACROLIMUS
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 - TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 - TELMISARTAN, TELMISARTAN
 - TERIFLUNOMIDE, TERIFLUNOMIDE
 - TETRABENAZINE, TETRABENAZINE
 - THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 - TIMOLOL MALEATE, TIMOLOL MALEATE
 - TRAVOPROST, TRAVOPROST
 - TRETINOIN, TRETINOIN
 - TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 - VALSARTAN, VALSARTAN
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 - VUSION, MICONAZOLE NITRATE
 - WIXELA INHUB, FLUTICASONE PROPIONATE
 - ZONALON, DOXEPIN HYDROCHLORIDE
- * MYLAN PHARMACEUTICALS INC A VIATRIS CO
 - DAPTOMYCIN, DAPTOMYCIN

MYLAN ASI

- * MYLAN ASI LLC
 - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 - ADENOSINE, ADENOSINE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

MYLAN INSTITUTIONAL

- * MYLAN INSTITUTIONAL INC
 - BUSULFAN, BUSULFAN
- * MYLAN INSTITUTIONAL LLC
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - ARGATROBAN, ARGATROBAN
 - BIVALIRUDIN, BIVALIRUDIN
 - CIDOFOVIR, CIDOFOVIR
 - COSYNTROPIN, COSYNTROPIN
 - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - DURACLON, CLONIDINE HYDROCHLORIDE
 - ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 - ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 - FOMEPIZOLE, FOMEPIZOLE
 - IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 - ISOSULFAN BLUE, ISOSULFAN BLUE
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 - NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 - OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 - PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 - RIMSO-50, DIMETHYL SULFOXIDE
 - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 - SOTRADECOL, SODIUM TETRADECYL SULFATE
 - THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 - TRANEXAMIC ACID, TRANEXAMIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN INSTITUTIONAL LLC

ULTIVA, REMIFENTANIL HYDROCHLORIDE

MYLAN IRELAND LTD

* MYLAN IRELAND LTD

ARIXTRA, FONDAPARINUX SODIUM

MIACALCIN, CALCITONIN SALMON

PRETOMANID, PRETOMANID

YUPELRI, REVEFENACIN

MYLAN LABS LTD

* MYLAN LABORATORIES LTD

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE

ADENOSINE, ADENOSINE

BACLOFEN, BACLOFEN

CIMDUO, LAMIVUDINE

CLOFARABINE, CLOFARABINE

CYANOCOBALAMIN, CYANOCOBALAMIN

DAPTOMYCIN, DAPTOMYCIN

DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL

DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE

DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE

DOCETAXEL, DOCETAXEL

DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE

DOXYCYCLINE, DOXYCYCLINE HYCLATE

DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE

EPTIFIBATIDE, EPTIFIBATIDE

ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE

ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL

ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

ETOMIDATE, ETOMIDATE

FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE

FAMOTIDINE, FAMOTIDINE

FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE

FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE

GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE

HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE

HALOPERIDOL, HALOPERIDOL LACTATE

HEPARIN SODIUM, HEPARIN SODIUM

IBANDRONATE SODIUM, IBANDRONATE SODIUM

LAMIVUDINE, LAMIVUDINE

LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM

LEVETIRACETAM, LEVETIRACETAM

LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

LEVONORGESTREL, LEVONORGESTREL (OTC)

LINEZOLID, LINEZOLID

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE

MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE

MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE

MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN

MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE

NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

NORETHINDRONE ACETATE, NORETHINDRONE ACETATE

NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL

NORETHINDRONE, NORETHINDRONE

OXALIPLATIN, OXALIPLATIN

PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

POSACONAZOLE, POSACONAZOLE

PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE

RIFAMPIN, RIFAMPIN

ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE

SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

SYMFI, EFAVIRENZ

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MYLAN LABORATORIES LTD**

TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

MYLAN PHARMS INC*** MYLAN PHARMACEUTICALS INC**

ABACAVIR SULFATE, ABACAVIR SULFATE
 ACYCLOVIR, ACYCLOVIR
 AMNESTEEM, ISOTRETINOIN
 ARMODAFINIL, ARMODAFINIL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AVITA, TRETINOIN
 CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 ESZOPICLONE, ESZOPICLONE
 FENOFIBRATE, FENOFIBRATE
 LANSOPRAZOLE, LANSOPRAZOLE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NEVIRAPINE, NEVIRAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VORICONAZOLE, VORICONAZOLE
 ZIDOVUDINE, ZIDOVUDINE

*** MYLAN PHARMACEUTICALS INC.**

FLUVASTATIN SODIUM, FLUVASTATIN SODIUM

MYLAN SPCLT VIATRIS*** MYLAN SPECIALTY LP A VIATRIS CO**

DIPENTUM, OLSALAZINE SODIUM

MYLAN SPECIALITY LP*** MYLAN SPECIALTY LP**

COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 CORTIFOAM, HYDROCORTISONE ACETATE
 DEPEN, PENICILLAMINE
 DYMISTA, AZELASTINE HYDROCHLORIDE
 EDLUAR, ZOLPIDEM TARTRATE
 ELESTRIN, ESTRADIOL
 EPIFOAM, HYDROCORTISONE ACETATE
 EPIPEN JR., EPINEPHRINE
 EPIPEN, EPINEPHRINE
 FELBATOL, FELBAMATE
 GASTROCROM, CROMOLYN SODIUM
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 MUSE, ALPROSTADIL
 PROCTOFOAM HC, HYDROCORTISONE ACETATE
 ROWASA, MESALAMINE
 SFROWASA, MESALAMINE
 SOMA, CARISOPRODOL
 TOBI PODHALER, TOBRAMYCIN
 TOBI, TOBRAMYCIN

MYLAN SPECLT*** MYLAN SPECIALTY LP**

PERFORMIST, FORMOTEROL FUMARATE

MYLAN TECH VIATRIS*** MYLAN TECHNOLOGIES INC A VIATRIS CO**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN TECHNOLOGIES INC A VIATRIS CO
METHYLPHENIDATE, METHYLPHENIDATE

MYLAN TECHNOLOGIES

* MYLAN TECHNOLOGIES INC
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
CLONIDINE, CLONIDINE
ESTRADIOL, ESTRADIOL
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-37, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-62, FENTANYL
FENTANYL-75, FENTANYL
FENTANYL-87, FENTANYL
LIDOCAINE, LIDOCAINE
NITROGLYCERIN, NITROGLYCERIN
RIVASTIGMINE, RIVASTIGMINE
SCOPOLAMINE, SCOPOLAMINE
XULANE, ETHINYL ESTRADIOL

MYOVANT SCIENCES

* MYOVANT SCIENCES GMBH
MYFEMBREE, ESTRADIOL

**** N ******NAARI PTE LTD**

* NAARI PTE LTD
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LEVONORGESTREL, LEVONORGESTREL (OTC)
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NORETHINDRONE, NORETHINDRONE
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

NABRIVA

* NABRIVA THERAPEUTICS IRELAND DAC
XENLETA, LEFAMULIN ACETATE

NAL PHARM

* NAL PHARMACEUTICAL GROUP LTD
LIDOCAINE, LIDOCAINE

NALPROPION

* NALPROPION PHARMACEUTICALS LLC
CONTRAVE, BUPROPION HYDROCHLORIDE

NANG KUANG PHARM CO

* NANG KUANG PHARMACEUTICAL CO LTD
ICATIBANT ACETATE, ICATIBANT ACETATE
LINEZOLID, LINEZOLID
PEMETREXED DISODIUM, PEMETREXED DISODIUM

NANJING

* NANJING SIMCERE DONGYUAN PHARMACEUTICAL CO LTD
CELECOXIB, CELECOXIB

NANJING KING-FRIEND

* NANJING KING-FRIEND BIOCHEMICAL PHARMACEUTICAL CO LTD
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE

NANOCOPOEIA

* NANOCOPOEIA LLC
PHYRAGO, DASATINIB

NAPO PHARMS INC

* NAPO PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NAPO PHARMACEUTICALS INC
MYTESI, CROFELEMER

NATCO

* NATCO PHARMA LTD
ALPRAZOLAM, ALPRAZOLAM
CARISOPRODOL, CARISOPRODOL
GEFITINIB, GEFITINIB
GLYCOPYRROLATE, GLYCOPYRROLATE
LANSOPRAZOLE, LANSOPRAZOLE (OTC)
NITROGLYCERIN, NITROGLYCERIN
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
TERIFLUNOMIDE, TERIFLUNOMIDE
TIPIRACIL HYDROCHLORIDE AND TRIFLURIDINE, TIPIRACIL HYDROCHLORIDE

NATCO PHARMA

* NATCO PHARMA LTD
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
SUNITINIB MALATE, SUNITINIB MALATE

NATCO PHARMA LTD

* NATCO PHARMA LIMITED
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE

* NATCO PHARMA LTD
ANASTROZOLE, ANASTROZOLE
ARMODAFINIL, ARMODAFINIL
AZACITIDINE, AZACITIDINE
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
LANSOPRAZOLE, LANSOPRAZOLE
LANTHANUM CARBONATE, LANTHANUM CARBONATE
LAPATINIB DITOSYLATE, LAPATINIB DITOSYLATE
LETROZOLE, LETROZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

NAVINTA LLC

* NAVINTA LLC
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CARGLUMIC ACID, CARGLUMIC ACID
CARMUSTINE, CARMUSTINE
FAMOTIDINE, FAMOTIDINE
FOMEPIZOLE, FOMEPIZOLE
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
METHOCARBAMOL, METHOCARBAMOL
PENICILLAMINE, PENICILLAMINE
RIBAVIRIN, RIBAVIRIN
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

NCM USA BRONX LLC

* NCM USA BRONX LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

NE RX PHARMA

* NE RX PHARMA LLC
ACETAZOLAMIDE, ACETAZOLAMIDE
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
METOLAZONE, METOLAZONE

NEOS THERAPS

* NEOS THERAPEUTICS
ADZENYS XR-ODT, AMPHETAMINE

NEOS THERAPS INC

* NEOS THERAPEUTICS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NEOS THERAPEUTICS INC
COTEMPLA XR-ODT, METHYLPHENIDATE

NEPHRON

* NEPHRON CORP
ALBUTEROL SULFATE, ALBUTEROL SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

* NEPHRON PHARMACEUTICALS CORP
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
BUDESONIDE, BUDESONIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION

NEURELIS INC

* NEURELIS INC
VALTOCO, DIAZEPAM

NEUROCRINE

* NEUROCRINE BIOSCIENCES INC
INGREZZA, VALBENZAZINE TOSYLATE
ONGENTYS, OPICAPONE

NEXTWAVE

* NEXTWAVE PHARMACEUTICALS INC A SUB OF TRIS PHARMA INC
QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE

NEXTWAVE PHARMS

* NEXTWAVE PHARMACEUTICALS INC
QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE

NEXUS

* NEXUS PHARMACEUTICALS LLC
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
BUSULFAN, BUSULFAN
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
EMERPHED, EPHEDRINE SULFATE
ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
NELARABINE, NELARABINE
POTASSIUM CHLORIDE 10MEQ, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 40MEQ, POTASSIUM CHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
TIROFIBAN HYDROCHLORIDE, TIROFIBAN HYDROCHLORIDE

NEXUS PHARMS

* NEXUS PHARMACEUTICALS INC
FLUORESCEIN SODIUM, FLUORESCEIN SODIUM
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

NIAGARA PHARMS

* NIAGARA PHARMACEUTICALS INC
PUR-WASH, PURIFIED WATER (OTC)

NIPPON SHINYAKU

* NIPPON SHINYAKU CO LTD
VILTEPSO, VILTOLARSEN

NIVAGEN PHARMS INC

* NIVAGEN PHARMACEUTICALS INC
CALCIUM GLUCONATE, CALCIUM GLUCONATE
DECITABINE, DECITABINE
EFINACONAZOLE, EFINACONAZOLE
TEMOZOLOMIDE, TEMOZOLOMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NOBELPHARMA**

* NOBELPHARMA CO LTD
HYFTOR, SIROLIMUS

NODEN PHARMA

* NODEN PHARMA DAC
TEKTURNA, ALISKIREN HEMIFUMARATE

NORTEC DEV ASSOC

* NORTEC DEVELOPMENT ASSOC INC
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

NORTHLAND

* NORTHLAND NUCLEAR MEDICINE LLC
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

NORTHSTAR HLTHCARE

* NORTHSTAR HEALTHCARE HOLDINGS LTD
ALLOPURINOL, ALLOPURINOL
BACLOFEN, BACLOFEN
GEMFIBROZIL, GEMFIBROZIL
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

NORTON WATERFORD

* NORTON WATERFORD LTD
QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

NORWICH

* NORWICH PHARMACEUTICALS INC
LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE

NOSTRUM LABS INC

* NOSTRUM LABORATORIES INC
ACETAZOLAMIDE, ACETAZOLAMIDE
BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
CALCIUM ACETATE, CALCIUM ACETATE
CARBAMAZEPINE, CARBAMAZEPINE
CARISOPRODOL, CARISOPRODOL
CLARITHROMYCIN, CLARITHROMYCIN
DAPSONE, DAPSONE
ELIXOPHYLLIN, THEOPHYLLINE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
HYDROCODONE, HYDROCODONE BITARTRATE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NITROFURANTOIN, NITROFURANTOIN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
SUCRALFATE, SUCRALFATE
THEOPHYLLINE, THEOPHYLLINE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

NOSTRUM PHARMS LLC

* NOSTRUM PHARMACEUTICALS LLC
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
THEOCHRON, THEOPHYLLINE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

NOVA LABS LTD

* NOVA LABORATORIES LTD
PURIXAN, MERCAPTOPYRINE

NOVADAQ TECH

* NOVADAQ TECHNOLOGIES ULC
SPY AGENT GREEN KIT, INDOCYANINE GREEN

NOVARTIS

* NOVARTIS PHARMACEUTICALS CORP
AFINITOR, EVEROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVARTIS PHARMACEUTICALS CORP
 ALOMIDE, LODOXAMIDE TROMETHAMINE
 BETOPTIC S, BETAXOLOL HYDROCHLORIDE
 COARTEM, ARTEMETHER
 DESFERAL, DEFEROXAMINE MESYLATE
 DIOVAN HCT, HYDROCHLOROTHIAZIDE
 DIOVAN, VALSARTAN
 EGATEN, TRICLABENDAZOLE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXFORGE, AMLODIPINE BESYLATE
 EXJADE, DEFERASIROX
 FABHALTA, IPTACOPAN HYDROCHLORIDE
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GLEEVEC, IMATINIB MESYLATE
 JADENU SPRINKLE, DEFERASIROX
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KISQALI, RIBOCICLIB SUCCINATE
 LEQVIO, INCLISIRAN SODIUM
 LOCAMETZ, GALLIUM GA-68 GOZETOTIDE
 MAYZENT, SIPONIMOD
 MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
 MYFORTIC, MYCOPHENOLIC SODIUM
 NEORAL, CYCLOSPORINE
 PIQRAY, ALPELISIB
 PLUVICTO, LUTETIUM LU-177 VIPIVOTIDE TETRAKETAN
 PROMACTA KIT, ELTROMBOPAG OLAMINE
 PROMACTA, ELTROMBOPAG OLAMINE
 RYDAPT, MIDOSTAURIN
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SCEMBLIX, ASCIMINIB HYDROCHLORIDE
 TAFINLAR, DABRAFENIB MESYLATE
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TOBRADEX, DEXAMETHASONE
 TOBREX, TOBRAMYCIN
 TRILEPTAL, OXCARBAZEPINE
 TYKERB, LAPATINIB DITOSYLATE
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
 VIJOICE, ALPELISIB
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 XIIDRA, LIFITEGRAST
 ZORTRESS, EVEROLIMUS
 ZYKADIA, CERITINIB

NOVARTIS PHARM

* NOVARTIS PHARMACEUTICAL CORP
 AFINITOR DISPERZ, EVEROLIMUS
 TABRECTA, CAPMATINIB HYDROCHLORIDE

NOVARTIS PHARMS

* NOVARTIS PHARMACEUTICALS CORP
 FEMARA, LETROZOLE

NOVARTIS PHARMS CORP

* NOVARTIS PHARMACEUTICALS CORP
 ENTRESTO, SACUBITRIL
 JADENU, DEFERASIROX

NOVAST LABS

* NOVAST LABORATORIES CHINA LTD
 NORETHINDRONE, NORETHINDRONE
 * NOVAST LABORATORIES INC
 DOCETAXEL, DOCETAXEL
 * NOVAST LABORATORIES LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVAST LABORATORIES LTD
 CARBOPLATIN, CARBOPLATIN
 CARISOPRODOL, CARISOPRODOL
 CHABELINA FE, ETHINYL ESTRADIOL
 CHLORTHALIDONE, CHLORTHALIDONE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DECITABINE, DECITABINE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOLISHALE, ETHINYL ESTRADIOL
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HER STYLE, LEVONORGESTREL (OTC)
 INDOMETHACIN, INDOMETHACIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LARIN 1.5/30, ETHINYL ESTRADIOL
 LARIN 1/20, ETHINYL ESTRADIOL
 LARIN 24 FE, ETHINYL ESTRADIOL
 LARIN FE 1.5/30, ETHINYL ESTRADIOL
 LARIN FE 1/20, ETHINYL ESTRADIOL
 LERIBANE, ETHINYL ESTRADIOL
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LO-MALMOREDE, ETHINYL ESTRADIOL
 MALMOREDE, ETHINYL ESTRADIOL
 MELAMISA, DROSPIRENONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 NIFEDIPINE, NIFEDIPINE
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE, NORETHINDRONE
 OXALIPLATIN, OXALIPLATIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PIMTREA, DESOGESTREL
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
 PROBENECID AND COLCHICINE, COLCHICINE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RANOLAZINE, RANOLAZINE
 SETLAKIN, ETHINYL ESTRADIOL
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLCAPONE, TOLCAPONE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRI-LO-LINYAH, ETHINYL ESTRADIOL
 TRIAZOLAM, TRIAZOLAM
 YAELA, DROSPIRENONE

NOVAST LABS LTD

* NOVAST LABORATORIES LTD
 DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL
 ELINEST, ETHINYL ESTRADIOL
 FALMINA, ETHINYL ESTRADIOL
 LEVONEST, ETHINYL ESTRADIOL
 MONO-LINYAH, ETHINYL ESTRADIOL
 PHILITH, ETHINYL ESTRADIOL
 TRI-LINYAH, ETHINYL ESTRADIOL
 WERA, ETHINYL ESTRADIOL

NOVATECH SA

* NOVATECH SA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVATECH SA
STERITALC, TALC

NOVEL LABS INC

* NOVEL LABORATORIES INC
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
CARBIDOPA, CARBIDOPA
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DESOXIMETASONE, DESOXIMETASONE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIAZEPAM, DIAZEPAM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
FAMOTIDINE, FAMOTIDINE
FLUCYTOSINE, FLUCYTOSINE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
LEVONORGESTREL, LEVONORGESTREL (OTC)
LINEZOLID, LINEZOLID
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC,
MORPHINE SULFATE, MORPHINE SULFATE
NITROFURANTOIN, NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND
PHENELZINE SULFATE, PHENELZINE SULFATE
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
TEMAZEPAM, TEMAZEPAM
TRIMETHOPRIM, TRIMETHOPRIM
VORICONAZOLE, VORICONAZOLE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

NOVELGENIX THERAPS

* NOVELGENIX THERAPEUTICS PVT LTD
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

NOVEN

* NOVEN PHARMACEUTICALS INC
MINIVELLE, ESTRADIOL

NOVEN PHARMS INC

* NOVEN PHARMACEUTICALS INC
COMBIPATCH, ESTRADIOL
DAYTRANA, METHYLPHENIDATE
XELSTRYM, DEXTROAMPHETAMINE

NOVITIUM PHARMA

* NOVITIUM PHARMA LLC
ACETAZOLAMIDE, ACETAZOLAMIDE
ACYCLOVIR, ACYCLOVIR
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALPRAZOLAM, ALPRAZOLAM
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
BETAINE, BETAINE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
CARBAMAZEPINE, CARBAMAZEPINE
CARGLUMIC ACID, CARGLUMIC ACID
CHLORZOXAZONE, CHLORZOXAZONE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVITIUM PHARMA LLC
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DAPSONE, DAPSONE
 DEXAMETHASONE, DEXAMETHASONE
 DIAZOXIDE, DIAZOXIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 ESTAZOLAM, ESTAZOLAM
 ESTRADIOL, ESTRADIOL
 FAMOTIDINE, FAMOTIDINE
 FELBAMATE, FELBAMATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 LEVOCARNITINE SF, LEVOCARNITINE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MELOXICAM, MELOXICAM
 METHSUXIMIDE, METHSUXIMIDE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 NAPROXEN, NAPROXEN
 NITISINONE, NITISINONE
 NITROFURANTOIN, NITROFURANTOIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RIFABUTIN, RIFABUTIN
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SIROLIMUS, SIROLIMUS
 SOVUNA, HYDROXYCHLOROQUINE SULFATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TADALAFIL, TADALAFIL
 THIOTHIXENE, THIOTHIXENE
 TRANILCYPROMINE SULFATE, TRANILCYPROMINE SULFATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 VALSARTAN, VALSARTAN

NOVO

* NOVO NORDISK INC
 OZEMPIC, SEMAGLUTIDE
 RIVFLOZA, NEDOSIRAN SODIUM
 RYBELSUS, SEMAGLUTIDE
 SAXENDA, LIRAGLUTIDE RECOMBINANT
 WEGOVY, SEMAGLUTIDE

NOVO NORDISK

* NOVO NORDISK PHARMACEUTICALS INC
 GLUCAGEN, GLUCAGON HYDROCHLORIDE

NOVO NORDISK INC

* NOVO NORDISK INC
 VAGIFEM, ESTRADIOL
 VICTOZA, LIRAGLUTIDE RECOMBINANT

NOVUGEN

* NOVUGEN ONCOLOGY SDN BHD
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 SUNITINIB MALATE, SUNITINIB MALATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVUGEN PHARMA MALAYSIA SDN BHD
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE

NUKEMED

* NUKEMED INC DBA SPECTRONRX
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

NUVO PHARM

* NUVO PHARMACEUTICAL INC
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIAZEPAM, DIAZEPAM
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

NUVO PHARMS INC

* NUVO PHAMACEUTICALS INC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CHLORDIAZEPOXIDE HYDROCHLORIDE AND CLIDINIUM BROMIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
NAPROXEN, NAPROXEN
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SULFASALAZINE, SULFASALAZINE

NXDC

* NX DEVELOPMENT CORP
GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE

**** O ******OCULAR THERAPEUTIX**

* OCULAR THERAPEUTIX INC
DEXTENZA, DEXAMETHASONE

OCUPHIRE

* OCUPHIRE PHARMA INC
RYZUMVI, PHENTOLAMINE MESYLATE

OHM LABS

* OHM LABORATORIES INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

OHM LABS INC

* OHM LABORATORIES INC
EZETIMIBE, EZETIMIBE
GUAIFENESIN, GUAIFENESIN (OTC)
VALSARTAN, VALSARTAN

OMNIVIUM PHARMS

* OMNIVIUM PHARMACEUTICALS LLC
ISONIAZID, ISONIAZID
NUMBRINO, COCAINE HYDROCHLORIDE
SODIUM BICARBONATE, SODIUM BICARBONATE

OMSAV PHARMA

* OMSAV PHARMA RESEARCH PRIVATE LTD
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN

ON TARGET LABS

* ON TARGET LABORATORIES INC
CYTALUX, PAFOLACIANINE SODIUM

ONYX PHARMS AMGEN

* ONYX PHARMACEUTICALS INC A WHOLLY OWNED SUB OF AMGEN INC
KYPROLIS, CARFILZOMIB

OPERAND PHARMS

* OPERAND PHARMACEUTICALS III LTD
EPHEDRINE SULFATE, EPHEDRINE SULFATE

OPTIMUS

* OPTIMUS PHARMA PRIVATE LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* OPTIMUS PHARMA PRIVATE LTD
AMINOCAPROIC ACID, AMINOCAPROIC ACID
BREXPIRAZOLE, BREXPIRAZOLE

OPTINOSE US INC

* OPTINOSE US INC
XHANCE, FLUTICASONE PROPIONATE

ORAPHARMA

* ORAPHARMA INC
ARESTIN, MINOCYCLINE HYDROCHLORIDE

ORASIS PHARMS

* ORASIS PHARMACEUTICALS LTD
QLOSI, PILOCARPINE HYDROCHLORIDE

ORBICULAR

* ORBICULAR PHARMACEUTICAL TECHNOLOGIES PRIVATE LTD
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE

ORBION PHARMS

* ORBION PHARMACEUTICALS PRIVATE LTD
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ARIPIRAZOLE, ARIPIRAZOLE
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
DESLORATADINE, DESLORATADINE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
ESZOPICLONE, ESZOPICLONE
FELODIPINE, FELODIPINE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
IBANDRONATE SODIUM, IBANDRONATE SODIUM
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN, LEVOFLOXACIN
MODAFINIL, MODAFINIL
NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
OLANZAPINE, OLANZAPINE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
RASAGILINE MESYLATE, RASAGILINE MESYLATE
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
ZALEPLON, ZALEPLON
ZOLMITRIPTAN, ZOLMITRIPTAN

OREXO US INC

* OREXO US INC
ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE

ORGANON

* ORGANON LLC A SUB OF ORGANON AND CO
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CLARINEX, DESLORATADINE
COZAAR, LOSARTAN POTASSIUM
DIPROLENE, BETAMETHASONE DIPROPIONATE
FOSAMAX, ALENDRONATE SODIUM
HYZAAR, HYDROCHLOROTHIAZIDE
MAXALT-MLT, RIZATRIPTAN BENZOATE
PROPECIA, FINASTERIDE
PROSCAR, FINASTERIDE
SINEMET, CARBIDOPA
SINGULAIR, MONTELUKAST SODIUM
VYTORIN, EZETIMIBE
ZETIA, EZETIMIBE
ZOCOR, SIMVASTATIN

* ORGANON USA LLC
NEXPLANON, ETNOGESTREL
REMERON, MIRTAZAPINE

ORGANON LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

*** ORGANON LLC**

ASMANEX HFA, MOMETASONE FUROATE
 ASMANEX TWISTHALER, MOMETASONE FUROATE
 CLARINEX-D 12 HOUR, DESLORATADINE
 DULERA, FORMOTEROL FUMARATE
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 MAXALT, RIZATRIPTAN BENZOATE

ORGANON USA ORGANON

* ORGANON USA LLC A SUB OF ORGANON AND CO
 GANIRELIX ACETATE, GANIRELIX ACETATE
 NUVARING, ETHINYL ESTRADIOL
 REMERON SOLTAB, MIRTAZAPINE

ORIENT PHARMA

* ORIENT PHARMA CO LTD
 EZETIMIBE, EZETIMIBE

ORIENT PHARMA CO LTD

* ORIENT PHARMA CO LTD
 CARISOPRODOL, CARISOPRODOL
 GLYBURIDE, GLYBURIDE
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ORION PHARMA

* ORION PHARMA
 COMTAN, ENTACAPONE
 STALEVO 100, CARBIDOPA
 STALEVO 125, CARBIDOPA
 STALEVO 150, CARBIDOPA
 STALEVO 200, CARBIDOPA
 STALEVO 50, CARBIDOPA
 STALEVO 75, CARBIDOPA

ORPHALAN

* ORPHALAN SA
 CUVRIOR, TRIENTINE TETRAHYDROCHLORIDE

OSMOTICA PHARM US

* OSMOTICA PHARMACEUTICAL US LLC
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NIFEDIPINE, NIFEDIPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 RELEXII, METHYLPHENIDATE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

OTSUKA

* OTSUKA PHARMACEUTICAL CO LTD
 ABILIFY ASIMTUFII, ARIPIPRAZOLE
 ABILIFY MYCITE KIT, ARIPIPRAZOLE
 ABILIFY, ARIPIPRAZOLE
 INQOVI, CEDAZURIDINE
 JYNARQUE, TOLVAPTAN
 REXULTI, BREXPIPRAZOLE
 SAMSCA, TOLVAPTAN

OTSUKA PHARM

* OTSUKA PHARMACEUTICAL CO LTD
 BUSULFEX, BUSULFAN

OTSUKA PHARM CO LTD

* OTSUKA PHARMACEUTICAL CO LTD
 ABILIFY MAINTENA KIT, ARIPIPRAZOLE

OTTER PHARMS

* OTTER PHARMACEUTICALS LLC
 OTREXUP, METHOTREXATE
 SYMPAZAN, CLOBAZAM

OVERSEAS

* OVERSEAS PHARMACEUTICALS LTD
 LEVETIRACETAM, LEVETIRACETAM

OXFORD PHARMS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** O ****

* OXFORD PHARMACEUTICALS LLC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BACLOFEN, BACLOFEN
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARISOPRODOL, CARISOPRODOL
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LORAZEPAM, LORAZEPAM
 METHOCARBAMOL, METHOCARBAMOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE
 RIMACTANE, RIFAMPIN
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

OYSTER POINT PHARMA

* OYSTER POINT PHARMA INC
 TYRVAYA, VARENICLINE TARTRATE

**** P ******P AND L**

* P AND L DEVELOPMENT LLC
 ADAPALENE, ADAPALENE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 DOCOSANOL, DOCOSANOL (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

P AND L DEV LLC

* P AND L DEVELOPMENT LLC DBA PLD DEVELOPMENTS LLC
 IBUPROFEN, IBUPROFEN (OTC)

PACIFIC PHARMA

* PACIFIC PHARMA
 TIMOLOL MALEATE, TIMOLOL MALEATE
 * PACIFIC PHARMA INC
 TIMOLOL MALEATE, TIMOLOL MALEATE

PACIRA PHARMS INC

* PACIRA PHARMACEUTICALS INC
 EXPAREL, BUPIVACAINE
 ZILRETTA, TRIAMCINOLONE ACETONIDE

PADAGIS ISRAEL

* PADAGIS ISRAEL PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PADAGIS ISRAEL PHARMACEUTICALS LTD
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BUDESONIDE, BUDESONIDE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE AND BETHAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DAPSONE, DAPSONE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ESTRADIOL, ESTRADIOL
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLURANDRENOLIDE, FLURANDRENOLIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GYNAZOLE-1, BUTOCONAZOLE NITRATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 IMIQUIMOD, IMIQUIMOD
 IVERMECTIN, IVERMECTIN
 KETOCONAZOLE, KETOCONAZOLE
 MESALAMINE, MESALAMINE
 METRONIDAZOLE, METRONIDAZOLE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE (OTC)
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PERMETHRIN, PERMETHRIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SUMATRIPTAN, SUMATRIPTAN
 TAZAROTENE, TAZAROTENE
 TERCONAZOLE, TERCONAZOLE
 TESTOSTERONE, TESTOSTERONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

PADAGIS US

* PADAGIS US LLC
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN, BACITRACIN
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BRINZOLAMIDE, BRINZOLAMIDE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CALCIUM ACETATE, CALCIUM ACETATE
 CENTANY, MUPIROCIN
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CICLOPIROX, CICLOPIROX
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLINDESSE, CLINDAMYCIN PHOSPHATE
 CLINDETS, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COMPRO, PROCHLORPERAZINE
 CYCLOSPORINE, CYCLOSPORINE
 DESONIDE, DESONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PADAGIS US LLC
 DESOXIMETASONE, DESOXIMETASONE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 ENTOCORT EC, BUDESONIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 EVAMIST, ESTRADIOL
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 HYDROCORTISONE, HYDROCORTISONE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NYSTATIN, NYSTATIN
 NYSTOP, NYSTATIN
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PODOFILOX, PODOFILOX
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 REPAGLINIDE, REPAGLINIDE
 SCOPOLAMINE, SCOPOLAMINE
 SELENIUM SULFIDE, SELENIUM SULFIDE
 STIE-CORT, HYDROCORTISONE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TAVABOROLE, TAVABOROLE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE

PAI HOLDINGS PHARM

* PAI HOLDINGS LLC DBA PHARMACEUTICAL ASSOCIATES INC
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LACTULOSE, LACTULOSE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LITHIUM CITRATE, LITHIUM CITRATE
 PHENYTOIN, PHENYTOIN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

PALATIN TECHNOLOGIES

* PALATIN TECHNOLOGIES INC
 VYLEESI (AUTOINJECTOR), BREMELANOTIDE ACETATE

PANACEA

* PANACEA BIOTEC PHARMA LTD
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TACROLIMUS, TACROLIMUS

PAR FORM

* PAR FORMULATIONS PRIVATE LTD
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

PAR PHARM

* PAR PHARMACEUTICAL
 EVEROLIMUS, EVEROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PAR PHARMACEUTICAL INC
 ALPRAZOLAM, ALPRAZOLAM
 ALVIMOPAN, ALVIMOPAN
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLONAZEPAM, CLONAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROXYUREA, HYDROXYUREA
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LAMOTRIGINE, LAMOTRIGINE
 MINOXIDIL, MINOXIDIL
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 PIMOZIDE, PIMOZIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TOPIRAMATE, TOPIRAMATE
 URSODIOL, URSODIOL

PAR PHARM INC

* PAR PHARMACEUTICAL INC
 AMBRISENTAN, AMBRISENTAN
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 CHLORZOXAZONE, CHLORZOXAZONE
 COLCHICINE, COLCHICINE
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 PENICILLAMINE, PENICILLAMINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PRAZIQUANTEL, PRAZIQUANTEL
 SAPROPTERIN DIHYDROCHLORIDE, SAPROPTERIN DIHYDROCHLORIDE
 TIOPRONIN, TIOPRONIN
 TOLCAPONE, TOLCAPONE
 TOLVAPTAN, TOLVAPTAN
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VARENICLINE TARTRATE, VARENICLINE TARTRATE
 VIGABATRIN, VIGABATRIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

PAR STERILE PRODUCTS

* PAR STERILE PRODUCTS LLC
 ADRENALIN, EPINEPHRINE
 ARGATROBAN, ARGATROBAN
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CALCITONIN-SALMON, CALCITONIN SALMON
 COLY-MYCIN M, COLISTIMETHATE SODIUM
 CORPHEDRA, EPHEDRINE SULFATE
 DANTRIUM, DANTROLENE SODIUM
 DELESTROGEN, ESTRADIOL VALERATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PAR STERILE PRODUCTS LLC
 KETALAR, KETAMINE HYDROCHLORIDE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 MICAfungin, MICAfungin SODIUM
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PITOCIN, OXYTOCIN
 POSACONAZOLE, POSACONAZOLE
 TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TREPROSTINIL, TREPROSTINIL
 VASOSTRICT, VASOPRESSIN

PARAGON BIOTECK

* PARAGON BIOTECK INC
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

PARAPRO LLC

* PARAPRO LLC
 NATROBA, SPINOSAD

PARATEK PHARMS INC

* PARATEK PHARMACEUTICALS INC
 NUZYRA, OMADACYCLINE TOSYLATE

PARKE DAVIS

* PARKE DAVIS DIV WARNER LAMBERT CO
 CELONTIN, METHSUXIMIDE
 CEREBYX, FOSPHENYTOIN SODIUM
 NARDIL, PHENELZINE SULFATE
 ZARONTIN, ETHOSUXIMIDE

PARKE-DAVIS

* PARKE-DAVIS DIVISION OF PFIZER INC
 ZARONTIN, ETHOSUXIMIDE

PATHEON SOFTGELS

* PATHEON SOFTGELS BV
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

PATRIN

* PATRIN PHARMA INC
 CALCITRIOL, CALCITRIOL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 FLAC, FLUOCINOLONE ACETONIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 PREGABALIN, PREGABALIN

PENN LIFE

* PENN LIFE SCIENCES LLC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CARMUSTINE, CARMUSTINE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 THIOTEPA, THIOTEPA

PERRIGO NEW YORK

* PERRIGO NEW YORK INC
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 PERMETHRIN, PERMETHRIN (OTC)

PERRIGO PHARMA INTL

* PERRIGO PHARMA INTERNATIONAL DAC
 DICLOFENAC SODIUM, DICLOFENAC SODIUM (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 MINOXIDIL, MINOXIDIL (OTC)
 NASONEX 24HR ALLERGY, MOMETASONE FUROATE (OTC)
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
 * PERRIGO PHARMA INTERNATIONAL DESIGNATED ACTIVITY CO
 LORATADINE, LORATADINE (OTC)
 PREVACID 24 HR, LANSOPRAZOLE (OTC)

PERRIGO R AND D

* PERRIGO R AND D CO
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******* PERRIGO R AND D CO**

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)

PETNET*** PETNET SOLUTIONS INC**

AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

PF PRISM CV*** PF PRISM CV**

BOSULIF, BOSUTINIB MONOHYDRATE
 INLYTA, AXITINIB
 PRISTIQ, DESVENLAFAXINE SUCCINATE
 RAPAMUNE, SIROLIMUS
 TORISEL, TEMSIROLIMUS
 TYGACIL, TIGECYCLINE
 VFEND, VORICONAZOLE
 XALKORI, CRIZOTINIB
 XELJANZ, TOFACITINIB CITRATE

PFIZER*** PFIZER CENTRAL RESEARCH**

DIFLUCAN, FLUCONAZOLE
 ZITHROMAX, AZITHROMYCIN

*** PFIZER CHEMICALS DIV PFIZER INC**

DIFLUCAN, FLUCONAZOLE
 ZITHROMAX, AZITHROMYCIN

*** PFIZER INC**

ALDACTAZIDE, HYDROCHLOROTHIAZIDE
 ALDACTONE, SPIRONOLACTONE
 AROMASIN, EXEMESTANE
 ARTHROTEC, DICLOFENAC SODIUM
 AZULFIDINE EN-TABS, SULFASALAZINE
 AZULFIDINE, SULFASALAZINE
 CAVERJECT IMPULSE, ALPROSTADIL
 CAVERJECT, ALPROSTADIL
 CIBINQO, ABROCITINIB
 CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLEOCIN T, CLINDAMYCIN PHOSPHATE
 CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLEOCIN, CLINDAMYCIN PHOSPHATE
 COLESTID, COLESTIPOL HYDROCHLORIDE
 CORVERT, IBUTILIDE FUMARATE
 CYKLOKAPRON, TRANEXAMIC ACID
 CYTOTEC, MISOPROSTOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

- * PFIZER INC
 DAURISMO, GLASDEGIB MALEATE
 DAYPRO, OXAPROZIN
 DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
 DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ESTRING, ESTRADIOL
 FLAGYL, METRONIDAZOLE
 FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
 FRAGMIN, DALTEPARIN SODIUM
 GLUCOTROL XL, GLIPIZIDE
 GLYNASE, GLYBURIDE
 GLYSET, MIGLITOL
 HALCION, TRIAZOLAM
 HEMABATE, CARBOPROST TROMETHAMINE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 IBRANCE, PALBOCICLIB
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 LITFULO, RITLECITINIB TOSYLATE
 LOMOTIL, ATROPINE SULFATE
 LORBRENA, LORLATINIB
 MEDROL, METHYLPREDNISOLONE
 MERREM, MEROPENEM
 MYCOBUTIN, RIFABUTIN
 NICOTROL, NICOTINE
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 NURTEC ODT, RIMEGEPANT SULFATE
 OGEN 5, ESTROPIPATE
 PAXLOVID (COPACKAGED), NIRMATRELVIR
 PREPIDIL, DINOPROSTONE
 PROCARDIA, NIFEDIPINE
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROVERA, MEDROXYPROGESTERONE ACETATE
 SONATA, ZALEPLON
 SYNAREL, NAFARELIN ACETATE
 TALZENNA, TALAZOPARIB TOSYLATE
 TESSALON, BENZONATATE
 TOVIAZ, FESOTERODINE FUMARATE
 UNASYN, AMPICILLIN SODIUM
 VELSIPITY, ETRASIMOD ARGININE
 VIZIMPRO, DACOMITINIB
 XELJANZ XR, TOFACITINIB CITRATE
 XELJANZ, TOFACITINIB CITRATE
 ZAVZPRET, ZAVEGEPANT HYDROCHLORIDE
 ZITHROMAX, AZITHROMYCIN
 ZYVOX, LINEZOLID
- * PFIZER LABORATORIES DIV PFIZER INC
 FELDENE, PIROXICAM
 MINIPRESS, PRAZOSIN HYDROCHLORIDE
 PFIZERPEN, PENICILLIN G POTASSIUM
 PROCARDIA XL, NIFEDIPINE
 VIBRAMYCIN, DOXYCYCLINE
 VIBRAMYCIN, DOXYCYCLINE CALCIUM
 VIBRAMYCIN, DOXYCYCLINE HYCLATE
 VISTARIL, HYDROXYZINE PAMOATE
- * PFIZER PHARMACEUTICALS PRODUCTION CORP LTD
 TIKOSYN, DOFETILIDE

PFIZER INC

- * PFIZER INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PFIZER INC
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 ELLENCE, EPIRUBICIN HYDROCHLORIDE
 NICOTROL, NICOTINE

PFIZER PHARMS

* PFIZER PHARMACEUTICALS LTD
 LOPID, GEMFIBROZIL

PHARM ASSOC

* PHARMACEUTICAL ASSOC INC
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LORAZEPAM, LORAZEPAM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE

* PHARMACEUTICAL ASSOCIATES INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LACTULOSE, LACTULOSE
 LEVETIRACETAM, LEVETIRACETAM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISOLONE, PREDNISOLONE
 PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 THEOPHYLLINE, THEOPHYLLINE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

PHARM SOURCING

* PHARMACEUTICAL SOURCING PARTNERS INC
 MESALAMINE, MESALAMINE

PHARMA RES SOFTWARE

* PHARMA RESEARCH SOFTWARE SOLUTION LLC
 POKONZA, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

PHARMACHEMIE BV

* PHARMACHEMIE BV
 CARBOPLATIN, CARBOPLATIN
 CISPLATIN, CISPLATIN
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

PHARMACIA

* PHARMACIA AND UPJOHN CO LLC
 CADUET, AMLODIPINE BESYLATE
 DILANTIN, PHENYTOIN

PHARMACIA AND UPJOHN

* PHARMACIA AND UPJOHN CO

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PHARMACIA AND UPJOHN CO
CORTEF, HYDROCORTISONE
EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
R-GENE 10, ARGININE HYDROCHLORIDE
SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE

PHARMACOSMOS AS

* PHARMACOSMOS AS
MONOFERRIC, FERRIC DERISOMALTOSE

PHARMACYCLICS LLC

* PHARMACYCLICS LLC
IMBRUVICA, IBRUTINIB

PHARMADAX INC

* PHARMADAX INC
LEVETIRACETAM, LEVETIRACETAM
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

PHARMALOGIC

* PHARMALOGIC SOUTH CAROLINA LLC
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

PHARMALOGIC HLDGS

* PHARMALOGIC HOLDINGS CORP
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

PHARMASCIENCE INC

* PHARMASCIENCE INC
BORTEZOMIB, BORTEZOMIB
BUSULFAN, BUSULFAN
DECITABINE, DECITABINE
GANCICLOVIR SODIUM, GANCICLOVIR SODIUM

PHARMING

* PHARMING TECHNOLOGIES BV
JOENJA, LENIOLISIB PHOSPHATE

PHATHOM

* PHATHOM PHARMACEUTICALS INC
VOQUEZNA DUAL PAK, AMOXICILLIN
VOQUEZNA TRIPLE PAK, AMOXICILLIN
VOQUEZNA, VONOPRAZAN FUMARATE

PHOTOCURE ASA

* PHOTOCURE ASA
CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE

PIERRE FABRE DERMA

* PIERRE FABRE DERMATOLOGIE
HEMANGEOL, PROPRANOLOL HYDROCHLORIDE

PIERREL

* PIERREL S.P.A.
ORABLOC, ARTICAINA HYDROCHLORIDE

PINNACLE BIOLGS

* PINNACLE BIOLOGICS INC
PHOTOFRIN, PORFIMER SODIUM

PIRAMAL CRITICAL

* PIRAMAL CRITICAL CARE INC
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
GABLOFEN, BACLOFEN
GLYCOPYRROLATE, GLYCOPYRROLATE
ISOFLURANE, ISOFLURANE
MITIGO, MORPHINE SULFATE
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
SOJOURN, SEVOFLURANE
* PIRAMAL CRITICAL CARE LTD
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM

PIRAMAL HLTHCARE UK

* PIRAMAL HEALTHCARE UK LTD
CLOBAZAM, CLOBAZAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PIRAMAL HEALTHCARE UK LTD
 DEFERASIROX, DEFERASIROX
 EPLERENONE, EPLERENONE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 RANOLAZINE, RANOLAZINE
 TETRABENAZINE, TETRABENAZINE

PIRAMAL PHARMA

* PIRAMAL PHARMA LTD
 ISOFLURANE, ISOFLURANE

PLD ACQUISITIONS

* PLD ACQUISITIONS LLC DBA AVEMA PHARMA SOLUTIONS
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)

PLD ACQUISITIONS LLC

* PLD ACQUISITIONS LLC
 LORATADINE, LORATADINE (OTC)
 ZOLMITRIPTAN, ZOLMITRIPTAN

PLIVA

* PLIVA INC
 AZITHROMYCIN, AZITHROMYCIN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

PLIVA PHARM IND

* PLIVA PHARMACEUTICAL INDUSTRY INC
 TORSEMIDE, TORSEMIDE

PLX PHARMA

* PLX PHARMA INC
 VAZALORE, ASPIRIN (OTC)

POHL BOSKAMP

* POHL BOSKAMP
 NITROLINGUAL PUMPSPRAY, NITROGLYCERIN

POLAREAN

* POLAREAN INC
 XENOVUE, XENON XE-129 HYPERPOLARIZED

POLYGEN PHARMS

* POLYGEN PHARMACEUTICALS INC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

POWDER PHARMS

* POWDER PHARMACEUTICALS INC
 ZINGO, LIDOCAINE HYDROCHLORIDE

PRASCO

* PRASCO LLC DBA PRASCO LABORATORIES
 DEXAMETHASONE, DEXAMETHASONE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 EPLERENONE, EPLERENONE
 ESTRADIOL, ESTRADIOL
 MIRTAZAPINE, MIRTAZAPINE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

PRAXGEN

* PRAXGEN PHARMACEUTICALS LLC
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 RANOLAZINE, RANOLAZINE

PRECISION DERMAT

* PRECISION DERMATOLOGY INC
 LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
 LOCOID, HYDROCORTISONE BUTYRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

PRECISION DOSE INC

* PRECISION DOSE INC
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE

PRECISION NUCLEAR

* PRECISION NUCLEAR LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

PRIMUS PHARMS

* PRIMUS PHARMACEUTICALS INC
IMPOYZ, CLOBETASOL PROPIONATE
SERNIVO, BETAMETHASONE DIPROPIONATE

PRINSTON INC

* PRINSTON PHARMACEUTICAL INC
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
CAPTOPRIL, CAPTOPRIL
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ENTECAVIR, ENTECAVIR
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
FEBUXOSTAT, FEBUXOSTAT
FENOFIBRATE, FENOFIBRATE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
FUROSEMIDE, FUROSEMIDE
GLIMEPIRIDE, GLIMEPIRIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
LEVETIRACETAM, LEVETIRACETAM
LEVOMILNACIPRAN HYDROCHLORIDE, LEVOMILNACIPRAN HYDROCHLORIDE
LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PAROXETINE MESYLATE, PAROXETINE MESYLATE
PAROXETINE, PAROXETINE HYDROCHLORIDE
PEMETREXED DISODIUM, PEMETREXED DISODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PRINSTON PHARMACEUTICAL INC
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREGABALIN, PREGABALIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROFLUMILAST, ROFLUMILAST
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SILODOSIN, SILODOSIN
 TADALAFIL, TADALAFIL
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VORICONAZOLE, VORICONAZOLE
 VYDUO, NEBIVOLOL HYDROCHLORIDE

PROF DSPLS

* PROFESSIONAL DISPOSABLES INC
 PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)

PROGENICS PHARMS INC

* PROGENICS PHARMACEUTICALS INC
 AZEDRA, IOBENGUANE I-131
 PYLARIFY, PIFLUFOLASTAT F-18

PROPEL PHARMA

* PROPEL PHARMA CORP
 VIGABATRIN, VIGABATRIN

PROTEGA PHARMS

* PROTEGA PHARMACEUTICALS INC
 ROXYBOND, OXYCODONE HYDROCHLORIDE

PROVELL

* PROVELL PHARMACEUTICALS LLC
 EUTHYROX, LEVOTHYROXINE SODIUM **

PROVENSIS

* PROVENSIS LTD
 VARITHENA, POLIDOCANOL

PROVEPHARM SAS

* PROVEPHARM SAS
 BAL, DIMERCAPROL
 BLUDIGO, INDIGOTINDISULFONATE SODIUM
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROVAYBLUE, METHYLENE BLUE
 TRANEXAMIC ACID, TRANEXAMIC ACID

PTC THERAP

* PTC THERAPEUTICS INC
 EMFLAZA, DEFLAZACORT

PULMOFLOW INC

* PULMOFLOW INC
 KITABIS PAK, TOBRAMYCIN

PUMA BIOTECH

* PUMA BIOTECHNOLOGY INC
 NERLYNX, NERATINIB MALEATE

PURACAP PHARM

* PURACAP PHARMACEUTICAL LLC
 MELOXICAM, MELOXICAM

PURACAP PHARM LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PURACAP PHARMACEUTICAL LLC
 BENZONATATE, BENZONATATE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 ETHOSUXIMIDE, ETHOSUXIMIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

PURDUE PHARMA LP

* PURDUE PHARMA LP
 BUTRANS, BUPRENORPHINE
 HYSINGLA ER, HYDROCODONE BITARTRATE
 MS CONTIN, MORPHINE SULFATE
 NALMEFENE HYDROCHLORIDE, NALMEFENE HYDROCHLORIDE
 OXYCONTIN, OXYCODONE HYDROCHLORIDE

PURE SOURCE

* PURE SOURCE LLC
 THEROXIDIL, MINOXIDIL (OTC)

PYROS PHARMS

* PYROS PHARMACEUTICALS INC
 VIGPODER, VIGABATRIN

**** Q ******Q BIOMED**

* Q BIOMED INC
 METASTRON, STRONTIUM CHLORIDE SR-89
 STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89

QILU

* QILU PHARMACEUTICAL CO LTD
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 EXEMESTANE, EXEMESTANE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE

QILU PHARM HAINAN

* QILU PHARMACEUTICAL HAINAN CO LTD
 BORTEZOMIB, BORTEZOMIB
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 GEFITINIB, GEFITINIB
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 TADALAFIL, TADALAFIL

QINGDAO BAHEAL PHARM

* QINGDAO BAHEAL PHARMACEUTICAL CO LTD
 CELECOXIB, CELECOXIB
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE

QOL MEDCL

* QOL MEDICAL LLC
 ETHAMOLIN, ETHANOLAMINE OLEATE

QUAGEN

* QUAGEN PHARMACEUTICALS LLC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Q ****

* QUAGEN PHARMACEUTICALS LLC
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

QUEEN HAMAMATSU PET

* QUEEN HAMAMATSU PET IMAGING CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**** R ******R-PHARM US LLC**

* R-PHARM US LLC
 IXEMPRA KIT, IXABEPILONE

RADIOMEDIX

* RADIOMEDIX INC
 DETECTNET, COPPER CU-64 DOTATATE

RADIUS

* RADIUS HEALTH INC
 BINOSTO, ALENDRONATE SODIUM
 TYMLOS, ABALOPARATIDE

RAYNER SURGICAL

* RAYNER SURGICAL INC
 OMIIDRIA, KETOROLAC TROMETHAMINE

RB HLTH

* RB HEALTH US LLC
 DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
 MUCINEX D, GUAIFENESIN (OTC)
 MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 MUCINEX, GUAIFENESIN (OTC)

REATA PHARMS

* REATA PHARMACEUTICALS INC
 SKYCLARYS, OMAVELOXOLONE

RECORDATI RARE

* RECORDATI RARE DISEASES INC
 CARBAGLU, CARGLUMIC ACID
 CHEMET, SUCCIMER
 COSMEGEN, DACTINOMYCIN
 CYSTADANE, BETAINE
 CYSTADROPS, CYSTEAMINE HYDROCHLORIDE
 ISTURISA, OSILODROSTAT PHOSPHATE
 NEOPROFEN, IBUPROFEN LYSINE
 SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
 SIGNIFOR, PASIREOTIDE DIASPARTATE

REDHILL

* REDHILL BIOPHARMA INC
 AEMCOLO, RIFAMYCIN SODIUM
 * REDHILL BIOPHARMA LTD
 TALICIA, AMOXICILLIN

RELIANCE LIFE

* RELIANCE LIFE SCIENCES PVT LTD
 CAPECITABINE, CAPECITABINE

REMPEX

* REMPEX PHARMACEUTICALS INC A WHOLLY OWNED SUB OF MELINTA THERAPEUTICS LLC
 MINOCIN, MINOCYCLINE HYDROCHLORIDE
 VABOMERE, MEROPENEM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ******RENATA**

* RENATA LTD
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 PREGABALIN, PREGABALIN
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

RENEW PHARMS

* RENEW PHARMACEUTICALS LTD
 INDOCYANINE GREEN, INDOCYANINE GREEN

RESILIA PHARMS

* RESILIA PHARMACEUTICALS INC
 ECOZA, ECONAZOLE NITRATE

REYOUNG

* REYOUNG CORPORATION
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE

RHODES PHARMS

* RHODES PHARMACEUTICALS LP
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DILAUDID, HYDROMORPHONE HYDROCHLORIDE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 MORPHINE SULFATE, MORPHINE SULFATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE

RHYTHM

* RHYTHM PHARMACEUTICALS INC
 IMCIVREE, SETMELANOTIDE ACETATE

RICONPHARMA LLC

* RICONPHARMA LLC
 BISMUTH SUBCITRATE POTASSIUM, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
 CARBAMAZEPINE, CARBAMAZEPINE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 FLUNISOLIDE, FLUNISOLIDE
 ISOSORBIDE DINITRATE AND HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 SCOPOLAMINE, SCOPOLAMINE

RIGEL PHARMS INC

* RIGEL PHARMACEUTICALS INC
 REZLIDHIA, OLUTASIDENIB
 TAVALLISSE, FOSTAMATINIB DISODIUM

RILEY CONSUMER

* RILEY CONSUMER CARE LLC DBA CARLIN CONSUMER HEALTH
 ZEGERID OTC, OMEPRAZOLE (OTC)

RISE PHARMA

* RISE PHARMA LLC
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHLORZOAZONE, CHLORZOAZONE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 KETOCONAZOLE, KETOCONAZOLE

RISING

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

- * RISING PHARMA HOLDING INC
 AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CAPECITABINE, CAPECITABINE
 CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CYTARABINE, CYTARABINE
 DIGOXIN, DIGOXIN
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FLUMAZENIL, FLUMAZENIL
 FLUNISOLIDE, FLUNISOLIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
- * RISING PHARMA HOLDINGS INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACETIC ACID, ACETIC ACID, GLACIAL
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ADENOSINE, ADENOSINE
 AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALFENTA, ALFENTANIL HYDROCHLORIDE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZATHIOPRINE, AZATHIOPRINE
 BACLOFEN, BACLOFEN
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 CALCITRIOL, CALCITRIOL
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHLORZOXAZONE, CHLORZOXAZONE
 CICLOPIROX, CICLOPIROX
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DAPSONE, DAPSONE
 DESOXIMETASONE, DESOXIMETASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 EPLERENONE, EPLERENONE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 EXEMESTANE, EXEMESTANE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCORTISONE, HYDROCORTISONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

- * RISING PHARMA HOLDINGS INC
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 INDAPAMIDE, INDAPAMIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LINEZOLID, LINEZOLID
 LORAZEPAM, LORAZEPAM
 METHIMAZOLE, METHIMAZOLE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NADOLOL, NADOLOL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NATEGLINIDE, NATEGLINIDE
 NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
 NITAZOXANIDE, NITAZOXANIDE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 PARICALCITOL, PARICALCITOL
 PERPHENAZINE, PERPHENAZINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PREGABALIN, PREGABALIN
 PROBENECID AND COLCHICINE, COLCHICINE
 PROBENECID, PROBENECID
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RANOLAZINE, RANOLAZINE
 RISPERIDONE, RISPERIDONE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 SULFAMYLON, MAFENIDE ACETATE
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 TEMOZOLOMIDE, TEMOZOLOMIDE
 THIOTHIXENE, THIOTHIXENE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOLMETIN SODIUM, TOLMETIN SODIUM
 TOREMIFENE CITRATE, TOREMIFENE CITRATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 TROPICACYL, TROPICAMIDE
 VORICONAZOLE, VORICONAZOLE
 ZILEUTON, ZILEUTON
- * RISING PHARMACEUTICALS
 URSODIOL, URSODIOL
- RISING PHARMS**
- * RISING PHARMACEUTICALS INC
 ZAFIRLUKAST, ZAFIRLUKAST
- RK PHARMA**
- * RK PHARMA INC
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ******* RK PHARMA INC**

CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MITOMYCIN, MITOMYCIN
 MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
 NADOLOL, NADOLOL
 PALIPERIDONE, PALIPERIDONE
 URSODIOL, URSODIOL

ROCHE PALO*** ROCHE PALO ALTO LLC**

CELLCEPT, MYCOPHENOLATE MOFETIL
 CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE

ROMARK*** ROMARK LABORATORIES**

ALINIA, NITAZOXANIDE

ROXANE*** ROXANE LABORATORIES INC**

METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE

RUBICON*** RUBICON RESEARCH PRIVATE LTD**

ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 BACLOFEN, BACLOFEN
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARVEDILOL, CARVEDILOL
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GLIPIZIDE, GLIPIZIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LITHIUM CITRATE, LITHIUM CITRATE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NITROGLYCERIN, NITROGLYCERIN
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* RUBICON RESEARCH PRIVATE LTD
 PREGABALIN, PREGABALIN
 PRIMIDONE, PRIMIDONE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

RVL PHARMS

* RVL PHARMACEUTICALS INC
 UPNEEQ, OXYMETAZOLINE HYDROCHLORIDE

**** S ******SAGE CHEMS**

* SAGE CHEMICALS INC
 APOMORPHINE HYDROCHLORIDE, APOMORPHINE HYDROCHLORIDE

SAGE PRODS

* SAGE PRODUCTS INC
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

SAGE THERAP

* SAGE THERAPEUTICS INC
 ZULRESSO, BREXANOLONE

SAGENT

* SAGENT PHARMACEUTICALS
 BUMETANIDE, BUMETANIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 VALPROATE SODIUM, VALPROATE SODIUM

SAGENT PHARMS

* SAGENT PHARMACEUTICALS INC
 CAFFEINE CITRATE, CAFFEINE CITRATE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 FLUMAZENIL, FLUMAZENIL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

SAGENT PHARMS INC

* SAGENT PHARMACEUTICALS INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DAPTOMYCIN, DAPTOMYCIN
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPTIFIBATIDE, EPTIFIBATIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SAGENT PHARMACEUTICALS INC
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUOROURACIL, FLUOROURACIL
 FULVESTRANT, FULVESTRANT
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLYDO, LIDOCAINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LINEZOLID, LINEZOLID
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MESNA, MESNA
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OXYTOCIN, OXYTOCIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROPOFOL, PROPOFOL
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

SALIX

* SALIX PHARMACEUTICALS INC
 APRISO, MESALAMINE
 FENOGLIDE, FENOFIBRATE
 PLENVU, ASCORBIC ACID
 RELISTOR, METHYLNALTREXONE BROMIDE
 TRULANCE, PLECANATIDE
 UCERIS, BUDESONIDE
 ZEGERID, OMEPRAZOLE

SALIX PHARMS

* SALIX PHARMACEUTICALS INC
 ANUSOL HC, HYDROCORTISONE
 DIURIL, CHLOROTHIAZIDE
 MOVIPREP, ASCORBIC ACID
 RELISTOR, METHYLNALTREXONE BROMIDE
 XIFAXAN, RIFAXIMIN

SAMSON MEDCL

* SAMSON MEDICAL TECHNOLOGIES LLC
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFOTIXIM SODIUM IN PLASTIC CONTAINER, CEFOTIXIM SODIUM
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

SANDOZ

* SANDOZ
 DOCETAXEL, DOCETAXEL

* SANDOZ INC
 ACETAMINOPHEN, ACETAMINOPHEN
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALPRAZOLAM, ALPRAZOLAM
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

* SANDOZ INC
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
 ANECTINE, SUCCINYLCHOLINE CHLORIDE
 ANGIOMAX, BIVALIRUDIN
 APREPITANT, APREPITANT
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARRANON, NELARABINE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 AZOPT, BRINZOLAMIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BETOPTIC, BETAXOLOL HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BIMATOPROST, BIMATOPROST
 BORTEZOMIB, BORTEZOMIB
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 CLARITHROMYCIN, CLARITHROMYCIN
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 COSYNTROPIN, COSYNTROPIN
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYCLOSPORINE, CYCLOSPORINE
 DECITABINE, DECITABINE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIGOXIN, DIGOXIN
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DUREZOL, DIFLUPREDNATE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPLERENONE, EPLERENONE
 ETODOLAC, ETODOLAC
 EXELON, RIVASTIGMINE
 EZETIMIBE, EZETIMIBE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FERUMOXYTOL, FERUMOXYTOL
 FLUMAZENIL, FLUMAZENIL
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FULVESTRANT, FULVESTRANT
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GATIFLOXACIN, GATIFLOXACIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GLATOPA, GLATIRAMER ACETATE
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL
 HEPARIN SODIUM, HEPARIN SODIUM
 HYCANTIN, TOPOTECAN HYDROCHLORIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ISONIAZID, ISONIAZID
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ITRACONAZOLE, ITRACONAZOLE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN (COPACKAGED), AMOXICILLIN
 LANSOPRAZOLE, LANSOPRAZOLE
 LATANOPROST, LATANOPROST
 LESCOL XL, FLUVASTATIN SODIUM
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LINEZOLID, LINEZOLID
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOTREL, AMLODIPINE BESYLATE
 MAXITROL, DEXAMETHASONE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYDRIACYL, TROPICAMIDE
 NADOLOL, NADOLOL
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEVIRAPINE, NEVIRAPINE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SANDOZ INC**

NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OMEPRAZOLE, OMEPRAZOLE
 OMNIPRED, PREDNISOLONE ACETATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXALIPLATIN, OXALIPLATIN
 OXAPROZIN, OXAPROZIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PENICILLIN G SODIUM, PENICILLIN G SODIUM
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PERPHENAZINE, PERPHENAZINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PIRFENIDONE, PIRFENIDONE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QOLIANA, BRIMONIDINE TARTRATE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RECLAST, ZOLEDRONIC ACID
 REGONOL, PYRIDOSTIGMINE BROMIDE
 RIBAVIRIN, RIBAVIRIN
 RISPERIDONE, RISPERIDONE
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILODOSIN, SILODOSIN
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TACROLIMUS, TACROLIMUS
 TAFLUPROST, TAFLUPROST
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBRADEX, DEXAMETHASONE
 TOBEX, TOBRAMYCIN
 TRAVATAN Z, TRAVOPROST
 TREPROSTINIL, TREPROSTINIL
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIAZOLAM, TRIAZOLAM
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 TRIFLURIDINE, TRIFLURIDINE
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VIVELLE-DOT, ESTRADIOL
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SANDOZ CANADA INC*** SANDOZ CANADA INC**

INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
 INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
 INFUVITE PEDIATRIC, ASCORBIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

SANDOZ INC

- * SANDOZ INC
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 - CEFTRIAXONE, CEFTRIAXONE SODIUM
 - CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 - CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 - CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 - ENALAPRIL MALEATE, ENALAPRIL MALEATE
 - ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - MONTELUKAST SODIUM, MONTELUKAST SODIUM
 - OLANZAPINE, OLANZAPINE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 - VORICONAZOLE, VORICONAZOLE
 - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

SANOFI

- * SANOFI AVENTIS US LLC
 - FEXINIDAZOLE, FEXINIDAZOLE
 - FLOMAX, TAMSULOSIN HYDROCHLORIDE
- * SANOFI GENZYME
 - HECTOROL, DOXERCALCIFEROL
 - RENVELA, SEVELAMER CARBONATE

SANOFI AVENTIS US

- * SANOFI AVENTIS US INC
 - JEVTANA KIT, CABAZITAXEL
- * SANOFI AVENTIS US LLC
 - AMARYL, GLIMEPIRIDE
 - AMBIEN CR, ZOLPIDEM TARTRATE
 - AMBIEN, ZOLPIDEM TARTRATE
 - ARAVA, LEFLUNOMIDE
 - AUBAGIO, TERIFLUNOMIDE
 - AVALIDE, HYDROCHLOROTHIAZIDE
 - AVAPRO, IRBESARTAN
 - DIABETA, GLYBURIDE
 - ELOXATIN, OXALIPLATIN
 - FERRLECIT, FERRIC OXYHYDROXIDE
 - LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 - LOVENOX, ENOXAPARIN SODIUM
 - MULTAQ, DRONEDARONE HYDROCHLORIDE
 - PLAVIX, CLOPIDOGREL BISULFATE
 - PRIFTIN, RIFAPENTINE
 - PRIMAQUINE, PRIMAQUINE PHOSPHATE
 - RIFADIN, RIFAMPIN
 - TAXOTERE, DOCETAXEL

SANTARUS INC

- * SANTARUS INC
 - GLUMETZA, METFORMIN HYDROCHLORIDE

SAPTALIS PHARMS

- * SAPTALIS PHARMACEUTICALS LLC
 - ACETIC ACID, ACETIC ACID, GLACIAL
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - LEVOCARNITINE, LEVOCARNITINE
 - LIKMEZ, METRONIDAZOLE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - VOSOL HC, ACETIC ACID, GLACIAL

SAREPTA THERAPS INC

- * SAREPTA THERAPEUTICS INC
 - AMONDYS 45, CASIMERSEN
 - EXONDYS 51, ETEPLIRSEN
 - VYONDYS 53, GOLODIRSEN

SARFE PHARMS

- * SARFEZ PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SARFEZ PHARMACEUTICALS INC
SOAANZ, TORSEMIDE

SAVIOR LIFETEC CORP

* SAVIOR LIFETEC CORP
ERTAPENEM SODIUM, ERTAPENEM SODIUM
MEROPENEM, MEROPENEM

SAWAI USA

* SAWAI USA INC
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

SCHERING

* SCHERING CORP
NOXAFIL, POSACONAZOLE

SCIARRA LABS

* SCIARRA LABORATORIES INC
SCLEROSOL, TALC
TALC, TALC

SCIECURE PHARMA INC

* SCIECURE PHARMA INC
CARBAMAZEPINE, CARBAMAZEPINE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

SCIEGEN PHARMS INC

* SCIEGEN PHARMACEUTICALS INC
ARIPIPIRAZOLE, ARIPIPIRAZOLE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARISOPRODOL, CARISOPRODOL
CELECOXIB, CELECOXIB
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CLOTRIMAZOLE, CLOTRIMAZOLE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DROXIDOPA, DROXIDOPA
ETHACRYNIC ACID, ETHACRYNIC ACID
EZETIMIBE, EZETIMIBE
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
HALOPERIDOL, HALOPERIDOL
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
LACOSAMIDE, LACOSAMIDE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LISINAPRIL, LISINAPRIL
METAXALONE, METAXALONE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NABUMETONE, NABUMETONE
NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
PHYTONADIONE, PHYTONADIONE
PIRFENIDONE, PIRFENIDONE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PREGABALIN, PREGABALIN
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** s **

* SCIEGEN PHARMACEUTICALS INC
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 VALPROIC ACID, VALPROIC ACID
 VALSARTAN, VALSARTAN

SCILEX HLDG

* SCILEX HOLDING CO
 ELYXYB, CELECOXIB

SCILEX PHARMS

* SCILEX PHARMACEUTICALS INC
 GLOPERBA, COLCHICINE
 ZTLIDO, LIDOCAINE

SCINOPHARM TAIWAN

* SCINOPHARM TAIWAN LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

SCPHARMACEUTICALS

* SCPHARMACEUTICALS INC
 FUROSCIX, FUROSEMIDE

SCYNEXIS

* SCYNEXIS INC
 BREXAFEMME, IBREXAFUNGERP CITRATE

SEAGEN

* SEAGEN INC
 TUKYSA, TUCATINIB

SEBELA IRELAND LTD

* SEBELA IRELAND LTD
 BRISDELLE, PAROXETINE MESYLATE
 IMURAN, AZATHIOPRINE
 LOTRONEX, ALOSETRON HYDROCHLORIDE
 MICORT-HC, HYDROCORTISONE ACETATE
 MOTOFEN, ATROPINE SULFATE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 PRAMOSONE, HYDROCORTISONE ACETATE
 RIDAURA, AURANOFIN

SECURA

* SECURA BIO INC
 COPIKTRA, DUVELISIB

SENORES PHARMS

* SENORES PHARMACEUTICALS INC
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

SENTISS

* SENTISS AG
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN
 ERYTHROMYCIN, ERYTHROMYCIN
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LOTEPREDNOL ETABONATE, LOTEPREDNOL ETABONATE
 OFLOXACIN, OFLOXACIN

SENTISS PHARMA

* SENTISS PHARMA PRIVATE LTD
 TIMOLOL MALEATE, TIMOLOL MALEATE

SENTYNL THERAPS INC

* SENTYNL THERAPEUTICS INC
 NULLIBRY, FOSDENOPTERIN HYDROBROMIDE

SEPTODONT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SEPTODONT INC**

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
DYCLOPRO, DYCLONINE HYDROCHLORIDE

SEPTODONT HOLDING*** SEPTODONT HOLDING SAS**

ORAVERSE, PHENTOLAMINE MESYLATE

SEPTODONT INC*** SEPTODONT INC**

LIDOCAINE, LIDOCAINE
PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE

SERVIER*** SERVIER PHARMACEUTICALS LLC**

TIBSOVO, IVOSIDENIB

SETON PHARM*** SETON PHARMACEUTICAL LLC**

PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE

SETON PHARMS*** SETON PHARMACEUTICALS LLC**

MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

SHANDONG*** SHANDONG ANXIN PHARMACEUTICAL CO LTD**

PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

*** SHANDONG NEW TIME PHARMACEUTICAL CO LTD**

MILRINONE LACTATE, MILRINONE LACTATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SEVOFLURANE, SEVOFLURANE

SHANDONG LUYE*** SHANDONG LUYE PHARMACEUTICAL CO LTD**

RYKINDO, RISPERIDONE

SHANDONG XINHUA*** SHANDONG XINHUA PHARMACEUTICAL CO LTD**

IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)

SHANGHAI HENGRUI*** SHANGHAI HENGRUI PHARMACEUTICAL CO LTD**

DESFLURANE, DESFLURANE
SEVOFLURANE, SEVOFLURANE

SHENZHEN TECHDOW*** SHENZHEN TECHDOW PHARMACEUTICAL CO LTD**

ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM

SHIELD TX*** SHIELD TX UK LTD**

ACCRUFER, FERRIC MALTOL

SHILPA*** SHILPA MEDICARE LTD**

APREMILAST, APREMILAST
BUSULFAN, BUSULFAN
CAPECITABINE, CAPECITABINE
DOCETAXEL, DOCETAXEL
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
PEMETREXED, PEMETREXED DISODIUM

SHILPA MEDICARE*** SHILPA MEDICARE LTD**

AZACITIDINE, AZACITIDINE

SHIONOGI INC*** SHIONOGI INC**

FETROJA, CEFIDEROCOL SULFATE TOSYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******SHORLA**

* SHORLA PHARMA LTD
 JYLAMVO, METHOTREXATE
 NELARABINE, NELARABINE

SIDMAK LABS INDIA

* SIDMAK LABORATORIES INDIA PVT LTD
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

SIGA TECHNOLOGIES

* SIGA TECHNOLOGIES INC
 TPOXX, TECOVIRIMAT

SIGMAPHARM LABS LLC

* SIGMAPHARM LABORATORIES LLC
 ACITRETIN, ACITRETIN
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 AMBRISENTAN, AMBRISENTAN
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 DISULFIRAM, DISULFIRAM
 DOFETILIDE, DOFETILIDE
 FLUCYTOSINE, FLUCYTOSINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE

SINOTHERAPEUTICS INC

* SINOTHERAPEUTICS INC
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 POSACONAZOLE, POSACONAZOLE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

SINTETICA US

* SINTETICA US LLC
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

SK LIFE

* SK LIFE SCIENCE INC
 XCOPRI, CENOBAMATE

SKYEPHARMA AG

* SKYEPHARMA AG
 TRIGLIDE, FENOFIBRATE

SLATE RUN PHARMA

* SLATE RUN PHARMACEUTICALS LLC
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 AZITHROMYCIN, AZITHROMYCIN
 BIVALIRUDIN, BIVALIRUDIN
 CILOSTAZOL, CILOSTAZOL
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DROXIDOPA, DROXIDOPA
 EPTIFIBATIDE, EPTIFIBATIDE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 METHOCARBAMOL, METHOCARBAMOL
 PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
 VORICONAZOLE, VORICONAZOLE

SLAYBACK PHARMA LLC

* SLAYBACK PHARMA LLC
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 DEOXYCHOLIC ACID, DEOXYCHOLIC ACID
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 LOTEPREDNOL ETABONATE, LOTEPREDNOL ETABONATE
 MERZEE, ETHINYL ESTRADIOL
 METARAMINOL BITARTRATE, METARAMINOL BITARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SLAYBACK PHARMA LLC**

METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 PREVDUO, GLYCOPYRROLATE
 TOFACITINIB CITRATE, TOFACITINIB CITRATE
 VIVIMUSTA, BENDAMUSTINE HYDROCHLORIDE

SOCIETAL CDMO*** SOCIETAL CDMO GAINESVILLE LLC**

VERELAN PM, VERAPAMIL HYDROCHLORIDE
 VERELAN, VERAPAMIL HYDROCHLORIDE

SOFGEN PHARMS*** SOFGEN PHARMACEUTICALS LLC**

IBUPROFEN, IBUPROFEN (OTC)
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 PROGESTERONE, PROGESTERONE

SOFIE*** SOFIE CO DBA SOFIE**

AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

*** SOFIE CO DBA SOFIE (FKA ZEVACOR PHARMA INC)**

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SOLA PHARMS*** SOLA PHARMACEUTICALS**

CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DIMETHYL FUMARATE, DIMETHYL FUMARATE
 TERIFLUNOMIDE, TERIFLUNOMIDE

SOLARIS PHARMA CORP*** SOLARIS PHARMA CORP**

CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 METRONIDAZOLE, METRONIDAZOLE
 TAZAROTENE, TAZAROTENE

SOLUBIOMIX*** SOLUBIOMIX LLC**

COXANTO, OXAPROZIN

SOMERSET*** SOMERSET PHARMACEUTICALS INC**

EMSAM, SELEGILINE

*** SOMERSET THERAPEUTICS LLC**

BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 LATANOPROST, LATANOPROST
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SOMERSET THERAPS LLC*** SOMERSET THERAPEUTICS LLC**

AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BIMATOPROST, BIMATOPROST
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 METHOCARBAMOL, METHOCARBAMOL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SOMERSET THERAPEUTICS LLC**

OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TROPICAMIDE, TROPICAMIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SPECGX LLC*** SPECGX LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
 ANEXSIA 5/325, ACETAMINOPHEN
 ANEXSIA 7.5/325, ACETAMINOPHEN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-37, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHADOSE, METHADONE HYDROCHLORIDE
 METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
 METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
 POSACONAZOLE, POSACONAZOLE
 RESTORIL, TEMAZEPAM
 ROXICODONE, OXYCODONE HYDROCHLORIDE
 TOFRANIL, IMIPRAMINE HYDROCHLORIDE

SPECTRA MDCL DEVICES*** SPECTRA MEDICAL DEVICES INC**

LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

SPIL*** SUN PHARMA INDUSTRIES LTD**

AMPHOTERICIN B, AMPHOTERICIN B
 ASPRUZYO SPRINKLE, RANOLAZINE
 KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

SPRINGWORKS*** SPRINGWORKS THERAPEUTICS INC**

OGSIVEO, NIROGACESTAT HYDROBROMIDE

SPROUT PHARMS*** SPROUT PHARMACEUTICALS INC**

ADDYI, FLIBANSERIN

SQUARE PHARMS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SQUARE PHARMACEUTICALS LTD**

ACYCLOVIR, ACYCLOVIR
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 CALCIUM ACETATE, CALCIUM ACETATE
 MIRTAZAPINE, MIRTAZAPINE
 VALSARTAN, VALSARTAN

SS PHARMA*** SS PHARMA LLC**

DRISDOL, ERGOCALCIFEROL
 HIPREX, METHENAMINE HIPPURATE
 PARLODEL, BROMOCRIPTINE MESYLATE
 ROCALTROL, CALCITRIOL

ST RENATUS*** ST RENATUS LLC**

KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE

STALLION LABS*** STALLION LABORATORIES PRIVATE LTD**

LEVETIRACETAM, LEVETIRACETAM

STAND HOMEOPATH*** STANDARD HOMEOPATHIC CO**

IVY BLOCK, BENTOQUATAM (OTC)

STASON PHARMS*** STASON PHARMACEUTICALS INC**

PURINETHOL, MERCAPTOPYRINE

STEMLINE THERAP*** STEMLINE THERAPEUTICS INC**

ORSERDU, ELACESTRANT DIHYDROCHLORIDE

STERISCIENCE*** STERISCIENCE PTE LTD**

ACETYLCYSTEINE, ACETYLCYSTEINE
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 FOCINVEZ, FOSAPREPITANT DIMEGLUMINE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NAFICILLIN SODIUM, NAFICILLIN SODIUM
 OXACILLIN SODIUM, OXACILLIN SODIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 SODIUM BICARBONATE, SODIUM BICARBONATE

STI PHARMA LLC*** STI PHARMA LLC**

MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE

STIEFEL*** STIEFEL LABORATORIES INC**

DUAC, BENZOYL PEROXIDE

STRIDES PHARMA*** STRIDES PHARMA GLOBAL PTE LTD**

ABACAVIR SULFATE, ABACAVIR SULFATE
 ACARBOSE, ACARBOSE
 ACCOLATE, ZAFIRLUKAST
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACYCLOVIR, ACYCLOVIR
 ALBENDAZOLE, ALBENDAZOLE
 ALPRAZOLAM, ALPRAZOLAM
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 BENZONATATE, BENZONATATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STRIDES PHARMA GLOBAL PTE LTD
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
 CABERGOLINE, CABERGOLINE
 CALCITRIOL, CALCITRIOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 COLCHICINE, COLCHICINE
 CYCLOSPORINE, CYCLOSPORINE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DOFETILIDE, DOFETILIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 DUTASTERIDE, DUTASTERIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ; EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FLUCYTOSINE, FLUCYTOSINE
 GABAPENTIN, GABAPENTIN
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCORTISONE, HYDROCORTISONE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 KETOCONAZOLE, KETOCONAZOLE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MELOXICAM, MELOXICAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHOXSALEN, METHOXSALEN
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NATEGLINIDE, NATEGLINIDE
 NEVIRAPINE, NEVIRAPINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PIROXICAM, PIROXICAM
 POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** s ****

- * STRIDES PHARMA GLOBAL PTE LTD
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 ROFLUMILAST, ROFLUMILAST
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TACROLIMUS, TACROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TESTOSTERONE, TESTOSTERONE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TORSEMIDE, TORSEMIDE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 URSODIOL, URSODIOL
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZILEUTON, ZILEUTON
- STRONGBRIDGE**
- * STRONGBRIDGE DUBLIN LTD
 RECORLEV, LEVOKETOCONAZOLE
- SUCAMPO PHARMA LLC**
- * SUCAMPO PHARMA AMERICAS LLC
 AMITIZA, LUBIPROSTONE
- SUMITOMO PHARMA**
- * SUMITOMO PHARMA SWITZERLAND GMBH
 ORGOVYX, RELUGOLIX
- SUMITOMO PHARMA AM**
- * SUMITOMO PHARMA AMERICA INC
 APTIOM, ESLICARBAZEPINE ACETATE
- SUN PHARM**
- * SUN PHARMACEUTICAL INDUSTRIES LTD
 ABSORICA LD, ISOTRETINOIN
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 AMBRISENTAN, AMBRISENTAN
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BOSENTAN, BOSENTAN
 BROMSITE, BROMFENAC SODIUM
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCITRIOL, CALCITRIOL
 CAPECITABINE, CAPECITABINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CARBOPLATIN, CARBOPLATIN
 CEQUA, CYCLOSPORINE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CIPROFLOXACIN AND DEXAMETHASONE, CIPROFLOXACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

* SUN PHARMACEUTICAL INDUSTRIES LTD
DALFAMPRIDINE, DALFAMPRIDINE
DECITABINE, DECITABINE
DEFERASIROX, DEFERASIROX
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
DOFETILIDE, DOFETILIDE
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DROXIDOPA, DROXIDOPA
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ENTACAPONE, ENTACAPONE
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
ESZOPICLONE, ESZOPICLONE
FEBUXOSTAT, FEBUXOSTAT
FENOFIBRATE, FENOFIBRATE
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
FINASTERIDE, FINASTERIDE
FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
FYREMADEL, GANIRELIX ACETATE
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
IBANDRONATE SODIUM, IBANDRONATE SODIUM
IMATINIB MESYLATE, IMATINIB MESYLATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LACOSAMIDE, LACOSAMIDE
LANSOPRAZOLE, LANSOPRAZOLE
LENALIDOMIDE, LENALIDOMIDE
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
LORATADINE, LORATADINE (OTC)
LOTEPREDNOL ETABONATE, LOTEPREDNOL ETABONATE
LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MESALAMINE, MESALAMINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
NIACIN, NIACIN
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
ODOMZO, SONIDEGIB PHOSPHATE
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OMEPRAZOLE, OMEPRAZOLE (OTC)
OPCICON ONE-STEP, LEVONORGESTREL (OTC)
PALIPERIDONE, PALIPERIDONE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
PREGABALIN, PREGABALIN
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
RANOLAZINE, RANOLAZINE
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SAXAGLIPTIN AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES LTD
 SAXAGLIPTIN, SAXAGLIPTIN HYDROCHLORIDE
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUNITINIB MALATE, SUNITINIB MALATE
 TADALAFIL, TADALAFIL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TOBRAMYCIN, TOBRAMYCIN
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 WINLEVI, CLASCOTERONE
 XELPROS, LATANOPROST
 YONSA, ABIRATERONE ACETATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SUN PHARM INDS

* SUN PHARMACEUTICAL INDUSTRIES LTD
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLANZAPINE, OLANZAPINE
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

SUN PHARM INDS (IN)

* SUN PHARMACEUTICAL INDUSTRIES LTD
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

SUN PHARM INDS INC

* SUN PHARMACEUTICAL INDUSTRIES INC
 ABSORICA, ISOTRETINOIN
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
 CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 CLONAZEPAM, CLONAZEPAM
 CLOZAPINE, CLOZAPINE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIGOXIN, DIGOXIN
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 HALOG, HALCINONIDE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 REPAGLINIDE, REPAGLINIDE
 SEZABY, PHENOBARBITAL SODIUM
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT
 TECHNETIUM TC99M MERTIATIDE KIT, TECHNETIUM TC-99M MERTIATIDE KIT
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES INC
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

SUN PHARM INDS LTD

* SUN PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DOXYCYCLINE, DOXYCYCLINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FELODIPINE, FELODIPINE
 FENOFIBRATE, FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LEVETIRACETAM, LEVETIRACETAM
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE REDIDOSE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 RILUZOLE, RILUZOLE
 RISPERIDONE, RISPERIDONE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TOPIRAMATE, TOPIRAMATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID

SUN PHARM INDUSTRIES

* SUN PHARMACEUTICAL INDUSTRIES INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALLOPURINOL, ALLOPURINOL
 BACTRIM DS, SULFAMETHOXAZOLE
 BACTRIM, SULFAMETHOXAZOLE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 CHLORTHALIDONE, CHLORTHALIDONE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ERGOLOID MESYLATES, ERGOLOID MESYLATES
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES INC
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NYSTATIN, NYSTATIN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PREDNISON, PREDNISON
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 QUALAQUIN, QUININE SULFATE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 TEMAZEPAM, TEMAZEPAM
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 ULTRAVATE, HALOBETASOL PROPIONATE

SUNNY

* SUNNY PHARMTECH INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 CARBOPROST TROMETHAMINE, CARBOPROST TROMETHAMINE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN

SUNOVION PHARMS INC

* SUNOVION PHARMACEUTICALS INC
 LATUDA, LURASIDONE HYDROCHLORIDE

SUNSHINE

* SUNSHINE LAKE PHARMA CO LTD
 ARIPIPRAZOLE, ARIPIPRAZOLE
 AZITHROMYCIN, AZITHROMYCIN
 CLARITHROMYCIN, CLARITHROMYCIN
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 FEBUXOSTAT, FEBUXOSTAT
 LINAGLIPTIN AND METFORMIN HYDROCHLORIDE, LINAGLIPTIN
 LINAGLIPTIN, LINAGLIPTIN
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 RANOLAZINE, RANOLAZINE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TADALAFIL, TADALAFIL

SUPERNUS PHARMS

* SUPERNUS PHARMACEUTICALS INC
 OXTELLAR XR, OXCARBAZEPINE
 QELBREE, VILOXAZINE HYDROCHLORIDE
 TROKENDI XR, TOPIRAMATE

SUVEN PHARMS

* SUVEN PHARMACEUTICALS LTD
 CALCIUM ACETATE, CALCIUM ACETATE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 MALATHION, MALATHION

SVC PHARMA

* SVC PHARMA LP
 DRONABINOL, DRONABINOL

SWEDISH ORPHAN

* SWEDISH ORPHAN BIOVITRUM AB PUBL
 ORFADIN, NITISINONE

SYNTHON PHARMS

* SYNTHON PHARMACEUTICALS INC
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

SYNTHON PHARMS INC

* SYNTHON PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SYNTHON PHARMACEUTICALS INC
PALBOCICLIB, PALBOCICLIB

**** T ******ACME LABS**

* THE ACME LABORATORIES LTD
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
METHOCARBAMOL, METHOCARBAMOL
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

GEN HOSP

* THE GENERAL HOSPITAL CORP
AMMONIA N 13, AMMONIA N-13

METHODIST

* THE METHODIST HOSP RESEARCH INSTITUTE
AMMONIA N 13, AMMONIA N-13

METHODIST HOSP RES

* THE METHODIST HOSP RESEARCH INSTITUTE
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

RITEDOSE CORP

* THE RITEDOSE CORP
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TAGI

* TAGI PHARMA INC
CHLORTHALIDONE, CHLORTHALIDONE
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
METHAZOLAMIDE, METHAZOLAMIDE

TAIHO ONCOLOGY

* TAIHO ONCOLOGY INC
LONSURF, TIPIRACIL HYDROCHLORIDE
LYTGOBI, FUTIBATINIB

TAKEDA PHARMS USA

* TAKEDA PHARMACEUTICALS USA INC
ACTOPLUS MET, METFORMIN HYDROCHLORIDE
ACTOS, PIOGLITAZONE HYDROCHLORIDE
ADDERALL XR 10, AMPHETAMINE ASPARTATE
ADDERALL XR 15, AMPHETAMINE ASPARTATE
ADDERALL XR 20, AMPHETAMINE ASPARTATE
ADDERALL XR 25, AMPHETAMINE ASPARTATE
ADDERALL XR 30, AMPHETAMINE ASPARTATE
ADDERALL XR 5, AMPHETAMINE ASPARTATE
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
ALUNBRIG, BRIGATINIB
CARBATROL, CARBAMAZEPINE
COLCRYS, COLCHICINE
DEXILANT, DEXLANSOPRAZOLE
DUETACT, GLIMEPIRIDE
EXKIVITY, MOBOCERTINIB SUCCINATE
FIRAZYR, ICATIBANT ACETATE
FOSRENOL, LANTHANUM CARBONATE
FRUZAQLA, FRUQUINTINIB
GATTEX KIT, TEDUGLUTIDE RECOMBINANT
ICLUSIG, PONATINIB HYDROCHLORIDE
INTUNIV, GUANFACINE HYDROCHLORIDE
KAZANO, ALOGLIPTIN BENZOATE
LIALDA, MESALAMINE
LIVTENCITY, MARIBAVIR
MOTTEGRITY, PRUCALOPRIDE SUCCINATE
MYDAYIS, AMPHETAMINE ASPARTATE
NESINA, ALOGLIPTIN BENZOATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TAKEDA PHARMACEUTICALS USA INC
 NINLARO, IXAZOMIB CITRATE
 OSENI, ALOGLIPTIN BENZOATE
 PENTASA, MESALAMINE
 PREVACID, LANSOPRAZOLE
 ROZEREM, RAMELTEON
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 ULORIC, FEBUXOSTAT
 VELCADE, BORTEZOMIB
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE

TAMARANG

* TAMARANG SA
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

TARO

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAZOLAMIDE, ACETAZOLAMIDE
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUMETANIDE, BUMETANIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CARBAMAZEPINE, CARBAMAZEPINE
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOCORTOLONE PIVALATE, CLOCORTOLONE PIVALATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 DEFERIPRONE, DEFERIPRONE
 DESONIDE, DESONIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FELBAMATE, FELBAMATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE, HYDROCORTISONE
 IMIQUIMOD, IMIQUIMOD
 INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 IVERMECTIN, IVERMECTIN (OTC)
 KETOCONAZOLE, KETOCONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 MELOXICAM, MELOXICAM
 METRONIDAZOLE, METRONIDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TARO PHARMACEUTICAL INDUSTRIES LTD
 - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 - MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 - NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 - NYSTATIN, NYSTATIN
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - OXCARBAZEPINE, OXCARBAZEPINE
 - PHENYTOIN, PHENYTOIN
 - SILDENAFIL CITRATE, SILDENAFIL CITRATE
 - TERIL, CARBAMAZEPINE
 - TOPICORT, DESOXIMETASONE
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 - WARFARIN SODIUM, WARFARIN SODIUM
- * TARO PHARMACEUTICALS INC
 - ACETIC ACID, ACETIC ACID, GLACIAL
 - ACYCLOVIR, ACYCLOVIR
 - ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 - ADAPALENE, ADAPALENE
 - ADAPALENE, ADAPALENE (OTC)
 - ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 - AMMONIUM LACTATE, AMMONIUM LACTATE
 - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
 - CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - DAPSONE, DAPSONE
 - DERMABET, BETAMETHASONE VALERATE
 - DOCOSANOL, DOCOSANOL (OTC)
 - IBUPROFEN, IBUPROFEN (OTC)
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 - NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 - OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 - TAVABOROLE, TAVABOROLE
 - TAZAROTENE, TAZAROTENE
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 - TERCONAZOLE, TERCONAZOLE
 - TOPICORT, DESOXIMETASONE
 - TRETINOIN, TRETINOIN
- * TARO PHARMACEUTICALS USA INC
 - AZELAIC ACID, AZELAIC ACID
 - CICLOPIROX, CICLOPIROX
 - CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - CLOTRIMAZOLE, CLOTRIMAZOLE
 - CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 - DAPSONE, DAPSONE
 - DESONIDE, DESONIDE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 - ECONAZOLE NITRATE, ECONAZOLE NITRATE
 - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 - FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 - FLUOCINONIDE, FLUOCINONIDE
 - GENTAMICIN SULFATE, GENTAMICIN SULFATE
 - HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 - HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 - HYDROCORTISONE, HYDROCORTISONE
 - IBUPROFEN, IBUPROFEN
 - IBUPROFEN, IBUPROFEN (OTC)
 - KETOZOLE, KETOCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TARO PHARMACEUTICALS USA INC
 LIDOCAINE, LIDOCAINE
 LORATADINE, LORATADINE (OTC)
 MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN
 MUPIROCIN, MUPIROCIN CALCIUM
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 PHENYTOIN, PHENYTOIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)

TARO PHARM INDS

* TARO PHARMACEUTICAL INDUSTRIES LTD
 AMCINONIDE, AMCINONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 LAMOTRIGINE, LAMOTRIGINE

TARSUS

* TARSUS PHARMACEUTICALS INC
 XDEMVI, LOTILANER

TASMAN PHARMA

* TASMAN PHARMA INC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 FLUCINOLONE ACETONIDE, FLUCINOLONE ACETONIDE
 KETOCONAZOLE, KETOCONAZOLE
 VERSACLOZ, CLOZAPINE

TCG FLUENT PHARMA

* TCG FLUENT PHARMA INVESTORS LP
 FLOLIPID, SIMVASTATIN

TEIKOKU PHARMA USA

* TEIKOKU PHARMA USA INC
 LIDODERM, LIDOCAINE

TELIGENT

* TELIGENT PHARMA INC
 FLURANDRENOLIDE, FLURANDRENOLIDE

TELIIX

* TELIIX PHARMACEUTICALS US INC
 ILLUCCIX, GALLIUM GA-68 GOZETOTIDE

TENSHI

* TENSHI KAIZEN PVT LTD
 LORATADINE, LORATADINE (OTC)

TERSERA

* TERSERA THERAPEUTICS LLC
 ERGOMAR, ERGOTAMINE TARTRATE
 PRIALT, ZICONOTIDE ACETATE
 VARUBI, ROLAPITANT HYDROCHLORIDE
 XERMELO, TELOTRISTAT ETIPRATE
 ZOLADEX, GOSERELIN ACETATE

TETRAPHASE PHARMS

* TETRAPHASE PHARMACEUTICALS INC
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******TEVA**

* TEVA NEUROSCIENCE INC
 AUSTEDO XR, DEUTETRABENAZINE
 AZILECT, RASAGILINE MESYLATE
 UZEDY, RISPERIDONE

* TEVA PHARMACEUTICALS USA INC
 ACYCLOVIR, ACYCLOVIR
 ADIPEX-P, PHENTERMINE HYDROCHLORIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN PEDIATRIC, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 ATENOLOL, ATENOLOL
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CALCITRIOL, CALCITRIOL
 CARVEDILOL, CARVEDILOL
 CEFACLOR, CEFACLOR
 CEFPROZIL, CEFPROZIL
 CELECOXIB, CELECOXIB
 CEPHALEXIN, CEPHALEXIN
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CILOSTAZOL, CILOSTAZOL
 CIMETIDINE, CIMETIDINE
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 EPITOL, CARBAMAZEPINE
 ESZOPICLONE, ESZOPICLONE
 ETODOLAC, ETODOLAC
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GALZIN, ZINC ACETATE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE, GLYBURIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOVASTATIN, LOVASTATIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 MUPIROCIN, MUPIROCIN
 NAPROXEN, NAPROXEN
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OFLOXACIN, OFLOXACIN
 OXAPROZIN, OXAPROZIN
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PENICILLIN-VK, PENICILLIN V POTASSIUM
 PIROXICAM, PIROXICAM
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRELONE, PREDNISOLONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TEVA BRANDED PHARM

* TEVA BRANDED PHARMACEUTICAL PRODUCTS R AND D INC
 ACTIGALL, URSODIOL
 AUSTEDO, DEUTETRABENAZINE
 CONDYLOX, PODOFILOX
 LOSEASONIQUE, ETHINYL ESTRADIOL
 MICROZIDE, HYDROCHLOROTHIAZIDE
 NOR-QD, NORETHINDRONE
 PROAIR DIGIHALER, ALBUTEROL SULFATE
 PROAIR HFA, ALBUTEROL SULFATE
 PROAIR RESPICLICK, ALBUTEROL SULFATE
 PROGLYCEM, DIAZOXIDE
 QNASL, BECLOMETHASONE DIPROPIONATE
 QUARTETTE, ETHINYL ESTRADIOL
 SEASONALE, ETHINYL ESTRADIOL
 SEASONIQUE, ETHINYL ESTRADIOL
 ZIAC, BISOPROLOL FUMARATE

TEVA PARENTERAL

* TEVA PARENTERAL MEDICINES INC
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TEVA PHARM

* TEVA PHARMACEUTICAL INDUSTRIES LTD
 AIRDUO DIGIHALER, FLUTICASONE PROPIONATE
 AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
 ARMONAIR DIGIHALER, FLUTICASONE PROPIONATE

TEVA PHARMS

* TEVA PHARMACEUTICALS DEVELOPMENT INC
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 LISDEXAMFETAMINE DIMESYLATE, LISDEXAMFETAMINE DIMESYLATE
 PYRIMETHAMINE, PYRIMETHAMINE

* TEVA PHARMACEUTICALS USA
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 BUDESONIDE, BUDESONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 FAMCICLOVIR, FAMCICLOVIR
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 IRBESARTAN, IRBESARTAN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******* TEVA PHARMACEUTICALS USA**

LANSOPRAZOLE, LANSOPRAZOLE
 LETROZOLE, LETROZOLE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREGABALIN, PREGABALIN
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 VANDAZOLE, METRONIDAZOLE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

TEVA PHARMS INC*** TEVA PHARMACEUTICALS INC**

ALVAIZ, ELTROMBOPAG CHOLINE
 ALYQ, TADALAFIL
 CETRORELIX ACETATE, CETRORELIX ACETATE
 PAZOPANIB HYDROCHLORIDE, PAZOPANIB HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE

TEVA PHARMS INTL*** TEVA PHARMACEUTICALS INTERNATIONAL GMBH**

AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE
 SYNRIBO, OMACETAXINE MEPESUCCINATE

TEVA PHARMS USA*** TEVA PHARMACEUTICALS USA**

ACITRETIN, ACITRETIN
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 ARFORMOTEROL TARTRATE, ARFORMOTEROL TARTRATE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BEXAROTENE, BEXAROTENE
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUDESONIDE, BUDESONIDE
 CLARAVIS, ISOTRETINOIN
 CLOZAPINE, CLOZAPINE
 COPAXONE, GLATIRAMER ACETATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 EPTIFIBATIDE, EPTIFIBATIDE
 ESTRADIOL, ESTRADIOL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 IBUPROFEN AND FAMOTIDINE, FAMOTIDINE
 ICOSAPENT ETHYL, ICOSAPENT ETHYL
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LOGILIA, ULIPRISTAL ACETATE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OMEPRAZOLE, OMEPRAZOLE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PARICALCITOL, PARICALCITOL
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUNITINIB MALATE, SUNITINIB MALATE
 TADALAFIL, TADALAFIL
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOBRAMYCIN, TOBRAMYCIN
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TREPROSTINIL, TREPROSTINIL
 VIGABATRIN, VIGABATRIN
 VILAZODONE HYDROCHLORIDE, VILAZODONE HYDROCHLORIDE
 ZANOSAR, STREPTOZOCIN

* TEVA PHARMACEUTICALS USA INC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ALYQ, TADALAFIL
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 CAPECITABINE, CAPECITABINE
 DAPTOMYCIN, DAPTOMYCIN
 DARUNAVIR, DARUNAVIR
 DEFERASIROX, DEFERASIROX
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 EFINACONAZOLE, EFINACONAZOLE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
 ESTRADIOL, ESTRADIOL
 EVEROLIMUS, EVEROLIMUS
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUTICASONE PROPIONATE AND SALMETEROL XINAFOATE, FLUTICASONE PROPIONATE
 ICATIBANT ACETATE, ICATIBANT ACETATE
 IVERMECTIN, IVERMECTIN
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 MESALAMINE, MESALAMINE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METRONIDAZOLE, METRONIDAZOLE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 ONSURA, ETHINYL ESTRADIOL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PENCICLOVIR, PENCICLOVIR
 PIRFENIDONE, PIRFENIDONE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TERIPARATIDE, TERIPARATIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TEVA PHARMS USA INC

* TEVA PHARMACEUTICALS USA INC
 CHLORZOAZONE, CHLORZOAZONE
 CYCLOSPORINE, CYCLOSPORINE
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHINYL ESTRADIOL; ETNOGESTREL, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 FINZALA, ETHINYL ESTRADIOL
 FORMOTEROL FUMARATE, FORMOTEROL FUMARATE
 FULVESTRANT, FULVESTRANT
 LUBIPROSTONE, LUBIPROSTONE
 MIFEPRISTONE, MIFEPRISTONE
 NALTREXONE, NALTREXONE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PLERIXAFOR, PLERIXAFOR
 POTASSIUM CITRATE, POTASSIUM CITRATE
 RISPERIDONE, RISPERIDONE
 SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
 TASIMELTEON, TASIMELTEON
 TIOPRONIN, TIOPRONIN

TEYRO LABS

* TEYRO LABS PRIVATE LTD
 CAPECITABINE, CAPECITABINE
 CARBOPLATIN, CARBOPLATIN
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

THE FEINSTEIN INST

* THE FEINSTEIN INSTITUTE FOR MEDICAL RESEARCH
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

THE J MOLNER

* THE J MOLNER CO OU
 DESOXIMETASONE, DESOXIMETASONE
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE

THEA PHARMA

* THEA PHARMA INC
 AKTEN, LIDOCAINE HYDROCHLORIDE
 AZASITE, AZITHROMYCIN
 BETIMOL, TIMOLOL
 COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
 COSOPT, DORZOLAMIDE HYDROCHLORIDE
 IYUZEH, LATANOPROST
 ZIOPTAN, TAFLUPROST

THEPHARMANETWORK LLC

* THEPHARMANETWORK LLC
 ISONIAZID, ISONIAZID
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 NIMODIPINE, NIMODIPINE
 THERMAZENE, SILVER SULFADIAZINE

THERACOSBIO

* THERACOSBIO LLC
 BRENZAVVY, BEXAGLIFLOZIN

THERAGNOSTICS

* THERAGNOSTICS INC
 NEPHROSCAN, TECHNETIUM TC-99M SUCCIMER

THERAVIA

* THERAVIA PHARMA
 SIKLOS, HYDROXYUREA

THINQ PHARM-CRO PVT

* THINQ PHARMA-CRO PRIVATE LTD
 FLUCONAZOLE, FLUCONAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 TINIDAZOLE, TINIDAZOLE

TIANJIN KINGYORK

* TIANJIN KINGYORK PHARMACEUTICALS CO LTD
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE

TIANJIN TIANYAO

* TIANJIN TIANYAO PHARMACEUTICALS CO LTD
 CELECOXIB, CELECOXIB

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TIANJIN TIANYAO PHARMACEUTICALS CO LTD
METHYLPREDNISOLONE, METHYLPREDNISOLONE

TILDE SCIENCES

* TILDE SCIENCES LLC
DARAPRIM, PYRIMETHAMINE

TOLMAR

* TOLMAR INC
ELIGARD KIT, LEUPROLIDE ACETATE
JATENZO, TESTOSTERONE UNDECANOATE
KETOCONAZOLE, KETOCONAZOLE

* TOLMAR INTERNATIONAL LTD
FENSOLVI KIT, LEUPROLIDE ACETATE

TONIX MEDS

* TONIX MEDICINES INC
TOSYMRA, SUMATRIPTAN
ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE

TOPROL

* TOPROL ACQUISITION LLC
TOPROL-XL, METOPROLOL SUCCINATE

TORPHARM

* TORPHARM INC
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

TORRENT

* TORRENT PHARMA INC
APREPITANT, APREPITANT
DICHLORPHENAMIDE, DICHLORPHENAMIDE
ERYTHROMYCIN, ERYTHROMYCIN
NITISINONE, NITISINONE

* TORRENT PHARMACEUTICALS LTD
ACYCLOVIR, ACYCLOVIR
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ARIPIRAZOLE, ARIPIRAZOLE
CELECOXIB, CELECOXIB
DAPSONE, DAPSONE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
ITRACONAZOLE, ITRACONAZOLE
LAMOTRIGINE, LAMOTRIGINE
LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
NEBIVOLOL HYDROCHLORIDE, NEBIVOLOL HYDROCHLORIDE
NYSTATIN, NYSTATIN
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
PENCICLOVIR, PENCICLOVIR
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SILDENAFIL CITRATE, SILDENAFIL CITRATE
SORAFENIB TOSYLATE, SORAFENIB TOSYLATE
TADALAFIL, TADALAFIL
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

TORRENT PHARMS

* TORRENT PHARMACEUTICALS LTD
CARBAMAZEPINE, CARBAMAZEPINE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TORRENT PHARMACEUTICALS LTD
RISPERIDONE, RISPERIDONE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TORRENT PHARMS LTD

* TORRENT PHARMACEUTICALS LTD
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FELODIPINE, FELODIPINE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
SILDENAFIL CITRATE, SILDENAFIL CITRATE

TRAVERE

* TRAVERE THERAPEUTICS INC
FILSPARI, SPARSENTAN

TREVENA

* TREVENA INC
OLINVYK, OLICERIDINE

TRIS PHARMA INC

* TRIS PHARMA INC
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
DYANAVAL XR 10, AMPHETAMINE
DYANAVAL XR 15, AMPHETAMINE
DYANAVAL XR 20, AMPHETAMINE
DYANAVAL XR 5, AMPHETAMINE
DYANAVAL XR, AMPHETAMINE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
IBUPROFEN, IBUPROFEN (OTC)
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
THEOPHYLLINE, THEOPHYLLINE

TRUSTEES UNIV PA

* TRUSTEES OF THE UNIV OF PENNSYLVANIA
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

TULEX PHARMS INC

* TULEX PHARMACEUTICALS INC
AMINOCAPROIC ACID, AMINOCAPROIC ACID
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
NAPROXEN, NAPROXEN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

TWI PHARMS

* TWI PHARMACEUTICALS INC
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIMETHYL FUMARATE, DIMETHYL FUMARATE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FORFIVO XL, BUPROPION HYDROCHLORIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
MEGESTROL ACETATE, MEGESTROL ACETATE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MYCOPHENOLIC SODIUM, MYCOPHENOLIC SODIUM
NAPRELAN, NAPROXEN SODIUM
NIFEDIPINE, NIFEDIPINE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TWI PHARMACEUTICALS INC
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TESTOSTERONE, TESTOSTERONE
 TOPIRAMATE, TOPIRAMATE
 ZESTRIL, LISINAPRIL

TWI PHARMS INC

* TWI PHARMACEUTICALS INC
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE

**** U ******UBI**

* UBI PHARMA INC
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 VORICONAZOLE, VORICONAZOLE

UCB INC

* UCB INC
 BRIVIACT, BRIVARACETAM
 FINTEPLA, FENFLURAMINE HYDROCHLORIDE
 KEPPRA XR, LEVETIRACETAM
 KEPPRA, LEVETIRACETAM
 NAYZILAM, MIDAZOLAM
 NEUPRO, ROTIGOTINE
 VIMPAT, LACOSAMIDE
 ZILBRYSQ, ZILUCOPLAN SODIUM

UCLA BIOMEDICAL

* UCLA BIOMEDICAL CYCLOTRON
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UCSF RODIOPHARM

* UCSF RODIOPHARMACEUTICAL FACILITY
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UIHC PET IMAGING

* UNIV IOWA HOSPS AND CLINICS PET IMAGING CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 GALLIUM GA 68 EDOTREOTIDE, GALLIUM GA-68 EDOTREOTIDE

ULTRAGENYX PHARM INC

* ULTRAGENYX PHARMACEUTICAL INC
 DOJOLVI, TRIHEPTANOIN

UMEDICA

* UMEDICA LABORATORIES PRIVATE LTD
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 CARBAMAZEPINE, CARBAMAZEPINE
 CELECOXIB, CELECOXIB
 CHLORTHALIDONE, CHLORTHALIDONE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 GLYCOPYRROLATE, GLYCOPYRROLATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TADALAFIL, TADALAFIL

UNICHEM

* UNICHEM LABORATORIES LTD
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALLOPURINOL, ALLOPURINOL
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UNICHEM LABORATORIES LTD
 ARIPIPRAZOLE, ARIPIPRAZOLE
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 BACLOFEN, BACLOFEN
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CHLORTHALIDONE, CHLORTHALIDONE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MELOXICAM, MELOXICAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PIROXICAM, PIROXICAM
 PRASUGREL HYDROCHLORIDE, PRASUGREL HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANOLAZINE, RANOLAZINE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 ZALEPLON, ZALEPLON
 ZONISAMIDE, ZONISAMIDE

UNICHEM LABS LTD

* UNICHEM LABORATORIES LIMITED
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 * UNICHEM LABORATORIES LTD
 LAMOTRIGINE, LAMOTRIGINE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE

UNIMARK REMEDIES LTD

* UNIMARK REMEDIES LTD
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

UNIQUE

* UNIQUE PHARMACEUTICAL LABORATORIES A DIV OF JB CHEMICALS AND PHARMACEUTICALS LTD
 ATENOLOL, ATENOLOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 GLIPIZIDE, GLIPIZIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

UNIQUE PHARM

* UNIQUE PHARMACEUTICAL LABORATORIES
 CARBAMAZEPINE, CARBAMAZEPINE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)

UNITED BIOMEDCL

* UNITED BIOMEDICAL INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UNITED BIOMEDICAL INC
TERBUTALINE SULFATE, TERBUTALINE SULFATE

UNITED GUARDIAN

* UNITED GUARDIAN INC
RENACIDIN, CITRIC ACID

UNITED RES LABS

* UNITED RESEARCH LABORATORIES LLC
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
OXAZEPAM, OXAZEPAM

UNITED THERAP

* UNITED THERAPEUTICS CORP
ORENITRAM, TREPROSTINIL DIOLAMINE
REMODULIN, TREPROSTINIL
TYVASO DPI, TREPROSTINIL
TYVASO, TREPROSTINIL

UNIV ALAHAMA BIRM

* UNIV ALABAMA AT BIRMINGHAM
AMMONIA N 13, AMMONIA N-13

UNIV CA LOS ANGELES

* UNIV CALIFORNIA LOS ANGELES
GALLIUM GA 68 GOZETOTIDE, GALLIUM GA-68 GOZETOTIDE

UNIV MICHIGAN

* UNIV MICHIGAN PET RADIOPHARMACEUTICAL PRODUCTION PROGRAM
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV OF CA SAN FRAN

* UNIV OF CALIFORNIA SAN FRANCISCO
GALLIUM GA 68 GOZETOTIDE, GALLIUM GA-68 GOZETOTIDE

UNIV SOUTHERN CA

* UNIV SOUTHERN CALIFORNIA DBA USC MOLECULAR IMAGING CENTER
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV TX MD ANDERSON

* UNIV TEXAS MD ANDERSON CANCER CENTER
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV TX SW MEDCTR

* UNIV TEXAS SOUTHWESTERN MEDCTR
AMMONIA N 13, AMMONIA N-13

UNIV UTAH CYCLOTRON

* UNIV UTAH CYCLOTRON RADIOCHEMISTRY LAB
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

UNIV WISCONSIN

* UNIV WISCONSIN SYSTEM
AMMONIA N 13, AMMONIA N-13

UPJOHN

* UPJOHN MANUFACTURING IRELAND UNLTD
LIPITOR, ATORVASTATIN CALCIUM
RELPAX, ELETRIPTAN HYDROBROMIDE

* UPJOHN US 2 LLC
CELEBREX, CELECOXIB
DETROL LA, TOLTERODINE TARTRATE
DETROL, TOLTERODINE TARTRATE
EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
INSPRA, EPLERENONE
LYRICA CR, PREGABALIN
LYRICA, PREGABALIN
XALATAN, LATANOPROST
XANAX XR, ALPRAZOLAM
XANAX, ALPRAZOLAM

UPSHER SMITH LABS

* UPSHER SMITH LABORATORIES LLC
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BACLOFEN, BACLOFEN
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UPSHER SMITH LABORATORIES LLC
 BEXAROTENE, BEXAROTENE
 BRIMONIDINE TARTRATE AND TIMOLOL MALEATE, BRIMONIDINE TARTRATE
 BUMETANIDE, BUMETANIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORZOXAZONE, CHLORZOXAZONE
 CLOBAZAM, CLOBAZAM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DROXIDOPA, DROXIDOPA
 ELIGLUSTAT TARTRATE, ELIGLUSTAT TARTRATE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 EXEMESTANE, EXEMESTANE
 FAMOTIDINE, FAMOTIDINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 HALOPERIDOL, HALOPERIDOL
 ISOTRETINOIN, ISOTRETINOIN
 JANTOVEN, WARFARIN SODIUM
 Klor-CON M10, POTASSIUM CHLORIDE
 Klor-CON M15, POTASSIUM CHLORIDE
 Klor-CON M20, POTASSIUM CHLORIDE
 Klor-CON, POTASSIUM CHLORIDE
 LAMIVUDINE, LAMIVUDINE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYORISAN, ISOTRETINOIN
 NYSTATIN, NYSTATIN
 ORVATEN, MIDODRINE HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PACERONE, AMIODARONE HYDROCHLORIDE
 PREVALITE, CHOLESTYRAMINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 QUDEXY XR, TOPIRAMATE
 RAMELTEON, RAMELTEON
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 VIGABATRIN, VIGABATRIN
 VOGELXO, TESTOSTERONE

URL LABS

* URL LABORATORIES LLC
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

UROGEN PHARMA

* UROGEN PHARMA LTD
 JELMYTO, MITOMYCIN

UROVANT

* UROVANT SCIENCES GMBH
 GEMTESA, VIBEGRON

US ANTIBIOTICS

* US ANTIBIOTICS LLC
 AMOXIL, AMOXICILLIN
 AUGMENTIN '125', AMOXICILLIN
 AUGMENTIN '250', AMOXICILLIN
 AUGMENTIN '875', AMOXICILLIN
 AUGMENTIN ES-600, AMOXICILLIN
 LAROTID, AMOXICILLIN

USPHARMA

* USPHARMA LTD
 NITRO-DUR, NITROGLYCERIN

USPHARMA WINDLAS

* USPHARMA WINDLAS LLC
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE

USV

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

- * USV PRIVATE LTD
 - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE (OTC)
 - ZOLEDRONIC ACID, ZOLEDRONIC ACID

USWM

- * USWM LLC
 - CORGARD, NADOLOL
 - IWILFIN, EFLORNITHINE HYDROCHLORIDE
 - LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 - REVONTO, DANTROLENE SODIUM

**** V ******VALEANT**

- * VALEANT PHARMACEUTICALS INTERNATIONAL
 - BONTRIL PDM, PHENDIMETRAZINE TARTRATE
 - MYSOLINE, PRIMIDONE

VALEANT BERMUDA

- * VALEANT INTERNATIONAL BERMUDA
 - RETIN-A, TRETINOIN

VALEANT INTL

- * VALEANT INTERNATIONAL BARBADOS SRL
 - RETIN-A, TRETINOIN
- * VALEANT INTERNATIONAL SRL
 - BENZAMYCIN, BENZOYL PEROXIDE

VALEANT LUXEMBOURG

- * VALEANT PHARMACEUTICALS LUXEMBOURG SARL
 - TARGRETIN, BEXAROTENE
 - VISUDYNE, VERTEPORFIN

VALEANT PHARM INTL

- * VALEANT PHARMACEUTICALS INTERNATIONAL
 - ANDROID 25, METHYLTESTOSTERONE
 - LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE

VALEANT PHARMS

- * VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 - PENTOXIFYLLINE, PENTOXIFYLLINE

VALEANT PHARMS INTL

- * VALEANT PHARMACEUTICALS INTERNATIONAL
 - COLAZAL, BALSALAZIDE DISODIUM
 - CUPRIMINE, PENICILLAMINE

VALEANT PHARMS NORTH

- * VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 - CARAC, FLUOROURACIL
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - FENOFIBRATE, FENOFIBRATE
 - NIFEDIPINE, NIFEDIPINE
 - RENOVA, TRETINOIN
 - RETIN-A, TRETINOIN
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

VALIDUS PHARMS

- * VALIDUS PHARMACEUTICALS LLC
 - ANZEMET, DOLASETRON MESYLATE
 - BUMEX, BUMETANIDE
 - EQUETRO, CARBAMAZEPINE
 - LASIX, FUROSEMIDE
 - LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
 - LOPRESSOR, METOPROLOL TARTRATE
 - LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 - LOTENSIN, BENAZEPRIL HYDROCHLORIDE
 - NORPRAMIN, DESIPRAMINE HYDROCHLORIDE

VALIDUS PHARMS INC

- * VALIDUS PHARMACEUTICALS INC
 - MARPLAN, ISOCARBOXAZID

VALINOR

- * VALINOR PHARMA LLC
 - MOVANTIK, NALOXEGOL OXALATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ******VANCOCIN ITALIA**

* VANCOCIN ITALIA SRL
MULPLETA, LUSUTROMBOPAG

VANDA PHARMS INC

* VANDA PHARMACEUTICALS INC
FANAPT, ILOPERIDONE
HETLIOZ LQ, TASIMELTEON
HETLIOZ, TASIMELTEON

VELOXIS PHARMS INC

* VELOXIS PHARMACEUTICALS INC
ENVARUS XR, TACROLIMUS

VERITY

* VERITY PHARMACEUTICALS INC
TRELSTAR, TRIPTORELIN PAMOATE

VERO BIOTECH INC

* VERO BIOTECH INC
GENOSYL, NITRIC OXIDE

VEROSCIENCE

* VEROSCIENCE LLC
CYCLOSET, BROMOCRIPTINE MESYLATE

VERRICA PHARMS

* VERRICA PHARMACEUTICALS INC
YCANTH, CANTHARIDIN

VERTEX PHARMS

* VERTEX PHARMACEUTICALS INC
KALYDECO, IVACAFTOR

VERTEX PHARMS INC

* VERTEX PHARMACEUTICALS INC
KALYDECO, IVACAFTOR
ORKAMBI, IVACAFTOR
SYMDEKO (COPACKAGED), IVACAFTOR
TRIKAFTA (COPACKAGED), ELEXACAFTOR, IVACAFTOR, TEZACAFTOR

VERTICAL PHARMS

* VERTICAL PHARMACEUTICALS LLC
DIVIGEL, ESTRADIOL
DSUVIA, SUFENTANIL CITRATE

VIATRIS

* VIATRIS SPECIALTY LLC
CARDURA XL, DOXAZOSIN MESYLATE
CARDURA, DOXAZOSIN MESYLATE
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
GEODON, ZIPRASIDONE HYDROCHLORIDE
GEODON, ZIPRASIDONE MESYLATE
NEURONTIN, GABAPENTIN
NITROSTAT, NITROGLYCERIN
NORVASC, AMLODIPINE BESYLATE
REVATIO, SILDENAFIL CITRATE
VIAGRA, SILDENAFIL CITRATE
ZOLOFT, SERTRALINE HYDROCHLORIDE

VICURON HOLDINGS

* VICURON HOLDINGS LLC
ERAXIS, ANIDULAFUNGIN

VIFOR FRESENIUS

* VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA FRANCE
VELPHORO, FERRIC OXYHYDROXIDE

VIFOR PHARMA

* VIFOR PHARMA INC
VELTASSA, PATIROMER SORBITE X CALCIUM

VIIV HLTHCARE

* VIIV HEALTHCARE CO
APRETUDE, CABOTEGRAVIR
CABENUVA KIT, CABOTEGRAVIR
COMBIVIR, LAMIVUDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VIIV HEALTHCARE CO
 DOVATO, DOLUTEGRAVIR SODIUM
 EPIVIR, LAMIVUDINE
 EPZICOM, ABACAVIR SULFATE
 JULUCA, DOLUTEGRAVIR SODIUM
 LEXIVA, FOSAMPRENAVIR CALCIUM
 RETROVIR, ZIDOVUDINE
 RUKOBIA, FOSTEMSAVIR TROMETHAMINE
 SELZENTRY, MARAVIROC
 TIVICAY PD, DOLUTEGRAVIR SODIUM
 TIVICAY, DOLUTEGRAVIR SODIUM
 TRIUMEQ PD, ABACAVIR SULFATE
 TRIUMEQ, ABACAVIR SULFATE
 VOCABRIA, CABOTEGRAVIR SODIUM
 ZIAGEN, ABACAVIR SULFATE

VINTAGE

* VINTAGE PHARMACEUTICALS LLC
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 DUTASTERIDE, DUTASTERIDE

VINTAGE PHARMS

* VINTAGE PHARMACEUTICALS
 GILDAGIA, ETHINYL ESTRADIOL
 GILDESS 24 FE, ETHINYL ESTRADIOL
 KIMIDESS, DESOGESTREL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

* VINTAGE PHARMACEUTICALS INC
 ALLOPURINOL, ALLOPURINOL
 BACLOFEN, BACLOFEN
 IBUPROFEN, IBUPROFEN (OTC)
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 PERPHENAZINE, PERPHENAZINE
 PREDNISONE, PREDNISONE

VINTAGE PHARMS LLC

* VINTAGE PHARMACEUTICALS LLC
 FELODIPINE, FELODIPINE
 MYZILRA, ETHINYL ESTRADIOL
 PERCOCET, ACETAMINOPHEN
 PREVIFEM, ETHINYL ESTRADIOL

VIRTUS

* VIRTUS PHARMACEUTICALS LLC
 CROMOLYN SODIUM, CROMOLYN SODIUM
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PROMETRIUM, PROGESTERONE

VIRTUS PHARM

* VIRTUS PHARMACEUTICAL INC
 ACARBOSE, ACARBOSE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE

VISTA PHARMS

* VISTA PHARMACEUTICALS INC
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

VISTAPHARM

* VISTAPHARM INC
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 PHENYTOIN, PHENYTOIN
 SUCRALFATE, SUCRALFATE

VISUM PHARM

* VISUM PHARMACEUTICAL CO LTD
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

VITRUVIAS

* VITRUVIAS THERAPEUTICS INC
 ENTECAVIR, ENTECAVIR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ******VITRUVIAS THERAP**

* VITRUVIAS THERAPEUTICS LLC
CYANOCOBALAMIN, CYANOCOBALAMIN

VIVUS

* VIVUS INC
QSYMIA, PHENTERMINE HYDROCHLORIDE

VIWIT PHARM

* VIWIT PHARMACEUTICAL CO LTD
LEVETIRACETAM, LEVETIRACETAM
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE

VKT PHARMA

* VKT PHARMA PRIVATE LTD
FAMOTIDINE, FAMOTIDINE (OTC)
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANOLAZINE, RANOLAZINE
TADALAFIL, TADALAFIL

VPI PHARMS INC

* VPI PHARMACEUTICALS INC
ETHACRYNATE SODIUM, ETHACRYNATE SODIUM

VPNA

* VALEANT PHARMACEUTICALS NORTH AMERICA
DICLOFENAC SODIUM, DICLOFENAC SODIUM

**** W ******WA UNIV SCH MED**

* WASHINGTON UNIV SCHOOL MEDICINE
AMMONIA N 13, AMMONIA N-13

WANBANG BIOPHARMS

* WANBANG BIOPHARMACEUTICALS
CARISOPRODOL, CARISOPRODOL
LEFLUNOMIDE, LEFLUNOMIDE

WATSON LABS

* WATSON LABORATORIES
FOLIC ACID, FOLIC ACID
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

* WATSON LABORATORIES INC
ACARBOSE, ACARBOSE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALLOPURINOL, ALLOPURINOL
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMOXAPINE, AMOXAPINE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
CARISOPRODOL, CARISOPRODOL
CHLORZOXAZONE, CHLORZOXAZONE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
COL-PROBENECID, COLCHICINE
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
ESTAZOLAM, ESTAZOLAM
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GLIPIZIDE, GLIPIZIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LISINAPRIL, LISINAPRIL
LORAZEPAM, LORAZEPAM
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METRONIDAZOLE, METRONIDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WATSON LABORATORIES INC
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 NABUMETONE, NABUMETONE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NATEGLINIDE, NATEGLINIDE
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 NIZATIDINE, NIZATIDINE
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISOLONE, PREDNISOLONE
 PREDNISONE, PREDNISONE
 PRIMIDONE, PRIMIDONE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 QUASENSE, ETHINYL ESTRADIOL
 RAMIPRIL, RAMIPRIL
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SULFASALAZINE, SULFASALAZINE
 SULINDAC, SULINDAC
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOVIA 1/50E-28, ETHINYL ESTRADIOL

* WATSON LABS INC
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

WATSON LABS INC

* WATSON LABORATORIES INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMBRISENTAN, AMBRISENTAN
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 BOSENTAN, BOSENTAN
 BRINZOLAMIDE, BRINZOLAMIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 METRONIDAZOLE, METRONIDAZOLE
 MODAFINIL, MODAFINIL
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 PENICILLAMINE, PENICILLAMINE
 PERPHENAZINE, PERPHENAZINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 TADALAFIL, TADALAFIL

WATSON LABS TEVA

* WATSON LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ALVIMOPAN, ALVIMOPAN
 BICALUTAMIDE, BICALUTAMIDE
 BUPRENORPHINE, BUPRENORPHINE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 GLIPIZIDE, GLIPIZIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 ISRADIPINE, ISRADIPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WATSON LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 PROBENECID, PROBENECID
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

WATSON PHARMS INC

* WATSON PHARMACEUTICALS INC
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

WAVERLEY PHARMA INC

* WAVERLEY PHARMA INC
 BORTEZOMIB, BORTEZOMIB
 PEMETREXED DISODIUM, PEMETREXED DISODIUM

WAYLIS THERAP

* WAYLIS THERAPEUTICS LLC
 FLUTAMIDE, FLUTAMIDE
 LEUKERAN, CHLORAMBUCIL
 MYLERAN, BUSULFAN
 THIOGUANINE, THIOGUANINE
 VALIUM, DIAZEPAM

WES PHARMA INC

* WES PHARMA INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

WEST WARD

* WEST WARD PHARMACEUTICAL CORP
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

WEST WARD PHARM CORP

* WEST WARD PHARMACEUTICAL CORP
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

WEST-WARD PHARMS INT

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 BUMETANIDE, BUMETANIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DIGOXIN, DIGOXIN
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DOXYCYCLINE HCLATE, DOXYCYCLINE HCLATE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PENTOSTATIN, PENTOSTATIN
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, FERRIC OXYHYDROXIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA

WESTMINSTER PHARMS

* WESTMINSTER PHARMACEUTICALS LLC
 EPLERENONE, EPLERENONE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MIGLITOL, MIGLITOL
 POSACONAZOLE, POSACONAZOLE
 RISPERIDONE, RISPERIDONE

WILSHIRE PHARMS INC

* WILSHIRE PHARMACEUTICALS INC
 CARISOPRODOL, CARISOPRODOL
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 PERPHENAZINE, PERPHENAZINE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE

WINDER LABS LLC

* WINDER LABORATOIRES LLC
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE

WISCONSIN

* WISCONSIN MEDICAL RADIOPHARMACY LLC
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WOCKHARDT

* WOCKHARDT LTD
 CAPTOPRIL, CAPTOPRIL
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, LANSOPRAZOLE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

WOCKHARDT BIO AG

* WOCKHARDT BIO AG
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACETAMINOPHEN, ACETAMINOPHEN
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DECITABINE, DECITABINE
 ENTACAPONE, ENTACAPONE
 GENERLAC, LACTULOSE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LISINAPRIL, LISINAPRIL
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 NYSTATIN, NYSTATIN
 OXACILLIN SODIUM, OXACILLIN SODIUM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WOCKHARDT BIO AG
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

WOCKHARDT LTD

* WOCKHARDT LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE

WOODWARD

* WOODWARD PHARMA SERVICES LLC
 ACIPHEX, RABEPRAZOLE SODIUM
 AVODART, DUTASTERIDE
 COREG CR, CARVEDILOL PHOSPHATE
 COREG, CARVEDILOL
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 JALYN, DUTASTERIDE
 LOVAZA, OMEGA-3-ACID ETHYL ESTERS
 LUNESTA, ESZOPICLONE
 MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
 MIVACURIUM CHLORIDE, MIVACURIUM CHLORIDE

WRASER PHARMS

* WRASER PHARMACEUTICALS LLC
 CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE

WUSM CYCLOTRON

* WASHINGTON UNIV SCH MEDICINE CYCLOTRON FACILITY
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WYETH PHARMS

* WYETH PHARMACEUTICALS LLC
 DUAVEE, BAZEDOXIFENE ACETATE
 PREMARIN, ESTROGENS, CONJUGATED
 PREMPHASE 14/14, ESTROGENS, CONJUGATED
 PREMPRO, ESTROGENS, CONJUGATED
 PROTONIX IV, PANTOPRAZOLE SODIUM
 PROTONIX, PANTOPRAZOLE SODIUM
 TRECATOR, ETHIONAMIDE

**** X ******X-GEN PHARMS INC**

* X-GEN PHARMACEUTICALS INC
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

XELLIA PHARMS APS

* XELLIA PHARMACEUTICALS APS
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DAPTOMYCIN, DAPTOMYCIN
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

XERIS

* XERIS PHARMACEUTICALS INC
 GVOKE HYPOPEN, GLUCAGON
 GVOKE KIT, GLUCAGON
 GVOKE PFS, GLUCAGON
 KEVEYIS, DICHLORPHENAMIDE

XGEN PHARMS

* XGEN PHARMACEUTICALS DJB INC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 AMPHOTERICIN B, AMPHOTERICIN B
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DACTINOMYCIN, DACTINOMYCIN
 FOLIC ACID, FOLIC ACID
 IBUPROFEN LYSINE, IBUPROFEN LYSINE
 LINCOMYCIN HYDROCHLORIDE, LINCOMYCIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ******* XGEN PHARMACEUTICALS DJB INC**

LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID

XIAMEN LP PHARM CO*** XIAMEN LP PHARMACEUTICAL CO LTD**

CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

XIROMED*** XIROMED PHARMA ESPANA SL**

ACYCLOVIR, ACYCLOVIR
 ALTAVERA, ETHINYL ESTRADIOL
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 ENILLORING, ETHINYL ESTRADIOL
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 ESTARYLLA, ETHINYL ESTRADIOL
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 FLUOCINONIDE, FLUOCINONIDE
 FOSFOMYCIN TROMETHAMINE, FOSFOMYCIN TROMETHAMINE
 FULVESTRANT, FULVESTRANT
 GEMMILY, ETHINYL ESTRADIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 INTROVALE, ETHINYL ESTRADIOL
 ISIBLOOM, DESOGESTREL
 JAIMIESS, ETHINYL ESTRADIOL
 KETOCONAZOLE, KETOCONAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LORYNA, DROSPIRENONE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 OMEPRAZOLE, OMEPRAZOLE
 PROGESTERONE, PROGESTERONE
 RAMELTEON, RAMELTEON
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SYEDA, DROSPIRENONE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE, TESTOSTERONE
 TRI-ESTARYLLA, ETHINYL ESTRADIOL
 TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
 VIENVA, ETHINYL ESTRADIOL
 VOLNEA, DESOGESTREL

XSPIRE PHARMA*** XSPIRE PHARMA**

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 NALFON, FENOPROFEN CALCIUM
 TREZIX, ACETAMINOPHEN
 ZONTIVITY, VORAPAXAR SULFATE

*** XSPIRE PHARMA LLC**

DEXAMETHASONE, DEXAMETHASONE
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM

XTTRIUM*** XTTRIUM LABORATORIES INC**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ******XTTRIUM LABS INC**

* XTTRIUM LABORATORIES INC
LACTULOSE, LACTULOSE

**** Y ******YABAO PHARM**

* YABAO PHARMACEUTICAL CO LTD BEIJING
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
SORAFENIB TOSYLATE, SORAFENIB TOSYLATE

YAOPHARMA CO LTD

* YAOPHARMA CO LTD
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

YICHANG HUMANWELL

* YICHANG HUMANWELL PHARMACEUTICAL CO LTD
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
FENOFIBRIC ACID, CHOLINE FENOFIBRATE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

YILING

* YILING PHARMACEUTICAL LTD
ACYCLOVIR, ACYCLOVIR
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CELECOXIB, CELECOXIB
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
FELODIPINE, FELODIPINE
LAMOTRIGINE, LAMOTRIGINE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PREGABALIN, PREGABALIN
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

YOUNGTECH PHARMS INC

* YOUNGTECH PHARMACEUTICALS INC
METOPROLOL TARTRATE, METOPROLOL TARTRATE

YUNG SHIN PHARM

* YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
AZITHROMYCIN, AZITHROMYCIN
CEFACLOR, CEFACLOR
CEPHALEXIN, CEPHALEXIN
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
FELODIPINE, FELODIPINE
MELOXICAM, MELOXICAM

**** Z ******ZAMBON SPA**

* ZAMBON SPA ITALY
MONUROL, FOSFOMYCIN TROMETHAMINE

ZEALAND PHARMA

* ZEALAND PHARMA US INC
ZEGALOGUE (AUTOINJECTOR), DASIGLUCAGON HYDROCHLORIDE
ZEGALOGUE, DASIGLUCAGON HYDROCHLORIDE

ZENNOVA

* ZENNOVA PHARMACEUTICALS CHENGDU CO LTD
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

ZHEJIANG HISUN PHARM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZHEJIANG HISUN PHARMACEUTICAL CO LTD
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

ZHEJIANG JINGXIN

* ZHEJIANG JINGXIN PHARMACEUTICAL CO LTD
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM

ZHEJIANG JIUZHOU

* ZHEJIANG JIUZHOU BIOPHARMA CO LTD
CARBAMAZEPINE, CARBAMAZEPINE

ZHEJIANG JUTAI PHARM

* ZHEJIANG JUTAI PHARMACEUTICAL CO LTD
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE

ZHEJIANG NOVUS PHARM

* ZHEJIANG NOVUS PHARMACEUTICALS CO LTD
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ZHEJIANG YONGTAI

* ZHEJIANG YONGTAI PHARMACEUTICAL CO LTD
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
GABAPENTIN, GABAPENTIN
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

ZR PHARMA

* ZR PHARMA AND GMBH
RUBRACA, RUCAPARIB CAMSYLATE

ZUREX PHARMA

* ZUREX PHARMA
ZURAGARD, ISOPROPYL ALCOHOL (OTC)

ZYDUS

* ZYDUS WORLDWIDE DMCC
ACETAZOLAMIDE, ACETAZOLAMIDE
AZITHROMYCIN, AZITHROMYCIN
BACLOFEN, BACLOFEN
BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
ERYTHROMYCIN, ERYTHROMYCIN
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
ICOSAPENT ETHYL, ICOSAPENT ETHYL
INDOMETHACIN, INDOMETHACIN
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
IVERMECTIN, IVERMECTIN
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LEFLUNOMIDE, LEFLUNOMIDE
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
METHYLENE BLUE, METHYLENE BLUE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
OXCARBAZEPINE, OXCARBAZEPINE
PHYTONADIONE, PHYTONADIONE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROPAPENONE HYDROCHLORIDE, PROPAPENONE HYDROCHLORIDE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
TOPIRAMATE, TOPIRAMATE
URSODIOL, URSODIOL
VARENICLINE TARTRATE, VARENICLINE TARTRATE
VIGABATRIN, VIGABATRIN
ZITUVIMET, METFORMIN HYDROCHLORIDE
ZITUVIO, SITAGLIPTIN

ZYDUS HLTHCARE

* ZYDUS HEALTHCARE USA LLC
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
LANSOPRAZOLE, LANSOPRAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******ZYDUS LIFESCIENCES**

* ZYDUS LIFESCIENCES LTD
 ACYCLOVIR, ACYCLOVIR
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 FEBUXOSTAT, FEBUXOSTAT
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 SUCRALFATE, SUCRALFATE
 TAVABOROLE, TAVABOROLE

ZYDUS NOVELTECH INC

* ZYDUS NOVELTECH INC
 ETHINYL ESTRADIOL AND NORELGESTROMIN, ETHINYL ESTRADIOL

ZYDUS PHARMS

* ZYDUS PHARMACEUTICALS USA INC
 ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ALBENDAZOLE, ALBENDAZOLE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALLOPURINOL, ALLOPURINOL
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMBRISENTAN, AMBRISENTAN
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BORTEZOMIB, BORTEZOMIB
 BOSENTAN, BOSENTAN
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA, CARBIDOPA
 CHLORTHALIDONE, CHLORTHALIDONE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLCHICINE, COLCHICINE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DARUNAVIR, DARUNAVIR
 DECITABINE, DECITABINE
 DEFERASIROX, DEFERASIROX
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE, DOXYCYCLINE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DROXIDOPA, DROXIDOPA
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 ENTECAVIR, ENTECAVIR
 ERLOTINIB HYDROCHLORIDE, ERLOTINIB HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESTRADIOL, ESTRADIOL
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 EXEMESTANE, EXEMESTANE
 EZETIMIBE, EZETIMIBE
 FAMOTIDINE, FAMOTIDINE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FINGOLIMOD HYDROCHLORIDE, FINGOLIMOD HYDROCHLORIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE, FLUOCINONIDE
 FULVESTRANT, FULVESTRANT
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INDOMETHACIN, INDOMETHACIN
 ISOTRETINOIN, ISOTRETINOIN
 ITRACONAZOLE, ITRACONAZOLE
 IVABRADINE HYDROCHLORIDE, IVABRADINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACOSAMIDE, LACOSAMIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LENALIDOMIDE, LENALIDOMIDE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LINEZOLID, LINEZOLID
 LUBIPROSTONE, LUBIPROSTONE
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MICAFUNGIN SODIUM, MICAFUNGIN SODIUM
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NADOLOL, NADOLOL
 NATEGLINIDE, NATEGLINIDE
 NELARABINE, NELARABINE
 NIFEDIPINE, NIFEDIPINE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PEMETREXED DISODIUM, PEMETREXED DISODIUM
 PERPHENAZINE, PERPHENAZINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 PLERIXAFOR, PLERIXAFOR
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RAMELTEON, RAMELTEON
 RIVASTIGMINE, RIVASTIGMINE
 ROFLUMILAST, ROFLUMILAST
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIROLIMUS, SIROLIMUS
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUGAMMADEX SODIUM, SUGAMMADEX SODIUM
 TADALAFIL, TADALAFIL
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIAZOLAM, TRIAZOLAM
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZINC SULFATE, ZINC SULFATE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ZYDUS PHARMS USA

* ZYDUS PHARMACEUTICALS USA INC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATENOLOL, ATENOLOL
 AZATHIOPRINE, AZATHIOPRINE
 BENZONATATE, BENZONATATE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL
 LAMOTRIGINE, LAMOTRIGINE
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 RIBAVIRIN, RIBAVIRIN
 RISPERIDONE, RISPERIDONE
 SIMVASTATIN, SIMVASTATIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

ZYDUS PHARMS USA INC

* ZYDUS PHARMACEUTICALS USA INC
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ANASTROZOLE, ANASTROZOLE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******* ZYDUS PHARMACEUTICALS USA INC**

CARVEDILOL, CARVEDILOL
DIPYRIDAMOLE, DIPYRIDAMOLE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
FINASTERIDE, FINASTERIDE
GABAPENTIN, GABAPENTIN
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
IRBESARTAN, IRBESARTAN
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
OMEPRAZOLE, OMEPRAZOLE
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
ZOLMITRIPTAN, ZOLMITRIPTAN

ZYLA*** ZYLA LIFE SCIENCES US INC**

INDOCIN, INDOMETHACIN
SPRIX, KETOROLAC TROMETHAMINE

APPENDIX C**UNIFORM TERMS*****DOSAGE FORMS***

AEROSOL, FOAM	PASTE
AEROSOL, METERED	PATCH
CAPSULE	PELLET
CAPSULE, DELAYED REL PELLETS	PELLETS
CAPSULE, DELAYED RELEASE	POWDER
CAPSULE, EXTENDED RELEASE	POWDER, EXTENDED RELEASE
CAPSULE, PELLETS	POWDER, METERED
CAPSULE, TABLET	RING
CAPSULE, TABLET, TABLET	SHAMPOO
CLOTH	SOLUTION
CONCENTRATE	SOLUTION FOR SLUSH
CREAM	SOLUTION, EXTENDED RELEASE
CREAM, AUGMENTED	SOLUTION, GEL FORMING/DROPS
CREAM, INSERT	SOLUTION, METERED
ELIXIR	SOLUTION/DROPS
EMULSION	SPONGE
ENEMA	SPRAY
FILM	SPRAY, METERED
FILM, EXTENDED RELEASE	SUPPOSITORY
FOAM	SUSPENSION
FOR SOLUTION	SUSPENSION, EXTENDED RELEASE
FOR SUSPENSION	SUSPENSION, LIPOSOMAL
FOR SUSPENSION, DELAYED RELEASE	SUSPENSION/DROPS
FOR SUSPENSION, EXTENDED RELEASE	SWAB
GAS	SYRUP
GEL	SYSTEM
GEL, AUGMENTED	TABLET
GEL, METERED	TABLET, CHEWABLE
GRANULE	TABLET, DELAYED RELEASE
GRANULE, DELAYED RELEASE	TABLET, EFFERVESCENT
GRANULES	TABLET, EXTENDED RELEASE
GRANULES, EXTENDED RELEASE	TABLET, EXTENDED RELEASE, CHEWABLE
GUM, CHEWING	TABLET, FOR SUSPENSION
IMPLANT	TABLET, ORALLY DISINTEGRATING
INHALANT	TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE
INJECTABLE	TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE
INJECTABLE, LIPID COMPLEX	TAPE
INJECTABLE, LIPOSOMAL	TROCHE/LOZENGE
INJECTION, EXTENDED RELEASE	
INSERT	
INSERT, EXTENDED RELEASE	
INTRAUTERINE DEVICE	
JELLY	
LIQUID	
LOTION	
LOTION, AUGMENTED	
LOTION/SHAMPOO	
OIL	
OIL/DROPS	
OINTMENT	
OINTMENT, AUGMENTED	

APPENDIX C**UNIFORM TERMS*****ROUTES OF ADMINISTRATION***

BUCCAL	N/A
CARDIAC	NASAL
DENTAL	OPHTHALMIC
ENDOCERVICAL	ORAL
ENDOTRACHEAL	ORAL-21
ENTERAL	ORAL-28
IMPLANTATION	OTIC
INFILTRATION	PERFUSION
INHALATION	PERIARTICULAR
INJECTION	PERIODONTAL
INTERSTITIAL	PYELOCALYCEAL
INTRA-ANAL	RECTAL
INTRA-ARTERIAL	SPINAL
INTRA-ARTICULAR	SUBCUTANEOUS
INTRACAMERAL	SUBLINGUAL
INTRACAVITARY	TOPICAL
INTRACRANIAL	TRANSDERMAL
INTRADERMAL	TRANSMUCOSAL
INTRAMUSCULAR	URETHRAL
INTRAOCULAR	VAGINAL
INTRAOSSEOUS	
INTRAPERITONEAL	
INTRAPLEURAL	
INTRATHECAL	
INTRAUTERINE	
INTRAVENOUS	
INTRAVESICAL	
INTRAVITREAL	
IRRIGATION	
IV (INFUSION)	

APPENDIX C**UNIFORM TERMS*****ABBREVIATIONS***

AMP	AMPULE
AMPICIL	AMPICILLIN
APPROX	APPROXIMATELY
BOT	BOTTLE
CI	CURIE
CSR	CAROTID SINUS REFLEX
CU	CLINICAL UNITS
DIPROP	DIPROPIONATE
ELECT	ELECTROLYTE
EQ	EQUIVALENT TO
ER	EXTENDED RELEASE
GM	GRAM
HBR	HYDROBROMIDE
HCL	HYDROCHLORIDE
HR	HOUR
IM	INTRAMUSCULAR
INH	INHALATION
IU	INTERNATIONAL UNITS
IV	INTRAVENOUS
KIU	KALLIKREIN INHIBITOR UNITS
MCG	MICROGRAM
MCI	MILLICURIE
MEQ	MILLIEQUIVALENT
MG	MILLIGRAM
ML	MILLILITER
N/A	NOT APPLICABLE
PPM	PARTS PER MILLION
REL	RELEASE
SC	SUBCUTANEOUS
SQ CM	SQUARE CENTIMETER
U	UNITS
UCI	MICROCURIE
UMOLAR	MICROMOLAR
USP	UNITED STATES PHARMACOPEIA

PATENT AND EXCLUSIVITY INFORMATION ADDENDUM

This *Addendum* identifies drugs that qualify under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) for periods of exclusivity and provides patent information that has been submitted to the Food and Drug Administration (FDA) concerning the listed drug products.

Exclusivity

This *Addendum* identifies:

- Drugs approved under section 505(c) of the FD&C Act that have qualified under the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments) for five-year and three-year periods of exclusivity pursuant to Section 505(c)(3)(E) and Section 505(j)(5)(F) of the FD&C Act
- Drugs that have qualified for Orphan Drug Exclusivity pursuant to Section 527 of the FD&C Act
- Drugs that have qualified for Pediatric Exclusivity pursuant to Section 505A of the FD&C Act
- Drugs that have qualified for Generating Antibiotic Incentives Now (GAIN) exclusivity pursuant to Section 505E of the FD&C Act
- Generic drugs approved under section 505(j) of the FD&C Act that have qualified for 180-day exclusivity pursuant to Section 505(j)(5)(B)(iv) of the FD&C Act
- Generic drugs approved under section 505(j) of the FD&C Act that have qualified for Competitive Generic Therapy (CGT) exclusivity pursuant to Section 505(j)(5)(B)(v) of the FD&C Act

This section is arranged in alphabetical order by established name of the active ingredient, followed by the proprietary name (brand name or trade name) of the drug product. Active ingredient headings for multiple active ingredient, fixed-combination drug products are arranged alphabetically.

Individual descriptions of the protected use have been added to each Orphan Drug Exclusivity entry listed in the Orange Book. Such descriptions of Orphan Drug Exclusivity were included beginning with the 38th edition of the Orange Book. The ODE* code means that the timing of approval of certain follow-on applications may be subject to delay due to ODE for another drug that has the same active moiety.

For an explanation of the codes used in the *Addendum*, see the *Patent and Exclusivity Terms* Section. The exclusivity codes are general shorthand descriptions and do not necessarily identify, with specificity, the actual scope of exclusivity.

Patent Information

The FD&C Act requires that patent information be filed with all newly submitted Section 505(b) drug applications. In addition, patent information must be filed on Form FDA 3542 within 30 days of the date of approval of a Section 505(b) drug application.¹ FDA publishes certain information from Form FDA 3542 in the Orange Book after approval of the new drug application (NDA) or supplement.

The Orange Book includes the patent submission date (i.e., the date on which the FDA receives patent information from the NDA holder on Form FDA 3542) for each newly listed patent to facilitate assessments of whether patent information is untimely filed with respect to a pending 505(b)(2) application or abbreviated new drug application (ANDA) and whether patent information was submitted before the date on which a 505(b)(2) application or ANDA (excluding an amendment or supplement to the 505(b)(2) application or ANDA) was submitted.²

The patents that FDA regards as covered by the statutory provisions for submission of patent information for listing in the Orange Book are:

- Patents that claim the drug for which the applicant submitted the application and are drug substance (active ingredient) patents or drug product (formulation or composition) patents; or
- Patents that claim a method of using such drug for which approval has been granted in the application.

This information, as provided by the NDA holder on Form FDA 3542, will be published as described above. An NDA holder submitting information on a patent that claims both the drug substance and the drug product (and is eligible for listing on either basis) is required only to specify that the patent claims either the drug substance or the drug product.

The Addendum lists patent and exclusivity information up to January of the Edition year. The monthly Cumulative Supplements to the Annual Edition list patent and exclusivity information changes since the Annual Edition Addendum. Since all parts of this publication are subject to changes, additions, or deletions, the [Orange Book](#), updated regularly, should be consulted for the most recent patent and exclusivity information.

¹ Please note that the date of approval for an NDA for a drug for which FDA intends to recommend controls under the Controlled Substances Act is the later of the date on the approval letter for the NDA or the date of issuance of the interim final rule controlling the drug (see Section 505(x)(1) and (2) of the FD&C Act).

² See 21 CFR 314.50(i)(4) and 314.94(a)(12)(vi); see 21 CFR 314.107(b)(3)(i). The submission date for patent information is determined in accordance with 21 CFR 314.53(d)(5).

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE - TRIUMEO</u>						
N 205551 001	8129385	Oct 05, 2027	DS DP		M-294	Jun 15, 2026
	8129385*PED	Apr 05, 2028			PED	Dec 15, 2026
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE - TRIUMEO PD</u>						
N 215413 001	8129385	Oct 05, 2027	DS DP		NPP	Jun 15, 2026
	8129385*PED	Apr 05, 2028			PED	Dec 15, 2026
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>ABALOPARATIDE - TYMLOS</u>						
N 208743 001	10996208	Apr 30, 2038	DP		I-907	Dec 19, 2025
	11255842	Jan 10, 2040	U-3322		M-270	Sep 20, 2024
	11680942	Jan 10, 2040	U-3322			
	11782041	Apr 30, 2038	DP U-2009			
	11782041	Apr 30, 2038	DP U-3543			
	7803770	Apr 28, 2031	U-2009			
	8148333	Nov 08, 2027	DP			
	8748382	Oct 03, 2027	U-2009			
	8748382	Oct 03, 2027	U-3543			
	RE49444	Apr 28, 2031	U-2009			
	RE49444	Apr 28, 2031	U-3543			
<u>ABAMETAPIR - XEGLYZE</u>						
N 206966 001	10292389	Dec 17, 2034	DP U-2863		NCE	Jul 24, 2025
	7812163	Oct 28, 2026	DP U-2863			
	8212038	Jul 16, 2024	DP U-2863			
	9357783	Jul 16, 2024	DP			
	9839631	Jul 16, 2024	DS DP U-2863			
<u>ABEMACICLIB - VERZENIO</u>						
N 208716 001	7855211	Dec 15, 2029	DS DP U-2132		I-877	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2135		NPP	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2251			
	7855211	Dec 15, 2029	DS DP U-3241			
	7855211	Dec 15, 2029	DS DP U-3242			
	7855211	Dec 15, 2029	DS DP U-3243			
	7855211	Dec 15, 2029	DS DP U-3265			
	7855211	Dec 15, 2029	DS DP U-3546			
<u>ABEMACICLIB - VERZENIO</u>						
N 208716 002	7855211	Dec 15, 2029	DS DP U-2132		I-877	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2135		NPP	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2251			
	7855211	Dec 15, 2029	DS DP U-3241			
	7855211	Dec 15, 2029	DS DP U-3242			
	7855211	Dec 15, 2029	DS DP U-3243			
	7855211	Dec 15, 2029	DS DP U-3265			
	7855211	Dec 15, 2029	DS DP U-3546			

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<u>ABEMACICLIB - VERZENIO</u>						
N 208716 003	7855211	Dec 15, 2029	DS DP U-2132		I-877	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2135		NPP	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2251			
	7855211	Dec 15, 2029	DS DP U-3241			
	7855211	Dec 15, 2029	DS DP U-3242			
	7855211	Dec 15, 2029	DS DP U-3243			
	7855211	Dec 15, 2029	DS DP U-3265			
	7855211	Dec 15, 2029	DS DP U-3546			
<u>ABEMACICLIB - VERZENIO</u>						
N 208716 004	7855211	Dec 15, 2029	DS DP U-1981		I-877	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2132		NPP	Oct 12, 2024
	7855211	Dec 15, 2029	DS DP U-2135			
	7855211	Dec 15, 2029	DS DP U-2251			
<u>ABIRATERONE ACETATE - YONSA</u>						
N 210308 001	10292990	May 20, 2034	U-2535			
	9889144	Mar 17, 2034	DP			
<u>ABIRATERONE ACETATE; NIRAPARIB TOSYLATE - AKEEGA</u>						
N 216793 001	11091459	Mar 27, 2038	DS DP		NP	Aug 11, 2026
	11207311	Jul 28, 2037	U-2830			
	11673877	Mar 27, 2038	DS DP			
	8071579	Aug 12, 2027	U-2830			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2830			
	8436185	Apr 24, 2029	DS DP			
	8859562	Aug 04, 2031	U-2830			
<u>ABIRATERONE ACETATE; NIRAPARIB TOSYLATE - AKEEGA</u>						
N 216793 002	11091459	Mar 27, 2038	DS DP		NP	Aug 11, 2026
	11207311	Jul 28, 2037	U-2830			
	11673877	Mar 27, 2038	DS DP			
	8071579	Aug 12, 2027	U-2830			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2830			
	8436185	Apr 24, 2029	DS DP			
	8859562	Aug 04, 2031	U-2830			
<u>ABROCITINIB - CIBINQO</u>						
N 213871 001	9035074	Feb 19, 2034	DS DP		NCE	Jan 14, 2027
	9545405	Feb 19, 2034	DS DP		NPP	Feb 09, 2026
	9549929	Feb 19, 2034	U-3195			
<u>ABROCITINIB - CIBINQO</u>						
N 213871 002	9035074	Feb 19, 2034	DS DP		NCE	Jan 14, 2027
	9545405	Feb 19, 2034	DS DP		NPP	Feb 09, 2026
	9549929	Feb 19, 2034	U-3195			
<u>ABROCITINIB - CIBINQO</u>						
N 213871 003	9035074	Feb 19, 2034	DS DP		NCE	Jan 14, 2027
	9545405	Feb 19, 2034	DS DP		NPP	Feb 09, 2026
	9549929	Feb 19, 2034	U-3195			

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<u>ACALABRUTINIB - CALQUENCE</u>						
N 210259 001	10167291	Jul 01, 2036	DP U-2145		ODE-175	Oct 31, 2024
	10167291	Jul 01, 2036	DP U-2666		ODE-274	Nov 21, 2026
	10167291	Jul 01, 2036	DP U-2667			
	10167291	Jul 01, 2036	DP U-2668			
	10167291	Jul 01, 2036	DP U-2669			
	10167291	Jul 01, 2036	DP U-2670			
	10167291	Jul 01, 2036	DP U-2671			
	10239883	Jul 11, 2032	U-2666			
	10239883	Jul 11, 2032	U-2668			
	10272083	Jan 21, 2035	U-2519			
	10272083	Jan 21, 2035	U-2682			
	10272083	Jan 21, 2035	U-2683			
	10272083	Jan 21, 2035	U-2684			
	10272083	Jan 21, 2035	U-2685			
	10272083	Jan 21, 2035	U-2686			
	10272083	Jan 21, 2035	U-2687			
	11771696	Jan 21, 2035	U-3710			
	7459554	Nov 24, 2026	DS			
	9290504	Jul 11, 2032	DS DP			
	9758524	Jul 11, 2032	U-2145			
	9796721	Jul 01, 2036	DS DP U-2145			
	9796721	Jul 01, 2036	DS DP U-2666			
	9796721	Jul 01, 2036	DS DP U-2667			
	9796721	Jul 01, 2036	DS DP U-2668			
	9796721	Jul 01, 2036	DS DP U-2669			
	9796721	Jul 01, 2036	DS DP U-2670			
	9796721	Jul 01, 2036	DS DP U-2671			
<u>ACALABRUTINIB MALEATE - CALQUENCE</u>						
N 216387 001	10239883	Jul 11, 2032	U-2666			
	10239883	Jul 11, 2032	U-2668			
	10272083	Jan 21, 2035	U-2519			
	10272083	Jan 21, 2035	U-2682			
	10272083	Jan 21, 2035	U-2683			
	10272083	Jan 21, 2035	U-2684			
	10272083	Jan 21, 2035	U-2685			
	10272083	Jan 21, 2035	U-2686			
	10272083	Jan 21, 2035	U-2687			
	11059829	Jul 01, 2036	DS DP U-2145			
	11059829	Jul 01, 2036	DS DP U-2666			
	11059829	Jul 01, 2036	DS DP U-2667			
	11059829	Jul 01, 2036	DS DP U-2668			
	11059829	Jul 01, 2036	DS DP U-2669			
	11059829	Jul 01, 2036	DS DP U-2670			
	11059829	Jul 01, 2036	DS DP U-2671			
	11771696	Jan 21, 2035	U-3710			
	7459554	Nov 24, 2026	DS			
	9290504	Jul 11, 2032	DS DP			
	9758524	Jul 11, 2032	U-2145			

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<u>ACETAMINOPHEN - OFIRMEV</u>						
N 022450	001 10383834	Nov 13, 2028	U-2262			
	10383834	Nov 13, 2028	U-2621			
	9399012	Sep 11, 2031	U-2261			
	9399012	Sep 11, 2031	U-2262			
	9399012*PED	Mar 11, 2032				
	9610265	Nov 13, 2028	U-2263			
	9610265*PED	May 13, 2029				
	9987238	Nov 13, 2028	U-2261			
	9987238*PED	May 13, 2029				
<u>ACETAMINOPHEN - ACETAMINOPHEN</u>						
N 204767	001 8741959	Apr 19, 2030	DP			
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	001 8461137	Feb 22, 2031	DS DP			
	8748413	Jul 01, 2030	DS DP			
	8828978	Jul 01, 2030	DP			
	9132125	Jul 01, 2030	DS DP	U-2249		
	9549923	Jul 01, 2030	DS DP			
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	002 8461137	Feb 22, 2031	DS DP			
	8748413	Jul 10, 2030	DS DP			
	8828978	Jul 01, 2030	DP			
	9132125	Jul 01, 2030	DS DP	U-2249		
	9549923	Jul 01, 2030	DS DP			
<u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u>						
N 208653	003 8461137	Feb 22, 2031	DS DP			
	8748413	Jul 10, 2030	DS DP			
	8828978	Jul 01, 2030	DP			
	9132125	Jul 01, 2030	DS DP	U-2249		
	9549923	Jul 01, 2030	DS DP			
<u>ACETAMINOPHEN; IBUPROFEN - COMBOGESIC</u>						
N 209471	001 10532036	Sep 22, 2025	U-3553		NP	Mar 01, 2026
	11197830	Feb 27, 2039	DP U-3553			
	11534407	Feb 27, 2039	DP U-3553			
<u>ACETAMINOPHEN; IBUPROFEN SODIUM - COMBOGESIC IV</u>						
N 215320	001 11213498	Jan 14, 2036	DP		NP	Oct 17, 2026
	11389416	Jul 17, 2035	DP			
	11446266	Oct 26, 2031	U-3744			
	11446266	Oct 26, 2031	U-3745			
<u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u>						
N 204031	001 7976870	Jun 01, 2027	U-1498			
	8372432	Mar 11, 2029	DP U-1499			
	8377453	Nov 19, 2029	DP U-1499			
	8394408	Mar 11, 2029	DP			
	8597681	Dec 21, 2030	DP			
	8658631	May 16, 2032	DP			
	8668929	Mar 11, 2029	U-1499			
	8741885	May 16, 2032	DP U-1499			

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<u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u>						
N 204031	001	8980319	Dec 21, 2030	DP		
		8992975	May 16, 2032	DP		
		9050335	May 16, 2032	DP		
		9468636	May 16, 2032	U-1499		
<u>ACETYLCYSTEINE - ACETADOTE</u>						
N 021539	001	8148356	May 21, 2026	DP		
		8399445	Aug 24, 2025	U-1373		
		8653061	Aug 24, 2025	U-1373		
		8722738	Apr 06, 2032	U-1373		
		9327028	Jul 21, 2031	U-1839		
<u>ACETYLCYSTEINE - CETYLEV</u>						
N 207916	001	8747894	May 08, 2032	DP U-1373		
		9427421	May 08, 2032	DP		
		9561204	May 08, 2032	U-1373		
<u>ACETYLCYSTEINE - CETYLEV</u>						
N 207916	002	8747894	May 08, 2032	DP U-1373		
		9427421	May 08, 2032	DP		
		9561204	May 08, 2032	U-1373		
<u>ACLIDINIUM BROMIDE - TUDORZA PRESSAIR</u>						
N 202450	001	10085974	Mar 13, 2029	DP U-2513		
		11000517	Mar 13, 2029	DP U-2513		
		8051851	Apr 22, 2027	DP		
		RE46417	Feb 10, 2025	DS DP U-2513		
<u>ACLIDINIUM BROMIDE; FORMOTEROL FUMARATE - DUAKLIR PRESSAIR</u>						
N 210595	001	10085974	Mar 13, 2029	DP U-2513		
		11000517	Mar 13, 2029	DP U-2513		
		8051851	Apr 22, 2027	DP		
		RE46417	Feb 10, 2025	DS DP U-2513		
<u>ACYCLOVIR - AVACLYR</u>						
N 202408	001				ODE-235	Mar 29, 2026
<u>ACYCLOVIR - SITAVIG</u>						
N 203791	001	8592434	Jun 16, 2030	DP U-1460		
		8747896	Jun 03, 2027	DP U-1460		
		8791127	Mar 23, 2027	DP U-1460		
<u>ADAGRASIB - KRAZATI</u>						
N 216340	001	10689377	May 17, 2037	DS DP U-3490	NCE	Dec 12, 2027
					ODE-352	Dec 12, 2029
<u>ADAPALENE - DIFFERIN</u>						
N 021753	001	7579377	Feb 23, 2025	U-818		
		7737181	Aug 29, 2024	DP		
<u>ADAPALENE - DIFFERIN</u>						
N 022502	001	7998467	May 31, 2028	DP U-1078		
		8435502	Sep 15, 2026	DP U-1078		
		8709392	Sep 15, 2026	DP U-1078		

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<u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO</u>						
N 022320	001 7820186	Nov 23, 2025	DP			
	7964202	Sep 01, 2024	DP U-1078			
	8071644	Jul 18, 2027	DP U-1078			
	8080537	Jul 18, 2027	U-1078			
	8129362	Jul 18, 2027	U-1078			
<u>ADAPALENE; BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - CABTREO</u>						
N 216632	001 10220049	Jun 03, 2029	DP U-3713		NP	Oct 20, 2026
	10624918	Jun 03, 2029	U-3713			
	11389467	Dec 28, 2040	DP U-3713			
	8288434	Aug 05, 2029	DP U-3713			
	9561208	Jun 03, 2029	DP U-3713			
<u>AFAMELANOTIDE - SCENESSE</u>						
N 210797	001 10076555	Feb 11, 2025	U-2638		NCE	Oct 08, 2024
	8334265	Mar 11, 2029	U-2638		ODE-270	Oct 08, 2026
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	001 10004743	Jul 05, 2030	DP		M-276	Apr 07, 2025
	10004743*PED	Jan 05, 2031			ODE-230	Jan 12, 2025
	8426586	Oct 10, 2029	DS		PED	Jul 12, 2025
	8426586*PED	Apr 10, 2030			PED	Oct 07, 2025
	8545884	Dec 19, 2029	DP			
	8545884*PED	Jun 19, 2030				
	9539258	Nov 09, 2026	U-1950			
	9539258*PED	May 09, 2027				
	RE43431	Jan 13, 2026	DS			
	RE43431*PED	Jul 13, 2026				
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	002 10004743	Jul 05, 2030	DP		M-276	Apr 07, 2025
	10004743*PED	Jan 05, 2031			ODE-230	Jan 12, 2025
	8426586	Oct 10, 2029	DS		PED	Jul 12, 2025
	8426586*PED	Apr 10, 2030			PED	Oct 07, 2025
	8545884	Dec 19, 2029	DP			
	8545884*PED	Jun 19, 2030				
	9539258	Nov 09, 2026	U-1950			
	9539258*PED	May 09, 2027				
	RE43431	Jan 13, 2026	DS			
	RE43431*PED	Jul 13, 2026				
<u>AFATINIB DIMALEATE - GILOTRIF</u>						
N 201292	003 10004743	Jul 05, 2030	DP		M-276	Apr 07, 2025
	10004743*PED	Jan 05, 2031			ODE-230	Jan 12, 2025
	8426586	Oct 10, 2029	DS		PED	Jul 12, 2025
	8426586*PED	Apr 10, 2030			PED	Oct 07, 2025
	8545884	Dec 19, 2029	DP			
	8545884*PED	Jun 19, 2030				
	9539258	Nov 09, 2026	U-1950			
	9539258*PED	May 09, 2027				
	RE43431	Jan 13, 2026	DS			
	RE43431*PED	Jul 13, 2026				

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<u>AIR POLYMER-TYPE A - EXEM FOAM KIT</u>						
N 212279	001 9034300	Oct 15, 2030	DP U-2663		NCE	Nov 07, 2024
	9259494	May 04, 2035	DP U-2663			
	9849199	Feb 11, 2036	DP			
<u>ALBUTEROL SULFATE - PROAIR HEA</u>						
N 021457	001 10022509	May 18, 2031	DP			
	10022510	May 18, 2031	DP			
	10086156	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	11395889	May 18, 2031	DP			
	8132712	Sep 07, 2028	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			
<u>ALBUTEROL SULFATE - PROAIR RESPICLICK</u>						
N 205636	001 10022510	May 18, 2031	DP			
	10124131	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10765820	May 19, 2025	DP			
	8651103	Mar 26, 2028	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9731087	May 18, 2031	DP			
<u>ALBUTEROL SULFATE - PROAIR DIGIHALER</u>						
N 205636	002 10022510	May 18, 2031	DP			
	10124131	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10569034	Aug 16, 2036	DP			
	10765820	May 19, 2025	DP			
	10918816	Dec 14, 2035	DP			
	11000653	Dec 18, 2038	DP			
	11173259	Jul 06, 2040	DP			
	11266796	Feb 22, 2041	DP			
	11344685	Sep 26, 2039	DP			
	11351317	Feb 10, 2038	DP			
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	8651103	Mar 26, 2028	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9731087	May 18, 2031	DP			
	9782550	Aug 28, 2035	DP			
	9782551	Aug 28, 2035	DP			
<u>ALBUTEROL SULFATE; BUDESONIDE - AIRSUPRA</u>						
N 214070	001 9415009	May 28, 2030	U-3509		NP	Jan 10, 2026

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<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - COMBIVENT RESPIMAT</u>						
N 021747	001 7284474	Aug 26, 2024	DP			
	7396341	Oct 10, 2026	DP			
	7837235	Mar 13, 2028	DP			
	7896264	May 26, 2025	DP			
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
<u>ALCAFTADINE - LASTACAPT</u>						
N 022134	001 10617695	Mar 19, 2027	DP U-3267			
	8664215	Dec 23, 2027	U-3267			
<u>ALCOHOL - ABLYSINOL</u>						
N 207987	001				ODE-192	Jun 21, 2025
<u>ALCOHOL - ABLYSINOL</u>						
N 207987	002				ODE-192	Jun 21, 2025
<u>ALECTINIB HYDROCHLORIDE - ALECENSA</u>						
N 208434	001 10350214	Apr 24, 2035	DP		ODE-159	Nov 06, 2024
	11433076	Apr 24, 2035	DP			
	9126931	May 29, 2031	DS			
	9365514	Mar 04, 2032	DP			
	9440922	Jun 09, 2030	DP			
<u>ALENDRONATE SODIUM - BINOSTO</u>						
N 202344	001 9592195	Dec 05, 2031	DP			
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>						
N 021985	001 8617595	Feb 19, 2026	DP			
	8617595*PED	Aug 19, 2026				
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>						
N 021985	002 8617595	Feb 19, 2026	DP			
	8617595*PED	Aug 19, 2026				
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	001 8613949	Dec 21, 2029	DP			
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	002 8613949	Dec 21, 2029	DP			
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	003 8613949	Dec 21, 2029	DP			
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>						
N 022545	004 8613949	Dec 21, 2029	DP			
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	001 8618172	Jul 13, 2028	DP			
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	002 8618172	Jul 13, 2028	DP			
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	003 8618172	Jul 13, 2028	DP			
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N 022107	004 8618172	Jul 13, 2028	DP			

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<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N 022217	001	8168616	Jul 03, 2026	DP		
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N 022217	002	8168616	Jul 03, 2026	DP		
<u>ALLOPURINOL; LESINURAD - DUZALLO</u>						
N 209203	001	10183012	Nov 26, 2028		U-2104	
		8003681	Aug 25, 2025	DS		
		8084483	Aug 17, 2029		U-2104	
		8283369	Nov 26, 2028		U-2104	
		8357713	Dec 22, 2029	DP	U-2104	
		8546436	Feb 29, 2032	DS		
		8546437	Apr 29, 2029		U-2104	
		9216179	Aug 01, 2031		U-2104	
		9956205	Dec 28, 2031		U-2104	
<u>ALLOPURINOL; LESINURAD - DUZALLO</u>						
N 209203	002	10183012	Nov 26, 2028		U-2104	
		8003681	Aug 25, 2025	DS		
		8084483	Aug 17, 2029		U-2104	
		8283369	Nov 26, 2028		U-2104	
		8357713	Dec 22, 2029	DP	U-2104	
		8546436	Feb 29, 2032	DS		
		8546437	Apr 29, 2029		U-2104	
		9216179	Aug 01, 2031		U-2104	
		9956205	Dec 28, 2031		U-2104	
<u>ALOGLIPTIN BENZOATE - NESINA</u>						
N 022271	001	7807689	Jun 27, 2028	DS DP	U-1337	M-300 Jul 27, 2026
		8173663	Dec 02, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8697125	Jun 16, 2029	DP		
<u>ALOGLIPTIN BENZOATE - NESINA</u>						
N 022271	002	7807689	Jun 27, 2028	DS DP	U-1337	M-300 Jul 27, 2026
		8173663	Dec 02, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8697125	Jun 16, 2029	DP		
<u>ALOGLIPTIN BENZOATE - NESINA</u>						
N 022271	003	7807689	Jun 27, 2028	DS DP	U-1337	M-300 Jul 27, 2026
		8173663	Dec 02, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8697125	Jun 16, 2029	DP		
<u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u>						
N 203414	001	7807689	Jun 27, 2028	DS DP	U-1337	M-300 Jul 27, 2026
		8173663	Mar 15, 2025		U-1338	
		8288539	Jun 24, 2025	DS		
		8900638	May 24, 2029	DP		
<u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u>						
N 203414	002	7807689	Jun 27, 2028	DS DP	U-1337	M-300 Jul 27, 2026
		8173663	Mar 15, 2025		U-1338	

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<u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u>						
N 203414	002	8288539	Jun 24, 2025	DS		
		8900638	May 24, 2029	DP		
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426	001	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Mar 15, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8637079	Jun 04, 2029	DP		
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426	002	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Mar 15, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8637079	Jun 04, 2029	DP		
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426	003	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Mar 15, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8637079	Jun 04, 2029	DP		
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426	004	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Mar 15, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8637079	Jun 04, 2029	DP		
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426	005	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Mar 15, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8637079	Jun 04, 2029	DP		
<u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u>						
N 022426	006	7807689	Jun 27, 2028	DS DP	U-1337	
		8173663	Mar 15, 2025		U-1338	
		8288539	Mar 15, 2025	DS		
		8637079	Jun 04, 2029	DP		
<u>ALPELISIB - PIORAY</u>						
N 212526	001	8227462	Apr 29, 2033	DS DP	U-2539	
		8476268	Sep 10, 2029	DS DP		NCE May 24, 2024
<u>ALPELISIB - PIORAY</u>						
N 212526	002	8227462	Apr 29, 2033	DS DP	U-2539	
		8476268	Sep 10, 2029	DS DP		NCE May 24, 2024
<u>ALPELISIB - PIORAY</u>						
N 212526	003	8227462	Apr 29, 2033	DS DP	U-2539	
		8476268	Sep 10, 2029	DS DP		NCE May 24, 2024
<u>ALPELISIB - VIJOICE</u>						
N 215039	001	8227462	Apr 29, 2033	DS DP		NCE May 24, 2024
		8476268	Sep 10, 2029	DS DP		ODE-396 Apr 05, 2029

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<u>ALPELISIB - VIJOICE</u>						
N 215039	002 8227462	Apr 29, 2033	DS DP		NCE	May 24, 2024
	8476268	Sep 10, 2029	DS DP		ODE-396	Apr 05, 2029
<u>ALPELISIB - VIJOICE</u>						
N 215039	003 8227462	Apr 29, 2033	DS DP		NCE	May 24, 2024
	8476268	Sep 10, 2029	DS DP		ODE-396	Apr 05, 2029
<u>ALVIMOPAN - ENTEREG</u>						
N 021775	001 8946262	Feb 12, 2030		U-1655		
<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	001 10154971	Dec 04, 2034		U-2459	ODE-153	Aug 24, 2024
	10646456	Jun 17, 2034		U-2808		
	11065213	Aug 23, 2038	DP			
	11077073	Aug 23, 2038		U-2106		
	11077073	Aug 23, 2038		U-2224		
	11077073	Aug 23, 2038		U-3180		
	11197835	Dec 02, 2030		U-2106		
	8389578	Jan 22, 2028		U-2105		
	8741343	Dec 02, 2030		U-2106		
	8796337	Nov 23, 2025		U-2106		
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025		U-2106		
	8895616	Nov 23, 2025		U-2106		
	8895617	Nov 23, 2025		U-2106		
	8895618	Nov 23, 2025	DP			
	9867791	Dec 02, 2030		U-2106		
	9867792	Dec 02, 2030		U-2106		
	9867793	Dec 02, 2030		U-2106		
	9877933	Dec 02, 2030		U-2224		
<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	002 10154971	Dec 04, 2034		U-2459	ODE-153	Aug 24, 2024
	10646456	Jun 17, 2034		U-2808		
	11065213	Aug 23, 2038	DP			
	11077073	Aug 23, 2038		U-2106		
	11077073	Aug 23, 2038		U-2224		
	11077073	Aug 23, 2038		U-3180		
	11197835	Dec 02, 2030		U-2106		
	8389578	Jan 22, 2028		U-2105		
	8741343	Dec 02, 2030		U-2106		
	8796337	Nov 23, 2025		U-2106		
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025		U-2106		
	8895616	Nov 23, 2025		U-2106		
	8895617	Nov 23, 2025		U-2106		
	8895618	Nov 23, 2025	DP			
	9867791	Dec 02, 2030		U-2106		
	9867792	Dec 02, 2030		U-2106		
	9867793	Dec 02, 2030		U-2106		

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<u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u>						
N 208944	002 9877933	Dec 02, 2030	U-2224			
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	001 10213393	Feb 15, 2038	U-20			
	10213394	Feb 15, 2038	U-2497			
	10500170	Feb 15, 2038	U-20			
	10500171	Feb 15, 2038	U-2497			
	10500172	Feb 15, 2038	U-2497			
	10512617	Feb 15, 2038	U-2497			
	8252331	Mar 13, 2030	DP			
	8389578	Jan 22, 2028	U-219			
	8389578	Jan 22, 2028	U-3054			
	8574626	Nov 28, 2025	DP U-20			
	8796337	Nov 23, 2025	U-219			
	8796337	Nov 23, 2025	U-2497			
	8796337	Nov 23, 2025	U-3054			
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025	U-219			
	8895615	Nov 23, 2025	U-3054			
	8895616	Nov 23, 2025	U-219			
	8895616	Nov 23, 2025	U-3054			
	8895617	Nov 23, 2025	U-219			
	8895617	Nov 23, 2025	U-3054			
	8895618	Nov 23, 2025	DP			
	8987333	Nov 23, 2025	DP			
	9072697	Nov 23, 2025	U-219			
	9072697	Nov 23, 2025	U-3054			
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	002 10213393	Feb 15, 2038	U-20			
	10213394	Feb 15, 2038	U-2497			
	10500170	Feb 15, 2038	U-20			
	10500171	Feb 15, 2038	U-2497			
	10500172	Feb 15, 2038	U-2497			
	10512617	Feb 15, 2038	U-2497			
	8252331	Mar 13, 2030	DP			
	8389578	Jan 22, 2028	U-219			
	8389578	Jan 22, 2028	U-3054			
	8574626	Nov 28, 2025	DP U-20			
	8796337	Nov 23, 2025	U-219			
	8796337	Nov 23, 2025	U-2497			
	8796337	Nov 23, 2025	U-3054			
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025	U-219			
	8895615	Nov 23, 2025	U-3054			
	8895616	Nov 23, 2025	U-219			
	8895616	Nov 23, 2025	U-3054			
	8895617	Nov 23, 2025	U-219			
	8895617	Nov 23, 2025	U-3054			

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<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	002	8895618	Nov 23, 2025	DP		
		8987333	Nov 23, 2025	DP		
		9072697	Nov 23, 2025		U-219	
		9072697	Nov 23, 2025		U-3054	
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	003	10213393	Feb 15, 2038		U-20	
		10213394	Feb 15, 2038		U-2497	
		10500170	Feb 15, 2038		U-20	
		10500171	Feb 15, 2038		U-2497	
		10500172	Feb 15, 2038		U-2497	
		10512617	Feb 15, 2038		U-2497	
		8252331	Mar 13, 2030	DP		
		8389578	Jan 22, 2028		U-219	
		8389578	Jan 22, 2028		U-3054	
		8574626	Nov 28, 2025	DP	U-20	
		8796337	Nov 23, 2025		U-219	
		8796337	Nov 23, 2025		U-2497	
		8796337	Nov 23, 2025		U-3054	
		8889740	Nov 23, 2025	DP		
		8895614	Nov 23, 2025	DP		
		8895615	Nov 23, 2025		U-219	
		8895615	Nov 23, 2025		U-3054	
		8895616	Nov 23, 2025		U-219	
		8895616	Nov 23, 2025		U-3054	
		8895617	Nov 23, 2025		U-219	
		8895617	Nov 23, 2025		U-3054	
		8895618	Nov 23, 2025	DP		
		8987333	Nov 23, 2025	DP		
		9072697	Nov 23, 2025		U-219	
		9072697	Nov 23, 2025		U-3054	
<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	004	10213393	Feb 15, 2038		U-20	
		10213394	Feb 15, 2038		U-2497	
		10500170	Feb 15, 2038		U-20	
		10500171	Feb 15, 2038		U-2497	
		10500172	Feb 15, 2038		U-2497	
		10512617	Feb 15, 2038		U-2497	
		8252331	Mar 13, 2030	DP		
		8389578	Jan 22, 2028		U-219	
		8389578	Jan 22, 2028		U-3054	
		8574626	Nov 28, 2025	DP	U-20	
		8796337	Nov 23, 2025		U-219	
		8796337	Nov 23, 2025		U-2497	
		8796337	Nov 23, 2025		U-3054	
		8889740	Nov 23, 2025	DP		
		8895614	Nov 23, 2025	DP		
		8895615	Nov 23, 2025		U-219	
		8895615	Nov 23, 2025		U-3054	
		8895616	Nov 23, 2025		U-219	

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<u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u>						
N 209410	004	8895616	Nov 23, 2025	U-3054		
		8895617	Nov 23, 2025	U-219		
		8895617	Nov 23, 2025	U-3054		
		8895618	Nov 23, 2025	DP		
		8987333	Nov 23, 2025	DP		
		9072697	Nov 23, 2025	U-219		
		9072697	Nov 23, 2025	U-3054		
<u>AMBRISENTAN - LETAIRIS</u>						
N 022081	001	8377933	Dec 11, 2027	U-1754		
		9474752	Dec 11, 2027	U-1754		
		9549926	Oct 14, 2031	U-1965		
<u>AMBRISENTAN - LETAIRIS</u>						
N 022081	002	8377933	Dec 11, 2027	U-1754		
		9474752	Dec 11, 2027	U-1754		
		9549926	Oct 14, 2031	U-1965		
<u>AMIFAMPRIDINE PHOSPHATE - FIRDAPSE</u>						
N 208078	001	10626088	Feb 25, 2037	DP	NPP	Sep 29, 2025
		10793893	May 26, 2034	U-2956	ODE-223	Nov 28, 2025
		11060128	Jun 29, 2032	U-2956		
		11268128	Jun 29, 2032	U-2956		
		11274331	Jun 29, 2032	U-2956		
		11274332	Jun 29, 2032	U-2956		
<u>AMIKACIN SULFATE - ARIKAYCE KIT</u>						
N 207356	001	10251900	May 15, 2035	U-2414	ODE-214	Sep 28, 2025
		10751355	May 15, 2035	U-2414	GAIN	Sep 28, 2030
		11446318	May 15, 2035	U-2414		
		7718189	Jun 06, 2025	DP U-2415		
		8226975	Aug 15, 2028	DP		
		8632804	Dec 05, 2026	U-2416		
		8642075	Dec 05, 2026	DP		
		8679532	Dec 05, 2026	U-2415		
		8802137	Apr 08, 2024	DP U-2414		
		9566234	Jan 18, 2034	DP U-2415		
		9827317	Apr 08, 2024	DP U-2415		
		9895385	May 15, 2035	U-2417		
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - LEVULAN</u>						
N 020965	001	10357567	Jan 12, 2038	U-3163		
		11077192	Jan 12, 2038	U-3163		
		11135293	Jan 12, 2038	U-3163		
		11571478	Jan 12, 2038	U-3163		
		11690914	Jan 12, 2038	U-3163		
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - AMELUZ</u>						
N 208081	001	11235169	Oct 15, 2040	U-3303		
		11540981	Feb 07, 2028	DP		
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - GLEOLAN</u>						
N 208630	001				ODE-146	Jun 06, 2024

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<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325	001 7635773	Mar 13, 2029	DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325	002 7635773	Mar 13, 2029	DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
N 022325	003 7635773	Mar 13, 2029	DP			
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510	001 10525033	Mar 10, 2031	DP		NCE	Feb 26, 2025
	11357753	Feb 09, 2038	U-2754			
	9084765	Feb 26, 2034	U-1744			
	9084765	Feb 26, 2034	U-2754			
	9084765	Feb 26, 2034	U-3467			
	9545426	Mar 10, 2031	U-1744			
	9545426	Mar 10, 2031	U-2754			
	9889118	Mar 10, 2031	U-1744			
	9889118	Mar 10, 2031	U-2754			
<u>AMISULPRIDE - BARHEMSYS</u>						
N 209510	002 10525033	Mar 10, 2031	DP		NCE	Feb 26, 2025
	11357753	Feb 09, 2038	U-2754			
	9084765	Feb 26, 2034	U-1744			
	9084765	Feb 26, 2034	U-2754			
	9084765	Feb 26, 2034	U-3467			
	9545426	Mar 10, 2031	U-1744			
	9545426	Mar 10, 2031	U-2754			
	9889118	Mar 10, 2031	U-1744			
	9889118	Mar 10, 2031	U-2754			
<u>AMLODIPINE BENZOATE - KATERZIA</u>						
N 211340	001 10695329	Oct 16, 2037	DP			
	10799453	Apr 11, 2039	DP			
	10894039	Oct 06, 2037	U-185			
	10894039	Oct 06, 2037	U-3			
	10952998	Oct 06, 2037	DP			
	10959991	Oct 06, 2037	U-158			
	10959991	Oct 06, 2037	U-39			
	11364230	Oct 06, 2037	DP			
	11471409	Oct 06, 2037	U-3447			
	11471409	Oct 06, 2037	U-3448			
	11484498	Oct 06, 2037	DP			
	11701326	Oct 06, 2037	DP			
<u>AMLODIPINE BESYLATE - NORLIOVA</u>						
N 214439	001 11253474	Feb 24, 2041	DP	U-3309		
	11253474	Feb 24, 2041	DP	U-3310		
	11253474	Feb 24, 2041	DP	U-3311		
	11458095	Feb 24, 2041	DP	U-3309		
	11458095	Feb 24, 2041	DP	U-3310		
	11458095	Feb 24, 2041	DP	U-3311		
	11723866	Feb 24, 2041	DP	U-3309		
	11723866	Feb 24, 2041	DP	U-3310		

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<u>AMLODIPINE BESYLATE - NORLIOVA</u>						
N 214439	001 11723866	Feb 24, 2041	DP U-3311			
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	001 10350171	Jun 14, 2038	DP			
	10925835	Jun 14, 2038	U-2410			
	10945960	Jun 14, 2038	DP			
	9408837	Feb 28, 2030	U-2410			
	9662315	May 22, 2029	DP U-2410			
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	002 10350171	Jun 14, 2038	DP			
	10925835	Jun 14, 2038	U-2410			
	10945960	Jun 14, 2038	DP			
	9408837	Feb 28, 2030	U-2410			
	9662315	May 22, 2029	DP U-2410			
<u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u>						
N 210045	003 10350171	Jun 14, 2038	DP			
	10925835	Jun 14, 2038	U-2410			
	10945960	Jun 14, 2038	DP			
	9408837	Feb 28, 2030	U-2410			
	9662315	May 22, 2029	DP U-2410			
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	001 7846961	Oct 05, 2029	DS DP U-3			
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	002 7846961	Oct 05, 2029	DS DP U-3			
<u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u>						
N 205003	003 7846961	Oct 05, 2029	DS DP U-3			
<u>AMOXICILLIN - MOXATAG</u>						
N 050813	001 8299052	May 07, 2027	U-1304			
	8357394	Dec 08, 2026	DP			
	8778924	Dec 08, 2026	DS DP U-897			
<u>AMOXICILLIN; CLARITHROMYCIN; VONOPRAZAN FUMARATE - VOQUEZNA TRIPLE PAK</u>						
N 215152	001 7977488	Aug 11, 2028	DS		NCE	May 03, 2027
	9186411	Aug 11, 2030	DP		GAIN	May 03, 2032
<u>AMOXICILLIN; OMEPRAZOLE MAGNESIUM; RIFABUTIN - TALICIA</u>						
N 213004	001 10238606	Feb 12, 2034	DP		NP	Nov 01, 2022
	11135172	Feb 12, 2034	DP U-2660		GAIN	Nov 01, 2027
	9050263	Feb 12, 2034	DP U-2660			
	9498445	Feb 12, 2034	DP U-2660			
	9603806	Feb 12, 2034	DP U-2660			
<u>AMOXICILLIN; VONOPRAZAN FUMARATE - VOQUEZNA DUAL PAK</u>						
N 215153	001 7977488	Aug 11, 2028	DS		NCE	May 03, 2027
	9186411	Aug 11, 2030	DP		GAIN	May 03, 2032
<u>AMPHETAMINE - ADZENYS ER</u>						
N 204325	001 8709491	Jun 28, 2032	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			

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<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326 001	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326 002	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326 003	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326 004	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326 005	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<u>AMPHETAMINE - ADZENYS XR-ODT</u>						
N 204326 006	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 001	8846100	Aug 24, 2029	DP			
	9173857	May 12, 2026		U-2025		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 002	8846100	Aug 24, 2029	DP			
	9173857	May 12, 2026		U-2025		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 003	8846100	Aug 24, 2029	DP			
	9173857	May 12, 2026		U-2025		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 004	8846100	Aug 24, 2029	DP			
	9173857	May 12, 2026		U-2025		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905 001	10130580	Apr 19, 2024	DP			

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<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	001	10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	002	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	003	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	004	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE SULFATE - EVEKEO ODT</u>						
N 209905	005	10130580	Apr 19, 2024	DP		
		10441554	Mar 10, 2037	DP		
		11160772	Mar 10, 2037	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR</u>						
N 208147	001	10086087	Mar 15, 2027	DP		
		11590228	Sep 07, 2036	DP U-3538		
		8062667	Mar 29, 2029	DP		
		8597684	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		8883217	Mar 15, 2027	DP		
		9675703	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 5</u>						
N 210526	001	11590081	Sep 24, 2038	DP U-3538		
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 10</u>						
N 210526	002	11590081	Sep 24, 2038	DP U-3538		
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 15</u>						
N 210526	003	11590081	Sep 24, 2038	DP U-3538		
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		
<u>AMPHETAMINE; AMPHETAMINE ASPARTATE/DEXTROAMPHETAMINE SULFATE - DYANA VEL XR 20</u>						
N 210526	004	11590081	Sep 24, 2038	DP U-3538		
		8337890	Mar 15, 2027	DP		
		8747902	Mar 15, 2027	DP		
		9675704	Mar 15, 2027	DP		

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<u>AMPICILLIN/AMPICILLIN TRIHYDRATE - AMPICILLIN TRIHYDRATE</u>						
A 216554	001				CGT	Jun 03, 2024
<u>AMPICILLIN/AMPICILLIN TRIHYDRATE - AMPICILLIN TRIHYDRATE</u>						
A 216554	002				CGT	Jun 03, 2024
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	001	10028995	Dec 18, 2034	U-2338		
		10335451	Dec 16, 2029	U-2581		
		10493124	Dec 18, 2034	U-2679		
		10500247	Dec 16, 2029	U-2680		
		10500247	Dec 16, 2029	U-2681		
		10548943	Dec 16, 2029	U-2739		
		10548943	Dec 16, 2029	U-2740		
		11096983	Dec 18, 2034	U-3211		
		11096983	Dec 18, 2034	U-3212		
		11219662	Jan 06, 2037	U-3262		
		11559559	Dec 18, 2034	U-3514		
		9220745	Dec 18, 2034	U-2217		
		9220745	Dec 18, 2034	U-2218		
		9572856	Jul 18, 2031	U-2221		
		9867863	Dec 16, 2029	U-2231		
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	002	10028995	Dec 18, 2034	U-2338		
		10335451	Dec 16, 2029	U-2581		
		10493124	Dec 18, 2034	U-2679		
		10500247	Dec 16, 2029	U-2680		
		10500247	Dec 16, 2029	U-2681		
		10548943	Dec 16, 2029	U-2739		
		10548943	Dec 16, 2029	U-2740		
		11096983	Dec 18, 2034	U-3211		
		11096983	Dec 18, 2034	U-3212		
		11219662	Jan 06, 2037	U-3262		
		11559559	Dec 18, 2034	U-3514		
		9220745	Dec 18, 2034	U-2217		
		9220745	Dec 18, 2034	U-2218		
		9572856	Nov 20, 2030	U-2221		
		9867863	Dec 16, 2029	U-2231		
<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360	003	10028995	Dec 18, 2034	U-2338		
		10335451	Dec 16, 2029	U-2581		
		10493124	Dec 18, 2034	U-2679		
		10500247	Dec 16, 2029	U-2680		
		10500247	Dec 16, 2029	U-2681		
		10548943	Dec 16, 2029	U-2739		
		10548943	Dec 16, 2029	U-2740		
		11096983	Dec 18, 2034	U-3211		
		11096983	Dec 18, 2034	U-3212		
		11219662	Jan 06, 2037	U-3262		
		11559559	Dec 18, 2034	U-3514		
		9220745	Dec 18, 2034	U-2217		

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<u>ANGIOTENSIN II ACETATE - GIAPREZA</u>						
N 209360 003	9220745	Dec 18, 2034	U-2218			
	9572856	Jul 18, 2031	U-2221			
	9867863	Dec 16, 2029	U-2231			
<u>APALUTAMIDE - ERLEADA</u>						
N 210951 001	10052314	Sep 23, 2033	U-2381		Y	
	10052314	Sep 23, 2033	U-2382		Y	
	10702508	Apr 30, 2038	U-3012			
	10849888	Sep 23, 2033	U-3013			
	8445507	Sep 15, 2030	DS DP U-2237			
	8445507	Sep 15, 2030	DS DP U-2624			
	8802689	Mar 27, 2027	U-2237			
	8802689	Mar 27, 2027	U-2624			
	9388159	Mar 27, 2027	DS DP			
	9481663	Jun 04, 2033	DS DP U-2237			
	9481663	Jun 04, 2033	DS DP U-2624			
	9884054	Sep 23, 2033	U-2237			
	9987261	Mar 27, 2027	DP			
	RE49353	Sep 23, 2033	U-2381			
<u>APALUTAMIDE - ERLEADA</u>						
N 210951 002	10702508	Apr 30, 2038	U-3012			
	10849888	Sep 23, 2033	U-3013			
	8445507	Sep 15, 2030	DS DP U-2237			
	8445507	Sep 15, 2030	DS DP U-2624			
	8802689	Mar 27, 2027	U-2237			
	8802689	Mar 27, 2027	U-2624			
	9388159	Mar 27, 2027	DS DP			
	9481663	Jun 04, 2033	DS DP U-2237			
	9481663	Jun 04, 2033	DS DP U-2624			
	9884054	Sep 23, 2033	U-2237			
	9987261	Mar 27, 2027	DP			
	RE49353	Sep 23, 2033	U-2381			
<u>APIXABAN - ELIQUIS</u>						
N 202155 001	6967208	Nov 21, 2026	DS DP U-1167			
	6967208	Nov 21, 2026	DS DP U-1200			
	6967208	Nov 21, 2026	DS DP U-1301			
	6967208	Nov 21, 2026	DS DP U-1302			
	6967208	Nov 21, 2026	DS DP U-1323			
	6967208	Nov 21, 2026	DS DP U-1501			
	6967208	Nov 21, 2026	DS DP U-1502			
	6967208	Nov 21, 2026	DS DP U-1729			
	6967208	Nov 21, 2026	DS DP U-1730			
	9326945	Feb 24, 2031	DP			
<u>APIXABAN - ELIQUIS</u>						
N 202155 002	6967208	Nov 21, 2026	DS DP U-1200			
	6967208	Nov 21, 2026	DS DP U-1301			
	6967208	Nov 21, 2026	DS DP U-1302			
	6967208	Nov 21, 2026	DS DP U-1323			
	9326945	Feb 24, 2031	DP			

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<u>APIXABAN - ELIQUIS</u>						
N 202155	002	6967208	Nov 21, 2026	DS DP U-1200		
		6967208	Nov 21, 2026	DS DP U-1301		
		6967208	Nov 21, 2026	DS DP U-1302		
		6967208	Nov 21, 2026	DS DP U-1323		
		9326945	Feb 24, 2031	DP		
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	001	10420763	Jun 11, 2030	DP U-2825		
		10449146	Apr 19, 2036	U-2825		
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036	U-2825		
		11419769	Dec 16, 2031	DP U-2825		
		8414922	Dec 16, 2031	DP U-2825		
		8603514	Apr 03, 2024	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	002	10420763	Jun 11, 2030	DP U-2825		
		10449146	Apr 19, 2036	U-2825		
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036	U-2825		
		11419769	Dec 16, 2031	DP U-2825		
		8414922	Dec 16, 2031	DP U-2825		
		8603514	Apr 03, 2024	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		
		9669019	Jun 11, 2030	DP U-2825		
		9669021	Jun 11, 2030	U-2825		
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	003	10420763	Jun 11, 2030	DP U-2825		
		10449146	Apr 19, 2036	U-2825		
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036	U-2825		
		11419769	Dec 16, 2031	DP U-2825		
		8414922	Dec 16, 2031	DP U-2825		
		8603514	Apr 03, 2024	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP U-2825		
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP U-2825		
		9326981	Jun 11, 2030	U-2825		

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<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	003	9669019	Jun 11, 2030	DP	U-2825	
		9669021	Jun 11, 2030		U-2825	
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	004	10420763	Jun 11, 2030	DP	U-2825	
		10449146	Apr 19, 2036		U-2825	
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036		U-2825	
		11419769	Dec 16, 2031	DP	U-2825	
		8414922	Dec 16, 2031	DP	U-2825	
		8603514	Apr 03, 2024	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP	U-2825	
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP	U-2825	
		9326981	Jun 11, 2030		U-2825	
		9669019	Jun 11, 2030	DP	U-2825	
		9669021	Jun 11, 2030		U-2825	
<u>APOMORPHINE HYDROCHLORIDE - KYNMOBI</u>						
N 210875	005	10420763	Jun 11, 2030	DP	U-2825	
		10449146	Apr 19, 2036		U-2825	
		10821074	Aug 07, 2029	DP		
		10959943	Apr 19, 2036		U-2825	
		11419769	Dec 16, 2031	DP	U-2825	
		8414922	Dec 16, 2031	DP	U-2825	
		8603514	Apr 03, 2024	DP		
		8765167	Feb 20, 2024	DP		
		8846074	Dec 16, 2031	DP	U-2825	
		9044475	Jun 11, 2030	DP		
		9283219	Jun 11, 2030	DP	U-2825	
		9326981	Jun 11, 2030		U-2825	
		9669019	Jun 11, 2030	DP	U-2825	
		9669021	Jun 11, 2030		U-2825	
<u>APREMILAST - OTEZLA</u>						
N 205437	001	10092541	May 29, 2034	U-2403	I-884	Dec 20, 2024
		10092541	May 29, 2034	U-2659	M-299	Jul 20, 2026
		7427638	Feb 16, 2028	DS DP	ODE-248	Jul 19, 2026
		9872854	May 29, 2034	U-2232		
		9872854	May 29, 2034	U-2233		
<u>APREMILAST - OTEZLA</u>						
N 205437	002	10092541	May 29, 2034	U-2403	I-884	Dec 20, 2024
		10092541	May 29, 2034	U-2659	M-299	Jul 20, 2026
		7427638	Feb 16, 2028	DS DP	ODE-248	Jul 19, 2026
		9872854	May 29, 2034	U-2232		
		9872854	May 29, 2034	U-2233		
<u>APREMILAST - OTEZLA</u>						
N 205437	003	10092541	May 29, 2034	U-2403	I-884	Dec 20, 2024
		10092541	May 29, 2034	U-2659	M-299	Jul 20, 2026
		7427638	Feb 16, 2028	DS DP	ODE-248	Jul 19, 2026

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<u>APREMILAST - OTEZLA</u>						
N 205437	003 9872854	May 29, 2034	U-2232			
	9872854	May 29, 2034	U-2233			
<u>APREPITANT - EMEND</u>						
N 021549	001 8258132	Sep 26, 2027	DP U-1743			
	8258132	Sep 26, 2027	DP U-901			
<u>APREPITANT - EMEND</u>						
N 021549	002 8258132	Sep 26, 2027	DP U-1743			
	8258132	Sep 26, 2027	DP U-901			
<u>APREPITANT - EMEND</u>						
N 021549	003 8258132	Sep 26, 2027	DP U-1743			
	8258132	Sep 26, 2027	DP U-901			
<u>APREPITANT - EMEND</u>						
N 207865	001 8258132	Sep 26, 2027	DP U-1916			
<u>APREPITANT - CINVANTI</u>						
N 209296	001 10500208	Sep 18, 2035	DP			
	10624850	Sep 18, 2035	U-2161			
	10953018	Sep 18, 2035	U-2161			
	11173118	Sep 18, 2035	DP			
	11744800	Sep 18, 2035	DP			
	9561229	Sep 18, 2035	DP U-2161			
	9808465	Sep 18, 2035	U-2161			
	9974742	Sep 18, 2035	DP			
	9974793	Sep 18, 2035	DP			
	9974794	Sep 18, 2035	DP U-2161			
<u>APREPITANT - APONVIE</u>						
N 216457	001 10500208	Sep 18, 2035	DP			
	10624850	Sep 18, 2035	U-3440			
	10953018	Sep 18, 2035	U-3440			
	11173118	Sep 18, 2035	DP			
	11744800	Sep 18, 2035	DP U-3690			
	9561229	Sep 18, 2035	DP U-3440			
	9808465	Sep 18, 2035	U-3440			
	9974742	Sep 18, 2035	DP			
	9974793	Sep 18, 2035	DP			
	9974794	Sep 18, 2035	DP U-3440			
<u>ARGATROBAN - ARGATROBAN IN SODIUM CHLORIDE</u>						
N 022434	001 7589106	Sep 26, 2027	DP U-1163			
	7687516	Sep 26, 2027	DP U-1164			
<u>ARIPIPIRAZOLE - ABILIFY</u>						
N 021436	001 8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIPIRAZOLE - ABILIFY</u>						
N 021436	002 8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 002	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 003	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 004	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 005	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021436 006	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021713 001	8759350	Mar 02, 2027	U-1529			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729 002	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729 003	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
	8759350	Mar 02, 2027	U-1529			
	9125939	Jul 28, 2026	U-1749			
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729 004	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021729 005	8017615	Jun 16, 2024	DP			
	8017615*PED	Dec 16, 2024				
<u>ARIPIRAZOLE - ABILIFY</u>						
N 021866 001	7115587	Jul 21, 2024	DP U-764			
	7115587*PED	Jan 21, 2025				
	7550445	Jul 21, 2024	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 001	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	10980803	Sep 24, 2033	U-1632			
	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			
	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	7807680	Oct 19, 2024	DP			
	8030313	Oct 19, 2024	U-1632			
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 002	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	10980803	Sep 24, 2033	U-1632			
	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			
	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	7807680	Oct 19, 2024	DP			
	8030313	Oct 19, 2024	U-1632			
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 003	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 003	10980803	Sep 24, 2033	U-1632			
	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			
	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	7807680	Oct 19, 2024	DP			
	8030313	Oct 19, 2024	U-1632			
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
<u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u>						
N 202971 004	10525057	Mar 08, 2034	U-1632			
	10525057	Mar 08, 2034	U-2723			
	10525057	Mar 08, 2034	U-543			
	10980803	Sep 24, 2033	U-1632			
	10980803	Sep 24, 2033	U-543			
	11154553	Sep 24, 2033	U-1632			
	11154553	Sep 24, 2033	U-3245			
	11154553	Sep 24, 2033	U-814			
	11344547	Sep 24, 2033	U-1632			
	11344547	Sep 24, 2033	U-3245			
	11344547	Sep 24, 2033	U-814			
	11400087	Sep 24, 2033	U-1632			
	11400087	Sep 24, 2033	U-3245			
	11400087	Sep 24, 2033	U-814			
	11648347	Apr 06, 2034	DP			
	7807680	Oct 19, 2024	DP			
	8030313	Oct 19, 2024	U-1632			
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 001	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 001	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8674825	Apr 09, 2029	DP	U-2170		
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027		U-1529		
	8847766	Mar 29, 2030	DP	U-2167		
	8945005	Aug 19, 2029	DP	U-2167		
	8956288	Jul 06, 2029	DP	U-2167		
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026		U-1749		
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP	U-2168		
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 002	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8674825	Apr 09, 2029	DP	U-2170		
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027		U-1529		
	8847766	Mar 29, 2030	DP	U-2167		
	8945005	Aug 19, 2029	DP	U-2167		
	8956288	Jul 06, 2029	DP	U-2167		
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026		U-1749		
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP	U-2168		
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	002	9444503	Nov 19, 2027	DP	U-2169	
		9941931	Nov 04, 2030	DP		
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	003	10441194	Jul 26, 2029	DP		
		10517507	Jun 13, 2032	DP		
		11229378	Jul 11, 2031	DP		
		11464423	Sep 15, 2030	DP		
		11476952	Apr 28, 2026	DP		
		7978064	Sep 14, 2026	DP		
		8017615	Jun 16, 2024	DP		
		8114021	Jun 21, 2030	DP		
		8258962	Nov 25, 2030	DP		
		8545402	Apr 27, 2030	DP		
		8547248	Dec 18, 2030	DP	U-2167	
		8674825	Apr 09, 2029	DP	U-2170	
		8718193	Dec 05, 2029	DP		
		8759350	Mar 02, 2027		U-1529	
		8847766	Mar 29, 2030	DP	U-2167	
		8945005	Aug 19, 2029	DP	U-2167	
		8956288	Jul 06, 2029	DP	U-2167	
		8961412	Nov 17, 2030	DP		
		9060708	Mar 05, 2029	DP		
		9119554	Dec 16, 2028	DP		
		9125939	Jul 28, 2026		U-1749	
		9149577	Dec 15, 2029	DP		
		9258035	Mar 05, 2029	DP		
		9268909	Oct 15, 2033	DP	U-2168	
		9320455	Dec 15, 2031	DP		
		9433371	Sep 15, 2029	DP		
		9444503	Nov 19, 2027	DP	U-2169	
		9941931	Nov 04, 2030	DP		
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202	004	10441194	Jul 26, 2029	DP		
		10517507	Jun 13, 2032	DP		
		11229378	Jul 11, 2031	DP		
		11464423	Sep 15, 2030	DP		
		11476952	Apr 28, 2026	DP		
		7978064	Sep 14, 2026	DP		
		8017615	Jun 16, 2024	DP		
		8114021	Jun 21, 2030	DP		
		8258962	Nov 25, 2030	DP		
		8545402	Apr 27, 2030	DP		
		8547248	Dec 18, 2030	DP	U-2167	
		8674825	Apr 09, 2029	DP	U-2170	
		8718193	Dec 05, 2029	DP		
		8759350	Mar 02, 2027		U-1529	
		8847766	Mar 29, 2030	DP	U-2167	
		8945005	Aug 19, 2029	DP	U-2167	
		8956288	Jul 06, 2029	DP	U-2167	

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<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 004	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026		U-1749		
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP	U-2168		
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 005	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP	U-2167		
	8674825	Apr 09, 2029	DP	U-2170		
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027		U-1529		
	8847766	Mar 29, 2030	DP	U-2167		
	8945005	Aug 19, 2029	DP	U-2167		
	8956288	Jul 06, 2029	DP	U-2167		
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026		U-1749		
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP	U-2168		
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP	U-2169		
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 006	10441194	Jul 26, 2029	DP			
	10517507	Jun 13, 2032	DP			
	11229378	Jul 11, 2031	DP			
	11464423	Sep 15, 2030	DP			
	11476952	Apr 28, 2026	DP			
	7978064	Sep 14, 2026	DP			
	8017615	Jun 16, 2024	DP			
	8114021	Jun 21, 2030	DP			

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<u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u>						
N 207202 006	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9320455	Dec 15, 2031	DP			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9941931	Nov 04, 2030	DP			
<u>ARIPIRAZOLE - ABILIFY ASIMTUFII</u>						
N 217006 001	10517951	Apr 23, 2033	DP U-3245			
	10517951	Apr 23, 2033	DP U-814			
	11097007	Apr 23, 2033	DP U-3245			
	11097007	Apr 23, 2033	DP U-814			
	11638757	Apr 23, 2033	DP U-3245			
	11638757	Apr 23, 2033	DP U-814			
	8338427	Mar 15, 2025	DP U-1530			
	8399469	Jun 29, 2025	DS			
<u>ARIPIRAZOLE - ABILIFY ASIMTUFII</u>						
N 217006 002	10517951	Apr 23, 2033	DP U-3245			
	10517951	Apr 23, 2033	DP U-814			
	11097007	Apr 23, 2033	DP U-3245			
	11097007	Apr 23, 2033	DP U-814			
	11638757	Apr 23, 2033	DP U-3245			
	11638757	Apr 23, 2033	DP U-814			
	8338427	Mar 15, 2025	DP U-1530			
	8399469	Jun 29, 2025	DS			
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533 001	10112903	Jun 24, 2030	DS U-543			
	10226458	Mar 19, 2032	U-543			
	10238651	Mar 19, 2035	U-2402			
	10813928	Mar 19, 2035	U-2983			
	11097006	Oct 24, 2033	DP U-764			
	11273158	Apr 06, 2039	U-543			
	11406632	Mar 19, 2035	U-2402			
	8431576	Oct 26, 2030	DS			
	8796276	Jun 24, 2030	U-543			

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<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533	001	9034867	Nov 07, 2032	DP	U-543	
		9193685	Oct 24, 2033	DP	U-543	
		9452131	Mar 19, 2035		U-2402	
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533	002	10112903	Jun 24, 2030	DS	U-543	
		10226458	Mar 19, 2032		U-543	
		10238651	Mar 19, 2035		U-2402	
		10813928	Mar 19, 2035		U-2983	
		11097006	Oct 24, 2033	DP	U-764	
		11273158	Apr 06, 2039		U-543	
		11406632	Mar 19, 2035		U-2402	
		8431576	Oct 26, 2030	DS		
		8796276	Jun 24, 2030		U-543	
		9034867	Nov 07, 2032	DP	U-543	
		9193685	Oct 24, 2033	DP	U-543	
		9452131	Mar 19, 2035		U-2402	
		9526726	Mar 19, 2035	DP		
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533	003	10112903	Jun 24, 2030	DS	U-543	
		10226458	Mar 19, 2032		U-543	
		10238651	Mar 19, 2035		U-2402	
		10813928	Mar 19, 2035		U-2402	
		11097006	Oct 24, 2033	DP	U-764	
		11273158	Apr 06, 2039		U-543	
		11406632	Mar 19, 2035		U-2402	
		8431576	Oct 26, 2030	DS		
		8796276	Jun 24, 2030		U-543	
		9034867	Nov 07, 2032	DP	U-543	
		9193685	Oct 24, 2033	DP	U-543	
		9452131	Mar 19, 2035		U-2402	
		9526726	Mar 19, 2035	DP		
<u>ARIPIRAZOLE LAUROXIL - ARISTADA</u>						
N 207533	004	10112903	Jun 24, 2030	DS	U-543	
		10226458	Mar 19, 2032		U-543	
		10238651	Mar 19, 2035		U-2402	
		10813928	Mar 19, 2035		U-2983	
		11097006	Oct 24, 2033	DP	U-764	
		11273158	Apr 06, 2039		U-543	
		11406632	Mar 19, 2035		U-2402	
		8431576	Oct 26, 2030	DS		
		8796276	Jun 24, 2030		U-543	
		9034867	Nov 07, 2032	DP	U-543	
		9193685	Oct 24, 2033	DP	U-543	
		9452131	Mar 19, 2035		U-2402	
<u>ARIPIRAZOLE LAUROXIL - ARISTADA INITIO KIT</u>						
N 209830	001	10016415	Sep 08, 2035	DP		
		10112903	Jun 24, 2030	DS	U-543	
		10688091	Aug 17, 2035	DP		

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<u>ARIPIRAZOLE LAUROXIL - ARISTADA INITIO KIT</u>						
N 209830	001	10849894	Aug 17, 2035	U-543		
		11154552	Aug 17, 2035	DP		
		11273158	Apr 06, 2039	U-543		
		8431576	Oct 26, 2030	DS		
		8796276	Jun 24, 2030	U-543		
<u>ARSENIC TRIOXIDE - TRISENOX</u>						
N 021248	001				ODE-167	Jan 12, 2025
<u>ARSENIC TRIOXIDE - TRISENOX</u>						
N 021248	002				ODE-167	Jan 12, 2025
<u>ARTESUNATE - ARTESUNATE</u>						
N 213036	001				NCE ODE-290	May 26, 2025 May 26, 2027
<u>ASCIMINIB HYDROCHLORIDE - SCEMBLIX</u>						
N 215358	001	11407735	May 14, 2040	DS	NCE	Oct 29, 2026
		8829195	May 13, 2033	DS	U-1374 ODE-381 ODE-382	Oct 29, 2028 Oct 29, 2028
<u>ASCIMINIB HYDROCHLORIDE - SCEMBLIX</u>						
N 215358	002	11407735	May 14, 2040	DS	NCE	Oct 29, 2026
		8829195	May 13, 2033	DS	U-1374 ODE-381 ODE-382	Oct 29, 2028 Oct 29, 2028
<u>ASCORBIC ACID - ASCOR</u>						
N 209112	001				ODE-160	Oct 02, 2024
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u>						
N 021881	001	7169381	Sep 01, 2024	DS DP		
		7658914	Sep 01, 2024	DS DP		
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - PLENVU</u>						
N 209381	001	10016504	Sep 10, 2033	DP		
		10646512	Mar 25, 2032	DP		
		10780112	Mar 09, 2032	DP		
		10792306	Mar 09, 2032	DP	U-2310	
		10918723	Sep 10, 2033		U-2310	
		11529368	Mar 09, 2032	DP	U-2310	
		8999313	Sep 10, 2033	DP		
		9326969	Sep 10, 2033		U-2310	
		9592252	Aug 11, 2032	DP	U-2310	
		9707297	Sep 10, 2033	DP		
<u>ASENAPINE - SECUADO</u>						
N 212268	001	10022445	Jul 25, 2033	DP		
		10583121	Jul 25, 2033	DP	U-2763	
		10814002	Jul 25, 2033	DP	U-2763	
		11123305	Jul 25, 2033	DP		
		11813364	Sep 22, 2033	DP		
		9687474	Jul 25, 2033	DP		

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<u>ASENAPINE - SECUADO</u>						
N 212268 002	10022445	Jul 25, 2033	DP			
	10583121	Jul 25, 2033	DP U-2763			
	10814002	Jul 25, 2033	DP U-2763			
	11123305	Jul 25, 2033	DP			
	11813364	Sep 22, 2033	DP			
	9687474	Jul 25, 2033	DP			
<u>ASENAPINE - SECUADO</u>						
N 212268 003	10022445	Jul 25, 2033	DP			
	10583121	Jul 25, 2033	DP U-2763			
	10814002	Jul 25, 2033	DP U-2763			
	11123305	Jul 25, 2033	DP			
	11813364	Sep 22, 2033	DP			
	9687474	Jul 25, 2033	DP			
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117 001	7741358	Apr 06, 2026	DS DP U-1064			
	7741358	Apr 06, 2026	DS DP U-1960			
	7741358	Apr 06, 2026	DS DP U-1961			
	7741358	Apr 06, 2026	DS DP U-1962			
	7741358	Apr 06, 2026	DS DP U-1963			
	7741358*PED	Oct 06, 2026				
	8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117 002	7741358	Apr 06, 2026	DS DP U-1064			
	7741358	Apr 06, 2026	DS DP U-1960			
	7741358	Apr 06, 2026	DS DP U-1961			
	7741358	Apr 06, 2026	DS DP U-1962			
	7741358	Apr 06, 2026	DS DP U-1963			
	7741358*PED	Oct 06, 2026				
	8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<u>ASENAPINE MALEATE - SAPHRIS</u>						
N 022117 003	7741358	Apr 06, 2026	DS DP U-1893			
	7741358	Apr 06, 2026	DS DP U-1966			
	7741358*PED	Oct 06, 2026				
	8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<u>ASPIRIN - VAZALORE</u>						
N 203697 001	10646431	Sep 29, 2032	DP			
	10786444	Sep 29, 2032	U-3559			
	9216150	Sep 29, 2032	DP			
	9226892	Sep 29, 2032	U-1731			
	9226892	Sep 29, 2032	U-1732			
	9226892	Sep 29, 2032	U-1733			
<u>ASPIRIN - VAZALORE</u>						
N 203697 002	10646431	Sep 29, 2032	DP			
	10786444	Sep 29, 2032	U-3559			

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<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103	001 9539214	Mar 13, 2033	U-1902			
	9987231	Jan 02, 2033	U-2324			
<u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u>						
N 205103	002 9539214	Mar 13, 2033	U-1902			
	9987231	Jan 02, 2033	U-2324			
<u>ATAZANAVIR SULFATE; COBICISTAT - EVOTAZ</u>						
N 206353	001 10039718	Oct 06, 2032	DP			
	8148374	Sep 03, 2029	DS DP U-1279			
<u>ATOGEFANT - QULIPTA</u>						
N 215206	001 10117836	Jan 30, 2035	DP		I-909	Apr 17, 2026
	8754096	Jul 19, 2032	DS DP U-3534		NCE	Sep 28, 2026
	9499545	Nov 10, 2031	DS DP U-3534			
	9850246	Mar 13, 2033	DS			
<u>ATOGEFANT - QULIPTA</u>						
N 215206	002 10117836	Jan 30, 2035	DP		I-909	Apr 17, 2026
	8754096	Jul 19, 2032	DS DP U-3534		NCE	Sep 28, 2026
	9499545	Nov 10, 2031	DS DP U-3534			
	9850246	Mar 13, 2033	DS			
<u>ATOGEFANT - QULIPTA</u>						
N 215206	003 10117836	Jan 30, 2035	DP		I-909	Apr 17, 2026
	8754096	Jul 19, 2032	DS DP U-3534		NCE	Sep 28, 2026
	9499545	Nov 10, 2031	DS DP U-3534			
	9850246	Mar 13, 2033	DS			
<u>ATORVASTATIN CALCIUM - ATORVALIO</u>						
N 213260	001 11369567	Jun 07, 2037	DP			
	11654106	Jun 07, 2037	DP U-3612			
	11654106	Jun 07, 2037	DP U-3613			
<u>AVACINCAPTAD PEGOL SODIUM - IZERVAY</u>						
N 217225	001 10947544	Feb 14, 2025	DS U-3673			
	11273171	Jul 11, 2034	U-3673			
	11491176	Jul 11, 2034	U-3673			
	7538211	Feb 14, 2025	DS			
	7579456	Feb 14, 2025	DS			
	7803931	Feb 14, 2025	DS			
	8236773	Nov 11, 2026	U-3673			
	9617546	Feb 14, 2025	DS U-3673			
<u>AVACOPAN - TAVNEOS</u>						
N 214487	001 11603356	May 29, 2041	DS DP U-3558		NCE	Oct 07, 2026
	8445515	Feb 03, 2031	DS DP		ODE-377	Oct 07, 2028
	8906938	Dec 21, 2029	DS DP			
<u>AVANAFIL - STENDRA</u>						
N 202276	001 6656935	Apr 27, 2025	DS DP U-155		M-282	Oct 18, 2025
<u>AVANAFIL - STENDRA</u>						
N 202276	002 6656935	Apr 27, 2025	DS DP U-155		M-282	Oct 18, 2025

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<u>AVANAFIL - STENDRA</u>						
N 202276	003 6656935	Apr 27, 2025	DS DP U-155		M-282	Oct 18, 2025
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	001 9200002	Oct 15, 2034	DS DP U-2726		I-863	Jun 16, 2024
	9200002	Oct 15, 2034	DS DP U-3168		I-864	Jun 16, 2024
	9944651	Oct 15, 2034	DS DP U-2726		NCE	Jan 09, 2025
	9944651	Oct 15, 2034	DS DP U-3168		ODE-356	Jun 16, 2028
	9994575	Oct 15, 2034	DS DP U-2726		ODE-366	Jan 09, 2027
	9994575	Oct 15, 2034	DS DP U-3168		ODE-434	May 22, 2030
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	002 9200002	Oct 15, 2034	DS DP U-2726		I-863	Jun 16, 2024
	9200002	Oct 15, 2034	DS DP U-3168		I-864	Jun 16, 2024
	9944651	Oct 15, 2034	DS DP U-2726		NCE	Jan 09, 2025
	9944651	Oct 15, 2034	DS DP U-3168		ODE-356	Jun 16, 2028
	9994575	Oct 15, 2034	DS DP U-2726		ODE-366	Jan 09, 2027
	9994575	Oct 15, 2034	DS DP U-3168		ODE-434	May 22, 2030
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	003 9200002	Oct 15, 2034	DS DP U-2726		NCE	Jan 09, 2025
	9944651	Oct 15, 2034	DS DP U-2726		ODE-356	Jun 16, 2028
	9994575	Oct 15, 2034	DS DP U-2726		ODE-366	Jan 09, 2027
					ODE-434	May 22, 2030
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	004 11827642	Oct 15, 2034	DS DP U-3506		I-863	Jun 16, 2024
	9200002	Oct 15, 2034	DS DP U-3168		I-864	Jun 16, 2024
	9200002	Oct 15, 2034	DS DP U-3506		I-912	May 22, 2026
	9944651	Oct 15, 2034	DS DP U-3168		NCE	Jan 09, 2025
	9944651	Oct 15, 2034	DS DP U-3506		ODE-356	Jun 16, 2028
	9994575	Oct 15, 2034	DS DP U-3168		ODE-366	Jan 09, 2027
	9994575	Oct 15, 2034	DS DP U-3506		ODE-434	May 22, 2030
<u>AVAPRITINIB - AYWAKIT</u>						
N 212608	005 9200002	Oct 15, 2034	DS DP U-3168		I-863	Jun 16, 2024
	9944651	Oct 15, 2034	DS DP U-3168		I-864	Jun 16, 2024
	9994575	Oct 15, 2034	DS DP U-3168		NCE	Jan 09, 2025
					ODE-356	Jun 16, 2028
					ODE-366	Jan 09, 2027
					ODE-434	May 22, 2030
<u>AVATROMBOPAG MALEATE - DOPTELET</u>						
N 210238	001 7638536	May 05, 2025	DS DP		ODE-246	Jun 26, 2026
<u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u>						
N 206494	001 7112592	Jan 07, 2026	DS DP U-2244		NCE	Feb 25, 2020
	7112592	Jan 07, 2026	DS DP U-2508		NPP	Dec 20, 2025
	7112592	Jan 07, 2026	DS DP U-282		GAIN	Feb 25, 2025
	7612087	Nov 12, 2026	DP			
	8471025	Aug 12, 2031	DS			
	8835455	Oct 08, 2030	DP			
	8969566	Jun 15, 2032	DS			
	9284314	Jun 15, 2032	DS			
	9695122	Jun 15, 2032	DS			

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<u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u>						
N 206494	001	7112592	Jan 07, 2026	DS DP U-2244	NCE	Feb 25, 2020
		7112592	Jan 07, 2026	DS DP U-2508	NPP	Dec 20, 2025
		7112592	Jan 07, 2026	DS DP U-282	GAIN	Feb 25, 2025
		7612087	Nov 12, 2026	DP		
		8471025	Aug 12, 2031	DS		
		8835455	Oct 08, 2030	DP		
		8969566	Jun 15, 2032	DS		
		9284314	Jun 15, 2032	DS		
		9695122	Jun 15, 2032	DS		
<u>AXITINIB - INLYTA</u>						
N 202324	001	10570202	Feb 03, 2035	U-2844		
		10869924	Jan 12, 2037	U-3044		
		6534524	Apr 29, 2025	DS DP		
		8791140	Dec 14, 2030	DS		
<u>AXITINIB - INLYTA</u>						
N 202324	002	10570202	Feb 03, 2035	U-2844		
		10869924	Jan 12, 2037	U-3044		
		6534524	Apr 29, 2025	DS DP		
		8791140	Dec 14, 2030	DS		
<u>AZACITIDINE - VIDAZA</u>						
N 050794	001				I-889	May 20, 2025
					ODE-399	May 20, 2029
<u>AZACITIDINE - ONUREG</u>						
N 214120	001	11571436	May 14, 2029	DP	ODE-320	Sep 01, 2027
		8846628	Jun 03, 2030	DP U-2950		
<u>AZACITIDINE - ONUREG</u>						
N 214120	002	11571436	May 14, 2029	DP	ODE-320	Sep 01, 2027
		8846628	Jun 03, 2030	DP U-2950		
<u>AZELAIC ACID - FINACEA</u>						
N 207071	001	10117812	Oct 18, 2027	DP U-1796		
		7700076	Sep 18, 2027	DP		
		8435498	Mar 01, 2024	U-1727		
		9211259	Feb 28, 2029	U-1796		
		9265725	Dec 08, 2027	DP		
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u>						
N 022203	001	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-1430		
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u>						
N 022203	002	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-1430		
		9919050	Nov 22, 2025	DP		
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO ALLERGY</u>						
N 213872	001	8071073	Jun 04, 2028	DP		
		8518919	Nov 22, 2025	U-3166		
		9919050	Nov 22, 2025	DP		

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<u>AZELASTINE HYDROCHLORIDE - CHILDREN'S ASTEPRO ALLERGY</u>						
N 213872	002 8071073	Jun 04, 2028	DP			
	8518919	Nov 22, 2025	U-3166			
	9919050	Nov 22, 2025	DP			
<u>AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE - DYMISTA</u>						
N 202236	001 8163723*PED	Feb 29, 2024				
	8168620	Feb 24, 2026	DP			
<u>AZILSARTAN KAMEDOXOMIL - EDARBI</u>						
N 200796	001 7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
<u>AZILSARTAN KAMEDOXOMIL - EDARBI</u>						
N 200796	002 7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>						
N 202331	001 7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
	9169238	Feb 04, 2030	DP			
	9387249	Jul 01, 2031	U-3			
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>						
N 202331	002 7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
	9169238	Feb 04, 2030	DP			
	9387249	Jul 01, 2031	U-3			
<u>AZITHROMYCIN - ZMAX</u>						
N 050797	001 6984403	Feb 14, 2024	DP U-282			
	7887844	Feb 14, 2024	DP			
<u>BACLOFEN - OZOBAX</u>						
N 208193	001 10610502	Aug 30, 2039	U-2779			
<u>BACLOFEN - LYVISPAH</u>						
N 215422	001 10792262	Jul 29, 2039	DP U-3263			
	11491125	Sep 29, 2041	DP U-3488			
	11491125	Sep 29, 2041	DP U-3489			
	11654124	Jul 29, 2039	DP			
<u>BACLOFEN - LYVISPAH</u>						
N 215422	002 10792262	Jul 29, 2039	DP U-3263			
	11491125	Sep 29, 2041	DP U-3488			
	11491125	Sep 29, 2041	DP U-3489			
	11654124	Jul 29, 2039	DP			
<u>BACLOFEN - LYVISPAH</u>						
N 215422	003 10792262	Jul 29, 2039	DP U-3263			
	11491125	Sep 29, 2041	DP U-3488			
	11491125	Sep 29, 2041	DP U-3489			
	11654124	Jul 29, 2039	DP			

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<u>BACLOFEN - LYVISPAN</u>						
N 215422	003	10792262	Jul 29, 2039	DP	U-3263	
		11491125	Sep 29, 2041	DP	U-3488	
		11491125	Sep 29, 2041	DP	U-3489	
		11654124	Jul 29, 2039	DP		
<u>BACLOFEN - FLEOSUVY</u>						
N 215602	001	11324696	Sep 29, 2037	DP		
		11446246	Sep 08, 2037		U-3433	
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	001	10392406	Apr 27, 2036	DS		
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		11261198	Sep 25, 2038	DP		
		11306106	Aug 09, 2037		U-2816	
		11306106	Aug 09, 2037		U-3000	
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	002	10392406	Apr 27, 2036	DS		
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		11261198	Sep 25, 2038	DP		
		11306106	Aug 09, 2037		U-2816	
		11306106	Aug 09, 2037		U-3000	
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 210854	003	10392406	Apr 27, 2036	DS		
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		11261198	Sep 25, 2038	DP		
		11306106	Aug 09, 2037		U-2816	
		11306106	Aug 09, 2037		U-3000	
		8927710	May 05, 2031	DP		
		8987441	Sep 21, 2031	DS DP		
		9815835	Jun 14, 2030	DP		
<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 214410	001	10392406	Apr 27, 2036	DS		
		10633397	Apr 27, 2036		U-2816	
		10633397	Apr 27, 2036		U-3000	
		10759814	Aug 09, 2037	DS DP		
		11261198	Sep 25, 2038	DP		
		11306106	Aug 09, 2037		U-2816	
		11306106	Aug 09, 2037		U-3000	

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<u>BALOXAVIR MARBOXIL - XOFLUZA</u>						
N 214410	001 8927710	May 05, 2031	DP			
	8987441	Sep 21, 2031	DS DP			
	9815835	Jun 14, 2030	DP			
<u>BALSALAZIDE DISODIUM - COLAZAL</u>						
N 020610	001 7452872	Aug 24, 2026	U-141			
	7625884	Aug 24, 2026	U-141			
<u>BALSALAZIDE DISODIUM - GIAZO</u>						
N 022205	001 7452872	Aug 24, 2026	U-1229			
	7625884	Aug 24, 2026	U-1229			
	8497256	Jun 23, 2031	U-1229			
	9192616	Aug 02, 2026	U-1229			
<u>BARICITINIB - OLUMIANT</u>						
N 207924	001 11045474	Nov 30, 2032	U-3372		I-890	Jun 13, 2025
	11806555	Nov 02, 2031	U-3500		I-891	May 10, 2025
	8158616	Jun 08, 2030	DS DP			
	8420629	Mar 10, 2029	U-247			
	9089574	Nov 30, 2032	U-3372			
	9737469	Nov 02, 2031	U-3500			
<u>BARICITINIB - OLUMIANT</u>						
N 207924	002 11045474	Nov 30, 2032	U-3372		I-890	Jun 13, 2025
	11806555	Nov 02, 2031	U-3500		I-891	May 10, 2025
	8158616	Jun 08, 2030	DS DP			
	8420629	Mar 10, 2029	U-247			
	9089574	Nov 30, 2032	U-3372			
	9737469	Nov 02, 2031	U-3500			
<u>BARICITINIB - OLUMIANT</u>						
N 207924	003 11045474	Nov 30, 2032	U-3372		I-890	Jun 13, 2025
	11806555	Nov 02, 2031	U-3500		I-891	May 10, 2025
	8158616	Jun 08, 2030	DS DP			
	9089574	Nov 30, 2032	U-3372			
	9737469	Nov 02, 2031	U-3500			
<u>BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED - DUAVEE</u>						
N 022247	001 6479535	May 06, 2024	DP U-594			
	6479535	May 06, 2024	DP U-904			
	7683051	Mar 10, 2027	DS DP U-594			
	7683051	Mar 10, 2027	DS DP U-904			
<u>BECLMETHASONE DIPROPIONATE - QVAR 80</u>						
N 020911	001 10022509	May 18, 2031	DP			
	10022510	May 18, 2031	DP			
	10086156	May 18, 2031	DP			
	10561808	Jan 01, 2032	DP			
	10695512	May 18, 2031	DP			
	11395889	May 18, 2031	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			

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<u>BECLOMETHASONE DIPROPIONATE - QVAR 40</u>						
N 020911	002	10022509	May 18, 2031	DP		
		10022510	May 18, 2031	DP		
		10086156	May 18, 2031	DP		
		10561808	Jan 01, 2032	DP		
		10695512	May 18, 2031	DP		
		11395889	May 18, 2031	DP		
		9463289	May 18, 2031	DP		
		9808587	May 18, 2031	DP		
<u>BECLOMETHASONE DIPROPIONATE - QNASL</u>						
N 202813	001	10188811	Oct 21, 2031	DP		
		7780038	Jan 24, 2027	DP		
<u>BECLOMETHASONE DIPROPIONATE - QNASL</u>						
N 202813	002	10188811	Oct 21, 2031	DP		
		7780038	Jan 24, 2027	DP		
<u>BECLOMETHASONE DIPROPIONATE - QVAR REDIHALER</u>						
N 207921	001	10022509	May 18, 2031	DP		
		10022510	May 18, 2031	DP		
		10086156	May 18, 2031	DP		
		10561808	Jan 01, 2032	DP		
		10695512	May 18, 2031	DP		
		10792447	Jan 25, 2039	DP		
		11395888	Jan 26, 2038	DP		
		11395889	May 18, 2031	DP		
		11559637	Jul 21, 2039	DP		
		11583643	Aug 19, 2041	DP		
		11793953	Jan 26, 2038	DP		
		8132712	Sep 07, 2028	DP		
		8931476	Jul 17, 2031	DP		
<u>BECLOMETHASONE DIPROPIONATE - QVAR REDIHALER</u>						
N 207921	002	10022509	May 18, 2031	DP		
		10022510	May 18, 2031	DP		
		10086156	May 18, 2031	DP		
		10561808	Jan 01, 2032	DP		
		10695512	May 18, 2031	DP		
		10792447	Jan 25, 2039	DP		
		11395888	Jan 26, 2038	DP		
		11395889	May 18, 2031	DP		
		11559637	Jul 21, 2039	DP		
		11583643	Aug 19, 2041	DP		
		11793953	Jan 26, 2038	DP		
		8132712	Sep 07, 2028	DP		
		8931476	Jul 17, 2031	DP		
<u>BEDAQUILINE FUMARATE - SIRTURO</u>						
N 204384	001	7498343	Dec 01, 2026	DS DP U-1321	ODE-251	Aug 09, 2026
		8546428	Mar 19, 2029	DS DP U-1321	ODE-307	May 27, 2027

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<u>BEDAQUILINE FUMARATE - SIRTURO</u>						
N 204384	002 7498343	Dec 01, 2026	DS DP U-1321		ODE-307	May 27, 2027
	8546428	Mar 19, 2029	DS DP U-1321			
<u>BELINOSTAT - BELEODAO</u>						
N 206256	001 6888027	Aug 10, 2026	DS DP U-1544			
	8835501	Oct 27, 2027	DP			
<u>BELUMOSUDIL MESYLATE - REZUROCK</u>						
N 214783	001 10183931	Oct 07, 2033		U-3246	NCE	Jul 16, 2026
	10696660	Oct 07, 2033		U-3246	ODE-362	Jul 16, 2028
	11311541	Apr 09, 2035		U-3369		
	8357693	Oct 30, 2029	DS DP U-3247			
	9815820	Oct 07, 2033		U-3247		
<u>BELZUTIFAN - WELIREG</u>						
N 215383	001 9908845	Sep 05, 2034	DS DP U-3201		NCE	Aug 13, 2026
	9908845	Sep 05, 2034	DS DP U-3780		ODE-364	Aug 13, 2028
	9969689	Sep 05, 2034	DS DP U-3201			
	9969689	Sep 05, 2034	DS DP U-3780			
<u>BEMPEDOIC ACID - NEXLETOL</u>						
N 211616	001 11613511	Jun 19, 2040	DS		NCE	Feb 21, 2025
	11760714	Jun 19, 2040	DP			
	7335799	Dec 03, 2025	DS			
<u>BEMPEDOIC ACID; EZETIMIBE - NEXLIZET</u>						
N 211617	001 10912751	Mar 14, 2036		U-3224	NCE	Feb 21, 2025
	11613511	Jun 19, 2040	DS			
	11744816	Mar 14, 2036		U-3692		
	11760714	Jun 19, 2040	DP			
	7335799	Dec 03, 2025	DS			
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249	001 8436190	Oct 26, 2030	DP			
	8436190*PED	Apr 26, 2031				
	8445524	Mar 26, 2029	DS DP U-1402			
	8445524*PED	Sep 26, 2029				
	8609863	Jan 12, 2026	DP			
	8609863*PED	Jul 12, 2026				
	8669279	Mar 26, 2029	DP U-1402			
	8669279*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
	8883836	Mar 26, 2029	DP U-1402			
	8883836*PED	Sep 26, 2029				
	8895756	Jan 12, 2026	DP			
	8895756*PED	Jul 12, 2026				
	9533955	Mar 26, 2029	DP U-1949			
	9533955	Mar 26, 2029	DP U-1952			
	9533955*PED	Sep 26, 2029				
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249	002 8436190	Oct 26, 2030	DP			
	8436190*PED	Apr 26, 2031				

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<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249 002	8445524	Mar 26, 2029	DS DP U-1402			
	8445524*PED	Sep 26, 2029				
	8609863	Jan 12, 2026	DP			
	8609863*PED	Jul 12, 2026				
	8669279	Mar 26, 2029	DP U-1402			
	8669279*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
	8883836	Mar 26, 2029	DP U-1402			
	8883836*PED	Sep 26, 2029				
	8895756	Jan 12, 2026	DP			
	8895756*PED	Jul 12, 2026				
	9533955	Mar 26, 2029	DP U-1949			
	9533955	Mar 26, 2029	DP U-1952			
	9533955*PED	Sep 26, 2029				
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249 003	8344006	Sep 23, 2029	DP U-1402			
	8344006*PED	Mar 23, 2030				
	8445524	Mar 26, 2029	DS			
	8445524*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N 022249 004	8344006	Sep 23, 2029	DP U-1402			
	8344006*PED	Mar 23, 2030				
	8445524	Mar 26, 2029	DS			
	8445524*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
<u>BENDAMUSTINE HYDROCHLORIDE - BELRAPZO</u>						
N 205580 001	10010533	Jan 28, 2031	DP			
	11103483	Jan 28, 2031	DP U-1971			
	11103483	Jan 28, 2031	DP U-1972			
	11844783	Jan 28, 2031	U-1542			
	11844783	Jan 28, 2031	U-1971			
	11844783	Jan 28, 2031	U-1972			
	8609707	Aug 11, 2031	DP U-1971			
	8609707	Aug 11, 2031	DP U-1972			
	8791270	Jan 12, 2026	DP U-1971			
	8791270	Jan 12, 2026	DP U-1972			
	9265831	Jan 28, 2031	DP			
	9572796	Jan 28, 2031	DP U-1971			
	9572796	Jan 28, 2031	DP U-1972			
	9572797	Jan 28, 2031	U-1971			
	9572797	Jan 28, 2031	U-1972			
<u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u>						
N 208194 001	10010533	Jan 28, 2031	DP			
	10052385	Mar 15, 2033	U-1971			

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<u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u>						
N 208194	001	10052385	Mar 15, 2033	U-1972		
		11103483	Jan 28, 2031	DP U-1971		
		11103483	Jan 28, 2031	DP U-1972		
		11844783	Jan 28, 2031	U-1542		
		11844783	Jan 28, 2031	U-1971		
		11844783	Jan 28, 2031	U-1972		
		8609707	Aug 11, 2031	DP U-1542		
		8791270	Jan 12, 2026	DP U-1790		
		9000021	Mar 15, 2033	U-1542		
		9034908	Mar 15, 2033	U-1542		
		9144568	Mar 15, 2033	U-1542		
		9265831	Jan 28, 2031	DP		
		9572796	Jan 28, 2031	DP U-1971		
		9572796	Jan 28, 2031	DP U-1972		
		9572797	Jan 28, 2031	U-1971		
		9572797	Jan 28, 2031	U-1972		
		9572887	Mar 15, 2033	U-1971		
		9572887	Mar 15, 2033	U-1972		
		9579384	Mar 15, 2033	U-1971		
		9579384	Mar 15, 2033	U-1972		
		9597397	Mar 15, 2033	U-1971		
		9597397	Mar 15, 2033	U-1972		
		9597398	Mar 15, 2033	U-1971		
		9597399	Mar 15, 2033	U-1971		
		9597399	Mar 15, 2033	U-1972		
<u>BENDAMUSTINE HYDROCHLORIDE - VIVIMUSTA</u>						
N 212209	001	11844784	Jul 29, 2042	DP		
<u>BENOXINATE HYDROCHLORIDE; FLUORESCEIN SODIUM - FLUORESCEIN SODIUM AND BENOXINATE HYDROCHLORIDE</u>						
N 211039	001	10293047	Nov 15, 2037	DP U-2755		
		10632197	Nov 15, 2037	DP U-2755		
		10842872	Nov 15, 2037	U-3001		
<u>BENZNIDAZOLE - BENZNIDAZOLE</u>						
N 209570	001				ODE-154	Aug 29, 2024
<u>BENZNIDAZOLE - BENZNIDAZOLE</u>						
N 209570	002				ODE-154	Aug 29, 2024
<u>BENZOYL PEROXIDE - EPSOLAY</u>						
N 214510	001	10933046	Feb 19, 2040	DP U-3357	NP	Apr 22, 2025
		10945987	Feb 19, 2040	U-3356		
		11426378	Aug 18, 2040	U-3356		
		11541026	Feb 19, 2040	U-3356		
		11628155	Dec 27, 2040	U-3356		
		9687465	Nov 27, 2032	DP U-3356		
		9868103	Aug 08, 2028	DP U-3356		
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE</u>						
A 208683	001				PC	Mar 30, 2024

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<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u>						
N 050819 001	10220049	Jun 03, 2029	DP U-916			
	10624918	Jun 03, 2029	U-916			
	8288434	Aug 05, 2029	DP U-124			
	8663699	Jun 03, 2029	U-124			
	8895070	Jun 03, 2029	U-124			
	9078870	Jun 03, 2029	DP			
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ONEXTON</u>						
N 050819 002	10137142	Jun 03, 2029	DP U-916			
	10220049	Jun 03, 2029	DP U-916			
	8288434	Aug 05, 2029	DP U-1033			
	8288434	Aug 05, 2029	DP U-124			
	8288434	Aug 05, 2029	DP U-134			
	8288434	Aug 05, 2029	DP U-818			
	8288434	Aug 05, 2029	DP U-916			
	8288434	Aug 05, 2029	DP U-921			
	9504704	Jun 03, 2029	DP U-124			
	9504704	Jun 03, 2029	DP U-134			
	9504704	Jun 03, 2029	DP U-818			
	9504704	Jun 03, 2029	DP U-916			
	9561208	Jun 03, 2029	DP U-916			
<u>BENZOYL PEROXIDE; TRETINOIN - TWYNEO</u>						
N 214902 001	10420743	Jul 12, 2038	U-3194		NC	Jul 26, 2024
	10653899	Dec 30, 2030	DP U-3194			
	11071878	Dec 30, 2030	DP			
	8617580	Feb 03, 2028	DP			
	9868103	Aug 08, 2028	DP U-3194			
<u>BENZYL ALCOHOL - ULESFIA</u>						
N 022129 001	7294342	May 19, 2024	U-970			
<u>BEPOTASTINE BESILATE - BEPREVE</u>						
N 022288 001	8784789	Jan 13, 2025	DP			
<u>BEROTRALSTAT HYDROCHLORIDE - ORLADEYO</u>						
N 214094 001	10125102	Apr 07, 2035	DS U-3010		NCE	Dec 03, 2025
	10329260	Mar 09, 2035	DS		ODE-333	Dec 03, 2027
	10662160	Nov 01, 2039	DS U-3010			
	10689346	Mar 09, 2035	U-3010			
	11117867	Nov 01, 2039	DP U-3010			
	11230530	Mar 09, 2035	U-3300			
	11618733	Nov 01, 2039	U-3300			
	11708333	Mar 09, 2035	U-3300			
<u>BEROTRALSTAT HYDROCHLORIDE - ORLADEYO</u>						
N 214094 002	10125102	Apr 07, 2035	DS U-3010		NCE	Dec 03, 2025
	10329260	Mar 09, 2035	DS		ODE-333	Dec 03, 2027
	10662160	Nov 01, 2039	DS U-3010			
	10689346	Mar 09, 2035	U-3010			
	11117867	Nov 01, 2039	DP U-3010			
	11230530	Mar 09, 2035	U-3300			
	11618733	Nov 01, 2039	U-3300			

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<u>BEROTRALSTAT HYDROCHLORIDE - ORLADEYO</u>						
N 214094	002 11708333	Mar 09, 2035	U-3300			
<u>BESIFLOXACIN HYDROCHLORIDE - BESIVANCE</u>						
N 022308	001 8415342	Nov 07, 2030	U-80			
	8481526	Jan 09, 2031	DS			
	8604020	Mar 12, 2030	DP			
	8937062	Nov 13, 2029	U-80			
<u>BETAMETHASONE DIPROPIONATE - SERNIVO</u>						
N 208079	001 10179137	Aug 31, 2030	DP U-1858			
	9364485	Aug 31, 2030	DP U-1858			
	9433630	Aug 31, 2030	DP U-1858			
	9439911	Aug 31, 2030	DP U-1858			
	9655907	Aug 31, 2030	DP U-1858			
	9775851	Aug 31, 2030	DP U-1858			
	9877974	Aug 31, 2030	DP U-1858			
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u>						
N 207589	001 10130640	Jun 10, 2031	DP			
	10130640*PED	Dec 10, 2031				
	10617698	Jun 10, 2031	DP			
	10660908	Jun 10, 2031	DP U-2627			
	10682364	Jun 10, 2031	DP			
	10688108	Jun 10, 2031	U-2627			
	10716799	Jun 10, 2031	DP			
	9119781	Jun 10, 2031	DP U-1761			
	9119781	Jun 10, 2031	DP U-2627			
	9119781*PED	Dec 10, 2031				
	9566286	Jun 10, 2031	DP			
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - WYNZORA</u>						
N 213422	001 10265265	Sep 27, 2027	DP			
	11696919	Mar 18, 2039	DP			
<u>BETRIXABAN - BEVYXXA</u>						
N 208383	001 7598276	Nov 08, 2026	DS			
	8404724	Mar 29, 2031	DP U-2034			
	8557852	Sep 08, 2028	U-1167			
	8557852	Sep 08, 2028	U-2030			
	8987463	Dec 28, 2030	DP			
	9555023	Nov 07, 2026	U-1502			
<u>BETRIXABAN - BEVYXXA</u>						
N 208383	002 7598276	Nov 08, 2026	DS			
	8404724	Mar 29, 2031	DP U-2034			
	8557852	Sep 08, 2028	U-1167			
	8557852	Sep 08, 2028	U-2030			
	8987463	Dec 28, 2030	DP			
	9555023	Nov 07, 2026	U-1502			
<u>BEXAGLIFLOZIN - BRENZAVVY</u>						
N 214373	001 10533032	Jul 03, 2031	U-2214		NCE	Jan 20, 2028
	10981942	Jun 13, 2031	DS DP			
	7838499	Jan 30, 2029	DS DP U-2214			

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<u>BEXAGLIFLOZIN - BRENZAVVY</u>						
N 214373	001 8106021	Aug 22, 2028	U-2214			
	8802637	Aug 22, 2028	DS DP U-2214			
	8987323	May 14, 2032	DS DP			
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	001 10385067	Jun 19, 2035	U-257		M-82	Feb 24, 2024
	10548846	Nov 08, 2036	DP		ODE-256	Jun 18, 2026
	11744802	Nov 08, 2036	DP			
	7390791	Apr 17, 2025	DS DP			
	7390791*PED	Oct 17, 2025				
	8754065	Aug 15, 2032	DS DP U-257			
	8754065*PED	Feb 15, 2033				
	9216996	Dec 19, 2033	DS DP			
	9296769	Aug 15, 2032	DS DP U-257			
	9296769*PED	Feb 15, 2033				
	9708342	Jun 19, 2035	DS DP			
	9732092	Dec 19, 2033	DS DP			
<u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u>						
N 210251	002 10385067	Jun 19, 2035	U-257		ODE-378	Oct 07, 2028
	7390791	Apr 17, 2025	DS DP			
	8754065	Aug 15, 2032	DS DP U-257			
	9216996	Dec 19, 2033	DS DP			
	9296769	Aug 15, 2032	DS DP U-257			
	9708342	Jun 19, 2035	DS DP			
	9732092	Dec 19, 2033	DS DP			
<u>BIMATOPROST - LUMIGAN</u>						
N 022184	001 7851504	Jun 13, 2027	DS DP			
	8278353	Mar 16, 2025	DP			
	8299118	Mar 16, 2025	U-1295			
	8309605	Mar 16, 2025	U-1293			
	8309605	Mar 16, 2025	U-1294			
	8338479	Mar 16, 2025	DP U-1295			
	8524777	Mar 16, 2025	U-1235			
	8586630	Mar 16, 2025	U-1458			
	8772338	Mar 16, 2025	DP U-1528			
	8933120	Mar 16, 2025	DP			
	8933127	Mar 16, 2025	DP			
	9155716	Mar 16, 2025	DP U-1528			
	9241918	Mar 16, 2025	DP U-1814			
<u>BIMATOPROST - LATISSE</u>						
N 022369	001 8101161	May 25, 2024	U-1217			
	8101161	May 25, 2024	U-1218			
<u>BIMATOPROST - DURYSTA</u>						
N 211911	001 10398707	Apr 30, 2024	U-2759			
	10441543	Dec 19, 2026	DP			
	7799336	Apr 24, 2029	DP			
	8206737	Apr 07, 2027	U-2759			
	8629185	Jul 15, 2031	DS DP			
	8673341	Feb 19, 2025	U-2759			

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<u>BIMATOPROST - DURYSTA</u>						
N 211911	001 9149428	Dec 19, 2026	DP			
	9492316	Oct 31, 2034	DP			
	9980974	Oct 31, 2034		U-2759		
<u>BINIMETINIB - MEKTOVI</u>						
N 210498	001 10005761	Aug 27, 2030		U-2331	I-928	Oct 11, 2026
	10005761	Aug 27, 2030		U-3737	ODE-194	Jun 27, 2025
	7777050	Mar 13, 2024	DS DP			
	9314464	Jul 04, 2031		U-2332		
	9314464	Jul 04, 2031		U-3737		
	9562016	Oct 18, 2033	DS DP			
	9598376	Oct 18, 2033		U-2330		
	9850229	Aug 27, 2030		U-2333		
	9980944	Oct 18, 2033		U-2334		
<u>BIVALIRUDIN - ANGIOMAX</u>						
N 020873	001 7582727	Jul 27, 2028	DP			
	7598343	Jul 27, 2028	DP			
<u>BOCEPREVIR - VICTRELIS</u>						
N 202258	001 7772178	Nov 11, 2027	DP	U-1128		
	8119602	Mar 17, 2027		U-1233		
	RE43298	Dec 22, 2024	DS DP	U-1128		
<u>BORTEZOMIB - BORTEZOMIB</u>						
N 205004	001 8962572	Nov 03, 2032	DP			
<u>BORTEZOMIB - BORTEZOMIB</u>						
N 215331	001 11679119	Sep 23, 2042		U-3632		
	11679119	Sep 23, 2042		U-3633		
	11752164	Sep 23, 2042		U-3632		
	11752164	Sep 23, 2042		U-3633		
<u>BORTEZOMIB - BORTEZOMIB</u>						
N 215331	002 11679119	Sep 23, 2042		U-3632		
	11679119	Sep 23, 2042		U-3633		
	11752164	Sep 23, 2042		U-3632		
	11752164	Sep 23, 2042		U-3633		
<u>BOSENTAN - TRACLEER</u>						
N 021290	001				ODE*	Sep 05, 2024
<u>BOSENTAN - TRACLEER</u>						
N 021290	002				ODE*	Sep 05, 2024
<u>BOSENTAN - TRACLEER</u>						
N 209279	001 7959945	Dec 28, 2027	DP		ODE-161	Sep 05, 2024
	8309126	May 15, 2026	DP			
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341	001 11103497	Feb 28, 2034		U-3216	I-923	Sep 26, 2026
	11103497	Feb 28, 2034		U-3217	ODE-163	Dec 19, 2024
	11103497*PED	Aug 28, 2034			ODE-444	Sep 26, 2030
	7417148	Dec 11, 2025		U-1283	PED	Jun 19, 2025
	7417148	Dec 11, 2025		U-3707	PED	Mar 26, 2027
	7417148	Dec 11, 2025		U-3708	PED	Mar 26, 2031

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<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 001	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
	RE42376	Apr 13, 2024	DS			
	RE42376*PED	Oct 13, 2024				
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 002	11103497	Feb 28, 2034	U-3216		I-923	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-163	Dec 19, 2024
	11103497*PED	Aug 28, 2034			ODE-444	Sep 26, 2030
	7417148	Dec 11, 2025	U-1283		PED	Jun 19, 2025
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3708		PED	Mar 26, 2031
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
	RE42376	Apr 13, 2024	DS			
	RE42376*PED	Oct 13, 2024				
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 203341 003	11103497	Feb 28, 2034	U-3216		I-923	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-163	Dec 19, 2024
	11103497*PED	Aug 28, 2034			ODE-444	Sep 26, 2030
	7417148	Dec 11, 2025	U-1283		PED	Jun 19, 2025
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3708		PED	Mar 26, 2031
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
	RE42376	Apr 13, 2024	DS			
	RE42376*PED	Oct 13, 2024				
<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 217729 001	11103497	Feb 28, 2034	U-3216		NP	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-444	Sep 26, 2030
	11103497*PED	Aug 28, 2034			PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2031
	7417148	Dec 11, 2025	U-3708			
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
	RE42376	Apr 13, 2024	DS DP			
	RE42376*PED	Oct 13, 2024				

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<u>BOSUTINIB MONOHYDRATE - BOSULIF</u>						
N 217729 002	11103497	Feb 28, 2034	U-3216		NP	Sep 26, 2026
	11103497	Feb 28, 2034	U-3217		ODE-444	Sep 26, 2030
	11103497*PED	Aug 28, 2034			PED	Mar 26, 2027
	7417148	Dec 11, 2025	U-3707		PED	Mar 26, 2031
	7417148	Dec 11, 2025	U-3708			
	7417148*PED	Jun 11, 2026				
	7767678	Nov 23, 2026	DS DP			
	7767678*PED	May 23, 2027				
	7919625	Dec 11, 2025	DP			
	7919625*PED	Jun 11, 2026				
	RE42376	Apr 13, 2024	DS DP			
	RE42376*PED	Oct 13, 2024				
<u>BREMELANOTIDE ACETATE - VYLEESI (AUTOINJECTOR)</u>						
N 210557 001	10286034	Nov 05, 2033	U-2568		NCE	Jun 21, 2024
	11590209	Apr 29, 2041	U-3539			
	6794489	Jun 28, 2025	DS DP			
	9352013	Nov 05, 2033	U-2568			
	9700592	Nov 05, 2033	U-2568			
<u>BREXANOLONE - ZULRESSO</u>						
N 211371 001	10117951	Mar 13, 2029	DP		NCE	Jun 17, 2024
	10251894	Nov 27, 2033	U-2552		NPP	Jun 16, 2025
	10322139	Jan 23, 2033	DP			
	10940156	Mar 08, 2037	U-2552			
	7635773	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9200088	Mar 13, 2029	DP			
	9750822	Mar 13, 2029	DP			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 001	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	7888362	Apr 12, 2026	DS	Y	NPP	Dec 27, 2024
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	RE48059	Dec 23, 2028	DS			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 002	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	7888362	Apr 12, 2026	DS	Y	NPP	Dec 27, 2024
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	RE48059	Dec 23, 2028	DS			

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<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 003	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	7888362	Apr 12, 2026	DS	Y	NPP	Dec 27, 2024
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	RE48059	Dec 23, 2028	DS			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 004	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	7888362	Apr 12, 2026	DS	Y	NPP	Dec 27, 2024
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	RE48059	Dec 23, 2028	DS			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 005	10307419	Oct 12, 2032	DP		I-913	May 10, 2026
	7888362	Apr 12, 2026	DS	Y	NPP	Dec 27, 2024
	8349840	Apr 12, 2026	DP U-1529			
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	RE48059	Dec 23, 2028	DS			
<u>BREXPIRAZOLE - REXULTI</u>						
N 205422 006	7888362	Apr 12, 2026	DS	Y	I-913	May 10, 2026
	8349840	Apr 12, 2026	DP U-1529		NPP	Dec 27, 2024
	8618109	Apr 12, 2026	U-3281			
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-3281			
	9839637	Apr 12, 2026	DP U-543			
	RE48059	Dec 23, 2028	DS			
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772 001	10385078	Nov 10, 2035	DS DP U-2837		ODE-142	Apr 28, 2024
	9012462	Apr 28, 2031	DS		ODE-300	May 22, 2027
	9273077	May 21, 2029	U-2837			
	9611283	Apr 10, 2034	U-2837			
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772 002	10385078	Nov 10, 2035	DS DP U-2837		ODE-142	Apr 28, 2024
	9012462	Apr 28, 2031	DS		ODE-300	May 22, 2027
	9273077	May 21, 2029	U-2837			
	9611283	Apr 10, 2034	U-2837			

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<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	002 10385078	Nov 10, 2035	DS DP U-2837		ODE-142	Apr 28, 2024
	9012462	Apr 28, 2031	DS		ODE-300	May 22, 2027
	9273077	May 21, 2029		U-2837		
	9611283	Apr 10, 2034		U-2837		
<u>BRIGATINIB - ALUNBRIG</u>						
N 208772	003 10385078	Nov 10, 2035	DS DP U-2837		ODE-142	Apr 28, 2024
	9012462	Apr 28, 2031	DS		ODE-300	May 22, 2027
	9273077	May 21, 2029		U-2837		
	9611283	Apr 10, 2034		U-2837		
<u>BRILLIANT BLUE G - TISSUEBLUE</u>						
N 209569	001				NCE	Dec 20, 2024
					ODE-282	Dec 20, 2026
<u>BRIMONIDINE TARTRATE - OOLIANA</u>						
N 021764	001 7265117	Aug 19, 2025	DP			
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>						
N 021770	001 8858961*PED	Mar 02, 2024				
<u>BRIMONIDINE TARTRATE - MIRVASO</u>						
N 204708	001 10201517	Jun 13, 2031	DP			
	7439241	Aug 25, 2025		U-1428		
	8053427	Jun 13, 2031	DP U-1428			
	8163725	Jun 13, 2031	DP			
	8231885	May 24, 2025	DP			
	8410102	May 24, 2025		U-1428		
	8426410	May 24, 2025		U-1428		
	8513247	Mar 25, 2031	DP U-1428			
	8513249	Mar 25, 2031	DP U-1428			
	8859551	May 25, 2024		U-1428		
	9861631	Mar 25, 2031		U-1428		
	9861632	Mar 25, 2031		U-1428		
<u>BRIMONIDINE TARTRATE - LUMIFY</u>						
N 208144	001 11596600	Jul 27, 2029		U-2222		
	11833245	Jul 27, 2029		U-2222		
	8293742	Jul 14, 2030		U-2222		
	9259425	Jul 14, 2030		U-2222		
<u>BRIMONIDINE TARTRATE; BRINZOLAMIDE - SIMBRINZA</u>						
N 204251	001 9044484	Oct 30, 2030	DP			
	9421265	Jun 17, 2030	DP			
<u>BRINCIDOFIVIR - TEMBEXA</u>						
N 214460	001 9303051	Aug 31, 2031	DS DP U-3165		NP	Jun 04, 2024
					ODE-354	Jun 04, 2028
<u>BRINCIDOFIVIR - TEMBEXA</u>						
N 214461	001 10112909	Oct 10, 2034		U-3165	NP	Jun 04, 2024
	10487061	Oct 10, 2034	DP U-3165		ODE-354	Jun 04, 2028
	8962829	Oct 10, 2034	DS DP			
	9303051	Aug 31, 2031	DS DP U-3165			
	9371344	Oct 10, 2034	DP			

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<u>BRINZOLAMIDE - BRINZOLAMIDE</u>						
A 211914	001				CGT	Apr 07, 2024
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	001	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	002	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	003	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	004	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205836	005	10729653	Apr 09, 2030	DP	NPP	Aug 27, 2024
	6911461	Feb 21, 2026	DS DP U-2295			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205837	001	6911461	Feb 21, 2026	DS DP U-1815	NPP	Aug 27, 2024
	6911461	Feb 21, 2026	DS DP U-2130			
<u>BRIVARACETAM - BRIVIACT</u>						
N 205838	001	6911461	Feb 21, 2026	DS DP U-2295	NPP	Aug 27, 2024
<u>BROMFENAC SODIUM - PROLENSA</u>						
N 203168	001	10085958	Nov 19, 2032	DP		
	8129431	Sep 11, 2025	DS DP			
	8669290	Jan 16, 2024	DP			
	8754131	Jan 16, 2024	DP			
	8871813	Jan 16, 2024	DP			
	8927606	Jan 16, 2024	U-100			
	8927606	Jan 16, 2024	U-1095			
	8927606	Jan 16, 2024	U-810			
	9144609	Jan 16, 2024	DP			
	9517220	Nov 11, 2033	U-1933			
	9561277	Jan 16, 2024	U-1933			
<u>BROMFENAC SODIUM - BROMSITE</u>						
N 206911	001	8778999	Aug 07, 2029	DP U-1834		
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866	001	10688094	Apr 30, 2032	U-2870		
	10688094	Apr 30, 2032	U-2871			
	10688094	Apr 30, 2032	U-2872			
	10688094	Apr 30, 2032	U-2873			
	10688094	Apr 30, 2032	U-2874			
	10688094	Apr 30, 2032	U-2875			
	10688094	Apr 30, 2032	U-2876			
	10688094	Apr 30, 2032	U-2877			
	10688094	Apr 30, 2032	U-2878			
	10688094	Apr 30, 2032	U-2879			

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<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	10688094	Apr 30, 2032	U-2880			
	10688094	Apr 30, 2032	U-2881			
	10688094	Apr 30, 2032	U-2882			
	10688094	Apr 30, 2032	U-2883			
	10688094	Apr 30, 2032	U-2884			
	10688094	Apr 30, 2032	U-2885			
	10688094	Apr 30, 2032	U-2886			
	10688094	Apr 30, 2032	U-2887			
	10688094	Apr 30, 2032	U-2888			
	10688155	Jun 07, 2030	U-2281			
	10688155	Jun 07, 2030	U-2890			
	10688155	Jun 07, 2030	U-2891			
	10688155	Jun 07, 2030	U-2892			
	10688155	Jun 07, 2030	U-2893			
	10688155	Jun 07, 2030	U-2894			
	10688155	Jun 07, 2030	U-2895			
	10688155	Jun 07, 2030	U-2896			
	10688155	Jun 07, 2030	U-2897			
	10688155	Jun 07, 2030	U-2898			
	10688155	Jun 07, 2030	U-2899			
	10688155	Jun 07, 2030	U-2900			
	10688155	Jun 07, 2030	U-2901			
	10688155	Jun 07, 2030	U-2902			
	10688155	Jun 07, 2030	U-2903			
	10688155	Jun 07, 2030	U-2904			
	10688155	Jun 07, 2030	U-2905			
	10688155	Jun 07, 2030	U-2906			
	10688155	Jun 07, 2030	U-2907			
	10688155	Jun 07, 2030	U-2908			
	10688155	Jun 07, 2030	U-2909			
	10688155	Jun 07, 2030	U-2910			
	10688155	Jun 07, 2030	U-2911			
	10688155	Jun 07, 2030	U-2912			
	10688155	Jun 07, 2030	U-2913			
	10688155	Jun 07, 2030	U-2914			
	10688155	Jun 07, 2030	U-2915			
	10688155	Jun 07, 2030	U-2916			
	10688155	Jun 07, 2030	U-2917			
	10688155	Jun 07, 2030	U-2918			
	10688155	Jun 07, 2030	U-2919			
	10688155	Jun 07, 2030	U-2920			
	10688155	Jun 07, 2030	U-2921			
	10688155	Jun 07, 2030	U-2922			
	10688155	Jun 07, 2030	U-2923			
	10688155	Jun 07, 2030	U-2924			
	10688155	Jun 07, 2030	U-2925			
	10688155	Jun 07, 2030	U-2926			
	10688155	Jun 07, 2030	U-2927			
	10688155	Jun 07, 2030	U-2928			

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<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866 001	10688155	Jun 07, 2030	U-2929			
	10688155	Jun 07, 2030	U-2930			
	10688155	Jun 07, 2030	U-2931			
	10688155	Jun 07, 2030	U-2932			
	10688155	Jun 07, 2030	U-2933			
	10688155	Jun 07, 2030	U-2934			
	10688155	Jun 07, 2030	U-2935			
	10688155	Jun 07, 2030	U-2936			
	10688155	Jun 07, 2030	U-2937			
	11000522	Apr 30, 2032	U-3119			
	11000522	Apr 30, 2032	U-3120			
	11000522	Apr 30, 2032	U-3121			
	11000522	Apr 30, 2032	U-3122			
	8431155	Apr 30, 2032	DP U-976			
	8613947	Apr 30, 2032	DP U-976			
	8877708	Jun 07, 2030	DP U-1706			
	9192576	Apr 30, 2032	DP U-976			
	9352025	Jun 07, 2030	U-2111			
	9352025	Jun 07, 2030	U-2112			
	9352025	Jun 07, 2030	U-2113			
	9352025	Jun 07, 2030	U-2114			
	9352025	Jun 07, 2030	U-2115			
	9352025	Jun 07, 2030	U-2116			
	9352025	Jun 07, 2030	U-2117			
	9352025	Jun 07, 2030	U-2118			
	9352025	Jun 07, 2030	U-2119			
	9522117	Apr 30, 2032	DP U-1939			
	9522117	Apr 30, 2032	DP U-976			
	9700555	Apr 30, 2032	DP U-2183			
	9700555	Apr 30, 2032	DP U-2184			
	9700555	Apr 30, 2032	DP U-2185			
	9700555	Apr 30, 2032	DP U-2186			
	9700555	Apr 30, 2032	DP U-2187			
	9700555	Apr 30, 2032	DP U-2188			
	9700555	Apr 30, 2032	DP U-2189			
	9700555	Apr 30, 2032	DP U-2190			
	9700555	Apr 30, 2032	DP U-2191			
	9700555	Apr 30, 2032	DP U-2192			
	9700555	Apr 30, 2032	DP U-2193			
	9700555	Apr 30, 2032	DP U-2194			
	9700555	Apr 30, 2032	DP U-2195			
	9700555	Apr 30, 2032	DP U-2196			
	9700555	Apr 30, 2032	DP U-2197			
	9700555	Apr 30, 2032	DP U-2198			
	9895422	Jun 07, 2030	U-2114			
	9895422	Jun 07, 2030	U-2116			
	9895422	Jun 07, 2030	U-2281			
	9895422	Jun 07, 2030	U-2282			
	9895422	Jun 07, 2030	U-2283			

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<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
N 020866	001	9895422	Jun 07, 2030	U-2284		
		9895422	Jun 07, 2030	U-2285		
		9895422	Jun 07, 2030	U-2286		
		9895422	Jun 07, 2030	U-2287		
		9993474	Apr 30, 2032	U-2384		
		9993474	Apr 30, 2032	U-2385		
		9993474	Apr 30, 2032	U-2386		
		9993474	Apr 30, 2032	U-2387		
		9993474	Apr 30, 2032	U-2388		
		9993474	Apr 30, 2032	U-2389		
		9993474	Apr 30, 2032	U-2390		
		9993474	Apr 30, 2032	U-2391		
		9993474	Apr 30, 2032	U-2392		
		9993474	Apr 30, 2032	U-2393		
<u>BUDESONIDE - UCERIS</u>						
N 203634	001	10307375	Sep 07, 2031	DP		
		10660858	Sep 07, 2031	DP		
		8895064	Sep 07, 2031	DP		
		9132093	Sep 07, 2031	DP		
		9192581	Sep 07, 2031	DP	U-1325	
<u>BUDESONIDE - ORTIKOS</u>						
N 211929	001	10172802	Sep 09, 2036	U-2554		
		9707182	Sep 09, 2036	DP	U-2554	
<u>BUDESONIDE - ORTIKOS</u>						
N 211929	002	10172802	Sep 09, 2036	U-2554		
		9707182	Sep 09, 2036	DP	U-2554	
<u>BUDESONIDE - TARPEYO</u>						
N 215935	001	8491932	May 07, 2029	DP	U-3269	
						NP Dec 15, 2024
					ODE-389	Dec 15, 2028
<u>BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT AEROSPHERE</u>						
N 216579	001	10716753	May 28, 2030	DP	U-3203	
		9415009	May 28, 2030		U-3203	
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929	001	7587988	Apr 10, 2026	DP		
		7587988*PED	Oct 10, 2026			
		8387615	Mar 26, 2027	DP		
		8387615*PED	Sep 26, 2027			
		8528545	Oct 16, 2028	DP		
		8528545*PED	Apr 16, 2029			
		8616196	Apr 07, 2029	DP		
		8616196*PED	Oct 07, 2029			
		8875699	Nov 10, 2024	DP		
		8875699*PED	May 10, 2025			
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929	002	7587988	Apr 10, 2026	DP		
		7587988*PED	Oct 10, 2026			
		8387615	Mar 26, 2027	DP		

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<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N 021929 002	8387615*PED	Sep 26, 2027				
	8528545	Oct 16, 2028	DP			
	8528545*PED	Apr 16, 2029				
	8616196	Apr 07, 2029	DP			
	8616196*PED	Oct 07, 2029				
	8875699	Nov 10, 2024	DP			
	8875699*PED	May 10, 2025				
<u>BUDESONIDE; FORMOTEROL FUMARATE; GLYCOPYRROLATE - BREZTRI AEROSPHERE</u>						
N 212122 001	10716753	May 28, 2030	DP U-2889			
	11331442	Oct 05, 2038	DP			
	11833292	Oct 05, 2038	DP			
	8324266	May 28, 2030		U-2889		
	8703806	May 28, 2030		U-2889		
	8808713	May 28, 2030	DP U-2889			
	8815258	Mar 17, 2031		U-2889		
	9415009	May 28, 2030		U-2889		
	9463161	May 28, 2030	DP U-2889			
<u>BUPIVACAINE - EXPAREL</u>						
N 022496 001	11033495	Jan 22, 2041	DP U-3182		NPP	Mar 22, 2024
	11179336	Jan 22, 2041	DP U-3250			
	11278494	Jan 22, 2041	DP U-3250			
	11304904	Jan 22, 2041	DP U-3346			
	11311486	Jan 22, 2041	DP U-3250			
	11357727	Jan 22, 2041	DP U-3380			
	11426348	Jan 22, 2041	DP U-3380			
	11452691	Jan 22, 2041	DP U-3439			
	11819574	Jan 22, 2041	DP U-3250			
	11819575	Jan 22, 2041	DP U-3250			
<u>BUPIVACAINE - EXPAREL</u>						
N 022496 002	11033495	Jan 22, 2041	DP U-3182		NPP	Mar 22, 2024
	11179336	Jan 22, 2041	DP U-3250			
	11278494	Jan 22, 2041	DP U-3250			
	11304904	Jan 22, 2041	DP U-3346			
	11311486	Jan 22, 2041	DP U-3250			
	11357727	Jan 22, 2041	DP U-3380			
	11426348	Jan 22, 2041	DP U-3380			
	11452691	Jan 22, 2041	DP U-3439			
	11819574	Jan 22, 2041	DP U-3250			
	11819575	Jan 22, 2041	DP U-3250			
<u>BUPIVACAINE - POSIMIR</u>						
N 204803 001	11400019	Jan 12, 2041	DP		NP	Feb 01, 2024
	11771624	Jan 12, 2041		U-3724		
	8153149	Sep 15, 2025	DP			
	8153661	Sep 15, 2025		U-3074		
	8753665	Sep 15, 2025	DP U-3074			
	8846072	Sep 15, 2025	DP U-3074			

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<u>BUPIVACAINE HYDROCHLORIDE - XARACOLL</u>						
N 209511	001 11746141	Jan 09, 2033	DP			
	RE47826	May 20, 2029	U-2949			
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	001 10098957	Apr 20, 2035	U-3118		NP	May 12, 2024
	10213510	Apr 20, 2035	DP U-3118			
	10398686	Mar 13, 2034	DP			
	10632199	Apr 20, 2035	DP U-3118			
	10898575	Apr 20, 2035	DP U-3118			
	10980886	Apr 20, 2035	DP			
	11083730	Apr 20, 2035	DP U-3118			
	11083797	Apr 20, 2035	DP U-3118			
	11253504	Mar 13, 2034	U-3118			
	11413350	Apr 20, 2035	U-3417			
	11844837	Apr 21, 2036	U-3417			
	9592227	Mar 13, 2034	DP U-3118			
	9694079	Apr 20, 2035	DP U-3118			
	9744163	Mar 13, 2034	DP			
	9801945	Apr 20, 2035	DP U-3118			
	9913909	Mar 13, 2034	U-3118			
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	002 10098957	Apr 20, 2035	U-3118		NP	May 12, 2024
	10213510	Apr 20, 2035	DP U-3118			
	10398686	Mar 13, 2034	DP			
	10632199	Apr 20, 2035	DP U-3118			
	10898575	Apr 20, 2035	DP U-3118			
	10980886	Apr 20, 2035	DP			
	11083730	Apr 20, 2035	DP U-3118			
	11083797	Apr 20, 2035	DP U-3118			
	11253504	Mar 13, 2034	U-3118			
	11413350	Apr 20, 2035	U-3417			
	11844837	Apr 21, 2036	U-3417			
	9592227	Mar 13, 2034	DP U-3118			
	9694079	Apr 20, 2035	DP U-3118			
	9744163	Mar 13, 2034	DP			
	9801945	Apr 20, 2035	DP U-3118			
	9913909	Mar 13, 2034	U-3118			
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	003 10098957	Apr 20, 2035	U-3118		NP	May 12, 2024
	10213510	Apr 20, 2035	DP U-3118			
	10398686	Mar 13, 2034	DP			
	10632199	Apr 20, 2035	DP U-3118			
	10898575	Apr 20, 2035	DP U-3118			
	10980886	Apr 20, 2035	DP			
	11083730	Apr 20, 2035	DP U-3118			
	11083797	Apr 20, 2035	DP U-3118			
	11253504	Mar 13, 2034	U-3118			
	11413350	Apr 20, 2035	U-3417			
	11844837	Apr 21, 2036	U-3417			

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<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	003	9592227	Mar 13, 2034	DP U-3118		
		9694079	Apr 20, 2035	DP U-3118		
		9744163	Mar 13, 2034	DP		
		9801945	Apr 20, 2035	DP U-3118		
		9913909	Mar 13, 2034	U-3118		
<u>BUPIVACAINE; MELOXICAM - ZYNRELEF KIT</u>						
N 211988	004	10098957	Apr 20, 2035	U-3118	NP	May 12, 2024
		10213510	Apr 20, 2035	DP U-3118		
		10398686	Mar 13, 2034	DP		
		10632199	Apr 20, 2035	DP U-3118		
		10898575	Apr 20, 2035	DP U-3118		
		10980886	Apr 20, 2035	DP		
		11083730	Apr 20, 2035	DP U-3118		
		11083797	Apr 20, 2035	DP U-3118		
		11253504	Mar 13, 2034	U-3118		
		11413350	Apr 20, 2035	U-3417		
		11844837	Apr 21, 2036	U-3417		
		9592227	Mar 13, 2034	DP U-3118		
		9694079	Apr 20, 2035	DP U-3118		
		9744163	Mar 13, 2034	DP		
		9801945	Apr 20, 2035	DP U-3118		
		9913909	Mar 13, 2034	U-3118		
<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819	001	10198218	Jun 06, 2031	U-2489		
		10558394	Jun 25, 2031	DP		
		10592168	Jun 06, 2031	U-2489		
		11000520	Nov 06, 2035	U-3111		
		11839611	Nov 06, 2035	U-3111		
		8921387	Jan 06, 2032	DP U-2173		
		8921387	Jan 06, 2032	DP U-2174		
		8975270	Sep 05, 2031	DP U-2175		
		8975270	Sep 05, 2031	DP U-2206		
		9272044	Jun 06, 2031	U-2176		
		9272044	Jun 06, 2031	U-2177		
		9272044	Jun 06, 2031	U-2178		
		9272044	Jun 06, 2031	U-2209		
		9498432	Jun 06, 2031	DP U-2179		
		9782402	Jun 06, 2031	DP U-2176		
		9782402	Jun 06, 2031	DP U-2180		
		9782402	Jun 06, 2031	DP U-2207		
		9782402	Jun 06, 2031	DP U-2208		
		9827241	Jun 06, 2031	DP U-2174		
		9827241	Jun 06, 2031	DP U-2181		
		9827241	Jun 06, 2031	DP U-2206		
		9827241	Jun 06, 2031	DP U-2210		
		9827241	Jun 06, 2031	DP U-2211		
<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819	002	10198218	Jun 06, 2031	U-2489		

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<u>BUPRENORPHINE - SUBLOCADE</u>						
N 209819 002	10558394	Jun 25, 2031	DP			
	10592168	Jun 06, 2031	U-2489			
	10646484	Jun 22, 2038	U-2489			
	11000520	Nov 06, 2035	U-3111			
	11839611	Nov 06, 2035	U-3111			
	8921387	Jan 06, 2032	DP U-2173			
	8921387	Jan 06, 2032	DP U-2174			
	8975270	Sep 05, 2031	DP U-2175			
	8975270	Sep 05, 2031	DP U-2206			
	9272044	Jun 06, 2031	U-2176			
	9272044	Jun 06, 2031	U-2177			
	9272044	Jun 06, 2031	U-2178			
	9272044	Jun 06, 2031	U-2209			
	9498432	Jun 06, 2031	DP U-2179			
	9782402	Jun 06, 2031	DP U-2176			
	9782402	Jun 06, 2031	DP U-2180			
	9782402	Jun 06, 2031	DP U-2207			
	9782402	Jun 06, 2031	DP U-2208			
	9827241	Jun 06, 2031	DP U-2174			
	9827241	Jun 06, 2031	DP U-2181			
	9827241	Jun 06, 2031	DP U-2206			
	9827241	Jun 06, 2031	DP U-2210			
	9827241	Jun 06, 2031	DP U-2211			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 001	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 002	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 003	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			

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<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 004	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 005	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 006	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE - BRIXADI</u>						
N 210136 007	10912772	Jul 26, 2032	U-3617		NP	May 23, 2026
	11110084	Jul 26, 2032	DP U-3616			
	11135215	Jul 26, 2032	DP			
	8236292	Jan 10, 2027	DP U-3619			
	8236755	Jul 31, 2026	DP U-3620			
	8545832	Jun 06, 2025	DP U-3619			
	9937164	Jul 26, 2032	DP U-3618			
<u>BUPRENORPHINE HYDROCHLORIDE - PROBUPHINE</u>						
N 204442 001	7736665	Apr 25, 2024	U-1878			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 001	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 002	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932 003	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
	9901539	Dec 21, 2032	U-1556			

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<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	004	8147866	Jul 23, 2027	DP U-1769		
		9655843	Jul 23, 2027	DP U-1556		
		9901539	Dec 21, 2032	U-1556		
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	005	8147866	Jul 23, 2027	DP U-1769		
		9655843	Jul 23, 2027	DP U-1556		
		9901539	Dec 21, 2032	U-1556		
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	006	8147866	Jul 23, 2027	DP U-1769		
		9655843	Jul 23, 2027	DP U-1556		
		9901539	Dec 21, 2032	U-1556		
<u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u>						
N 207932	007	8147866	Jul 23, 2027	DP U-1769		
		9655843	Jul 23, 2027	DP U-1556		
		9901539	Dec 21, 2032	U-1556		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	001	11135216	Aug 07, 2029	DP U-3111		
		8475832	Mar 26, 2030	DP U-1411		
		8603514	Apr 03, 2024	DP U-1464		
		9687454	Aug 07, 2029	DP U-1464		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	002	11135216	Aug 07, 2029	DP U-3111		
		8475832	Mar 26, 2030	DP U-1411		
		8603514	Apr 03, 2024	DP U-1464		
		9687454	Aug 07, 2029	DP U-1464		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	003	11135216	Aug 07, 2029	DP U-3111		
		8475832	Mar 26, 2030	DP U-1411		
		8603514	Apr 03, 2024	DP U-1464		
		9687454	Aug 07, 2029	DP U-1464		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u>						
N 022410	004	11135216	Aug 07, 2029	DP U-3111		
		8475832	Mar 26, 2030	DP U-1411		
		8603514	Apr 03, 2024	DP U-1464		
		9687454	Aug 07, 2029	DP U-1464		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242	001	10946010	Sep 18, 2032	DP		
		11020388	Sep 18, 2032	DP U-3131		
		11433066	Sep 18, 2032	U-3131		
		8470361	May 22, 2030	DP U-1425		
		8658198	Dec 03, 2027	DP U-1494		
		8940330	Sep 18, 2032	DP		
		9259421	Sep 18, 2032	DP		
		9439900	Sep 18, 2032	DP		

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<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 002	10874661	Sep 18, 2032	DP			
	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 003	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 004	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 005	10946010	Sep 18, 2032	DP			
	11020387	Sep 18, 2032	DP U-3131			
	11020388	Sep 18, 2032	DP U-3131			
	11433066	Sep 18, 2032	U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242 006	10946010	Sep 18, 2032	DP			
	11020388	Sep 18, 2032	DP U-3131			
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			

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<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u>						
N 204242	006	10946010	Sep 18, 2032	DP		
		11020388	Sep 18, 2032	DP U-3131		
		8470361	May 22, 2030	DP U-1425		
		8658198	Dec 03, 2027	DP U-1494		
		8940330	Sep 18, 2032	DP		
		9259421	Sep 18, 2032	DP		
		9439900	Sep 18, 2032	DP	Y	
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u>						
N 205637	001	8147866	Jul 23, 2027	DP U-1521		
		8703177	Aug 20, 2032	DP		
		9522188	Apr 24, 2035	DP		
		9655843	Jul 23, 2027	DP U-2017		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u>						
N 205637	002	8147866	Jul 23, 2027	DP U-1521		
		8703177	Aug 20, 2032	DP		
		9522188	Apr 24, 2035	DP		
		9655843	Jul 23, 2027	DP U-2017		
<u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u>						
N 205637	003	8147866	Jul 23, 2027	DP U-1521		
		8703177	Aug 20, 2032	DP		
		9522188	Apr 24, 2035	DP		
		9655843	Jul 23, 2027	DP U-2017		
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108	001	7241805	Jun 27, 2026	DP		
		7569610	Jun 27, 2026	U-997		
		7572935	Jun 27, 2026	DP		
		7585897	Jun 27, 2026	DP		
		7645802	Jun 27, 2026	DP		
		7649019	Jun 27, 2026	DP		
		7662407	Jun 27, 2026	DP		
		7671094	Jun 27, 2026	DP		
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108	002	7241805	Jun 27, 2026	DP		
		7569610	Jun 27, 2026	U-997		
		7572935	Jun 27, 2026	DP		
		7585897	Jun 27, 2026	DP		
		7645802	Jun 27, 2026	DP		
		7649019	Jun 27, 2026	DP		
		7662407	Jun 27, 2026	DP		
		7671094	Jun 27, 2026	DP		
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108	003	7241805	Jun 27, 2026	DP		
		7569610	Jun 27, 2026	U-997		
		7572935	Jun 27, 2026	DP		
		7585897	Jun 27, 2026	DP		
		7645802	Jun 27, 2026	DP		
		7649019	Jun 27, 2026	DP		

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<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N 022108	003 7662407	Jun 27, 2026	DP			
	7671094	Jun 27, 2026	DP			
<u>BUPROPION HYDROCHLORIDE - FORFIVO XL</u>						
N 022497	001 7674479	Jun 25, 2027	DP			
<u>BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE - AUVELITY</u>						
N 215430	001 10058518	Nov 05, 2034	U-3419		NP	Aug 18, 2025
	10064857	Nov 05, 2034	U-3419			
	10080727	Nov 05, 2034	U-3419			
	10092560	Nov 05, 2034	U-3419			
	10092561	Nov 05, 2034	U-3419			
	10105327	Nov 05, 2034	U-3419			
	10105361	Nov 05, 2034	U-3419			
	10251879	Nov 05, 2034	U-3419			
	10463634	Nov 05, 2034	U-3419			
	10512643	Nov 05, 2034	U-3419			
	10548857	Nov 05, 2034	U-3419			
	10596167	Nov 05, 2034	U-3419			
	10772850	Nov 05, 2034	U-3419			
	10780064	Jan 07, 2040	U-3419			
	10780066	Nov 09, 2034	U-3419			
	10786469	Nov 05, 2034	U-3419			
	10786496	Nov 05, 2034	U-3419			
	10799497	Nov 05, 2034	U-3419			
	10806710	Nov 05, 2034	U-3419			
	10864209	Nov 05, 2034	U-3419			
	10874663	Nov 05, 2034	U-3419			
	10874664	Nov 05, 2034	U-3419			
	10874665	Nov 05, 2034	U-3419			
	10881624	Nov 05, 2034	U-3419			
	10881657	Nov 05, 2034	U-3419			
	10894046	Nov 05, 2034	U-3419			
	10894047	Nov 05, 2034	U-3419			
	10898453	Nov 05, 2034	U-3419			
	10925842	Jan 07, 2040	U-3419			
	10933034	Nov 05, 2034	U-3419			
	10940124	Jan 07, 2040	U-3419			
	10945973	Nov 05, 2034	U-3419			
	10966941	Nov 05, 2034	U-3419			
	10966942	Jan 07, 2040	U-3419			
	10966974	Nov 05, 2034	U-3419			
	11020389	Nov 05, 2034	U-3419			
	11058648	Nov 05, 2034	U-3419			
	11090300	Nov 05, 2034	U-3419			
	11096937	Nov 05, 2034	U-3419			
	11123343	Nov 05, 2034	U-3419			
	11129826	Nov 05, 2034	U-3419			
	11141388	Nov 05, 2034	U-3419			
	11141416	Nov 05, 2034	U-3419			
	11147808	Nov 05, 2034	U-3419			

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<u>BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE - AUVELITY</u>						
N 215430 001	11185515	Nov 05, 2034	U-3419			
	11191739	Nov 05, 2034	DP U-3419			
	11197839	Nov 05, 2034	DP U-3419			
	11207281	Nov 05, 2034	U-3419			
	11213521	Nov 05, 2034	U-3419			
	11229640	Nov 05, 2034	U-3419			
	11234946	Nov 05, 2034	U-3419			
	11253491	Nov 05, 2034	U-3419			
	11253492	Nov 05, 2034	U-3419			
	11273133	Nov 05, 2034	U-3419			
	11273134	Nov 05, 2034	U-3419			
	11285118	Nov 05, 2034	U-3419			
	11285146	Nov 05, 2034	U-3419			
	11291638	Nov 05, 2034	U-3419			
	11291665	Nov 05, 2034	U-3419			
	11298351	Nov 05, 2034	U-3419			
	11298352	Nov 05, 2034	U-3419			
	11311534	Nov 05, 2034	U-3419			
	11344544	Nov 05, 2034	U-3419			
	11357744	Nov 05, 2034	U-3419			
	11364233	Nov 05, 2034	U-3419			
	11382874	Nov 05, 2034	U-3419			
	11419867	Nov 05, 2034	U-3419			
	11426370	Nov 05, 2034	U-3419			
	11426401	Nov 05, 2034	U-3419			
	11433067	Nov 05, 2034	DP U-3419			
	11439636	Nov 05, 2034	U-3419			
	11478468	Nov 05, 2034	U-3419			
	11497721	Nov 05, 2034	U-3419			
	11510918	Nov 05, 2034	U-3419			
	11517542	Nov 05, 2034	U-3419			
	11517543	Nov 05, 2034	U-3419			
	11524007	Nov 05, 2034	U-3419			
	11524008	Nov 05, 2034	U-3419			
	11534414	Nov 05, 2034	DP U-3419			
	11541021	Nov 05, 2034	U-3419			
	11541048	Nov 05, 2034	U-3419			
	11596627	Nov 05, 2034	U-3419			
	11617728	Nov 05, 2034	U-3419			
	11617747	Nov 05, 2034	U-3563			
	11717518	Jan 20, 2043	U-3419			
	11730706	Jan 23, 2043	U-3419			
	11752144	Feb 23, 2043	U-3419			
	11779579	Nov 05, 2034	U-3419			
	11839612	Mar 02, 2043	U-3419			
	11844797	Apr 20, 2043	U-3778			
	8569328	Oct 29, 2033	DP U-3419			
	9168234	Nov 05, 2034	U-3419			
	9198905	Nov 05, 2034	U-3419			

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<u>BUPROPION HYDROCHLORIDE; DEXTROMETHORPHAN HYDROBROMIDE - AUVELITY</u>						
N 215430	001	9205083	Nov 05, 2034			U-3419
		9238032	Nov 05, 2034			U-3419
		9278095	Nov 05, 2034			U-3419
		9314462	Nov 05, 2034			U-3419
		9370513	Nov 05, 2034			U-3419
		9375429	Nov 05, 2034			U-3419
		9408815	Nov 05, 2034			U-3419
		9421176	Nov 05, 2034			U-3419
		9457023	Nov 05, 2034			U-3419
		9457025	Nov 05, 2034			U-3419
		9474731	Nov 05, 2034			U-3419
		9486450	Nov 05, 2034			U-3419
		9700528	Nov 05, 2034			U-3419
		9700553	Nov 05, 2034			U-3419
		9707191	Nov 05, 2034			U-3419
		9763932	Nov 05, 2034			U-3419
		9861595	Nov 05, 2034			U-3419
		9867819	Nov 05, 2034			U-3419
		9968568	Nov 05, 2034			U-3419
<u>BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE - CONTRAVE</u>						
N 200063	001	10231964	Jul 02, 2034			U-1583
		10307376	Nov 08, 2027			U-1585
		10403170	Jun 05, 2033			U-1583
		10828294	Jul 02, 2034			U-1583
		10835527	Jul 02, 2034			U-1583
		11033543	Jan 10, 2031			U-1583
		11139056	Jun 05, 2033			U-1583
		11278544	Apr 21, 2024			U-1583
		11324741	May 29, 2029			U-1583
		7375111	Mar 26, 2025	DP		
		7462626	Jul 20, 2024			U-1583
		8088786	Feb 03, 2029	DP		
		8318788	Nov 08, 2027			U-1584
		8722085	Nov 08, 2027			U-1585
		8815889	Jul 20, 2024			U-1586
		8916195	Feb 02, 2030			U-1639
		9107837	Jun 04, 2027			U-1639
		9125868	Nov 08, 2027			U-1585
		9248123	Jan 13, 2032			U-1808
		9633575	Jun 25, 2033			U-1583
<u>CABAZITAXEL - JEVTANA KIT</u>						
N 201023	001	10583110	Oct 27, 2030			U-2753
		10716777	Oct 27, 2030			U-2856
		7241907	Dec 10, 2025	DS		
		7241907*PED	Jun 10, 2026			
		8927592	Oct 27, 2030			U-3200
		8927592*PED	Apr 27, 2031			

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<u>CABOTEGRAVIR - APRETUDE</u>						
N 215499 001	10927129	Apr 28, 2026	DS DP		NCE	Jan 21, 2026
	11224597	Sep 15, 2031	DP			
	8410103	Apr 28, 2026	DS DP			
<u>CABOTEGRAVIR SODIUM - VOCABRIA</u>						
N 212887 001	10927129	Apr 28, 2026	DS DP		M-273	Jan 31, 2025
	8410103	Apr 28, 2026	DS DP U-3061		NCE	Jan 21, 2026
	8410103	Apr 28, 2026	DS DP U-3348		NPP	Mar 29, 2025
<u>CABOTEGRAVIR; RILPIVIRINE - CABENUVA KIT</u>						
N 212888 001	10927129	Apr 28, 2026	DS DP		D-184	Jan 31, 2025
	11224597	Sep 15, 2031	DP U-3348		NCE	Jan 21, 2026
	11389447	Jun 30, 2027	U-3405		NPP	Mar 29, 2025
	7125879	Apr 21, 2025	DS DP U-3348			
	8410103	Apr 28, 2026	DS DP U-3348			
<u>CABOTEGRAVIR; RILPIVIRINE - CABENUVA KIT</u>						
N 212888 002	10927129	Apr 28, 2026	DS DP		D-184	Jan 31, 2025
	11224597	Sep 15, 2031	DP U-3348		NCE	Jan 21, 2026
	11389447	Jun 30, 2027	U-3405		NPP	Mar 29, 2025
	7125879	Apr 21, 2025	DS DP U-3348			
	8410103	Apr 28, 2026	DS DP U-3348			
<u>CABOZANTINIB S-MALATE - COMETRIO</u>						
N 203756 001	11091439	Jan 15, 2030	DS			
	11091440	Jan 15, 2030	DP			
	11098015	Jan 15, 2030	U-1617			
	11298349	Feb 10, 2032	DP			
	7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP U-1617			
	9717720	Feb 10, 2032	DP			
<u>CABOZANTINIB S-MALATE - COMETRIO</u>						
N 203756 002	11091439	Jan 15, 2030	DS			
	11091440	Jan 15, 2030	DP			
	11098015	Jan 15, 2030	U-1617			
	11298349	Feb 10, 2032	DP			
	7579473	Aug 14, 2026	DS DP			
	8877776	Oct 08, 2030	DS DP U-1617			
	9717720	Feb 10, 2032	DP			
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692 001	10034873	Jul 18, 2031	U-2488		I-854	Jan 22, 2024
	10039757	Jul 18, 2031	U-1480		I-873	Sep 17, 2024
	11091439	Jan 15, 2030	DS		ODE-227	Jan 14, 2026
	11091440	Jan 15, 2030	DP		ODE-375	Sep 17, 2028
	11098015	Jan 15, 2030	U-1220			
	11098015	Jan 15, 2030	U-1480			
	11098015	Jan 15, 2030	U-2488			
	11098015	Jan 15, 2030	U-3225			
	11298349	Feb 10, 2032	DP			
	7579473	Aug 14, 2026	DS DP			
	8497284	Sep 24, 2024	U-1220			

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<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	001	8497284	Sep 24, 2024	U-1480		
		8497284	Sep 24, 2024	U-2488		
		8877776	Oct 08, 2030	DS DP U-3225		
		9724342	Jul 09, 2033	DP		
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	002	10034873	Jul 18, 2031	U-2488	I-854	Jan 22, 2024
		10039757	Jul 18, 2031	U-1480	I-873	Sep 17, 2024
		11091439	Jan 15, 2030	DS	ODE-227	Jan 14, 2026
		11091440	Jan 15, 2030	DP	ODE-375	Sep 17, 2028
		11098015	Jan 15, 2030	U-1220		
		11098015	Jan 15, 2030	U-1480		
		11098015	Jan 15, 2030	U-2488		
		11098015	Jan 15, 2030	U-3225		
		11298349	Feb 10, 2032	DP		
		7579473	Aug 14, 2026	DS DP		
		8497284	Sep 24, 2024	U-1220		
		8497284	Sep 24, 2024	U-1480		
		8497284	Sep 24, 2024	U-2488		
		8877776	Oct 08, 2030	DS DP U-3225		
		9724342	Jul 09, 2033	DP		
<u>CABOZANTINIB S-MALATE - CABOMETYX</u>						
N 208692	003	10034873	Jul 18, 2031	U-2488	I-854	Jan 22, 2024
		10039757	Jul 18, 2031	U-1480	I-873	Sep 17, 2024
		11091439	Jan 15, 2030	DS	ODE-227	Jan 14, 2026
		11091440	Jan 15, 2030	DP	ODE-375	Sep 17, 2028
		11098015	Jan 15, 2030	U-1220		
		11098015	Jan 15, 2030	U-1480		
		11098015	Jan 15, 2030	U-2488		
		11098015	Jan 15, 2030	U-3225		
		11298349	Feb 10, 2032	DP		
		7579473	Aug 14, 2026	DS DP		
		8497284	Sep 24, 2024	U-1220		
		8497284	Sep 24, 2024	U-1480		
		8497284	Sep 24, 2024	U-2488		
		8877776	Oct 08, 2030	DS DP U-3225		
		9724342	Jul 09, 2033	DP		
<u>CALCIFEDIOL - RAYALDEE</u>						
N 208010	001	10213442	Feb 02, 2027	DP		
		10300078	Mar 14, 2034	DP		
		10357502	Mar 14, 2034	DP		
		11154509	Apr 25, 2028	U-3248		
		11253528	Mar 14, 2034	DP		
		11801253	Sep 07, 2030	DP U-3721		
		11801253	Sep 07, 2030	DP U-3722		
		11801253	Sep 07, 2030	DP U-3723		
		8207149	Apr 25, 2028	U-1871		
		8361488	Jul 19, 2028	DP		
		8426391	Aug 27, 2028	U-1872		

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<u>CALCIFEDIOL - RAYALDEE</u>						
N 208010	001 8778373	Apr 25, 2028	U-1873			
	8906410	Feb 02, 2027	DP			
	9408858	Apr 25, 2028	U-1888			
	9498486	Apr 25, 2028	U-1920			
	9861644	Mar 14, 2034	DP			
	9925147	Apr 25, 2028	DP U-2255			
	9925147	Apr 25, 2028	DP U-2256			
	9925147	Apr 25, 2028	DP U-2257			
	9925147	Apr 25, 2028	DP U-2258			
	9925147	Apr 25, 2028	DP U-2259			
	9943530	Feb 02, 2027	U-2274			
<u>CALCIPOTRIENE - SORILUX</u>						
N 022563	001 8263580	May 07, 2028	DP U-1280			
	8263580	May 07, 2028	DP U-2662			
	8629128	May 26, 2026	DP U-1280			
	8629128	May 26, 2026	DP U-1767			
	8629128	May 26, 2026	DP U-2662			
<u>CALCIUM ACETATE - PHOSLYRA</u>						
N 022581	001 8591938	Feb 23, 2030	DP U-1469			
	8592480	Jul 20, 2027	U-1469			
	9089528	Jul 20, 2027	U-1469			
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE</u>						
A 216541	001				CGT	Apr 30, 2024
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
A 217174	001				CGT	Mar 10, 2024
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
A 217174	002				CGT	Mar 10, 2024
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE</u>						
A 217689	001				CGT	Apr 17, 2024
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	001 10130646	Jul 25, 2037	DP			
	10342813	Jul 25, 2037	DP			
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	002 10130646	Jul 25, 2037	DP			
	10342813	Jul 25, 2037	DP			
<u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u>						
N 210906	003 10130646	Jul 25, 2037	DP			
	10342813	Jul 25, 2037	DP			
<u>CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE - XYWAV</u>						
N 212690	001 10195168	Jan 11, 2033	DP		I-870	Aug 12, 2024
	10213400	Mar 15, 2033	U-2499		ODE-361	Jul 21, 2027
	10213400*PED	Sep 15, 2033			ODE-369	Aug 12, 2028
	10675258	Jan 11, 2033	U-2938			
	10864181	Mar 15, 2033	U-3017			
	10864181*PED	Sep 15, 2033				
	11253494	Mar 15, 2033	U-3323			

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<u>CALCIUM OXYBATE; MAGNESIUM OXYBATE; POTASSIUM OXYBATE; SODIUM OXYBATE - XYWAV</u>						
N 212690 001	11253494	Mar 15, 2033	U-3324			
	11253494*PED	Sep 15, 2033				
	11426373	Sep 19, 2037	U-3432			
	11554102	Jan 11, 2033	DP			
	8591922	Jan 11, 2033	DP			
	8772306	Mar 15, 2033	U-1532			
	8772306	Mar 15, 2033	U-3198			
	8772306*PED	Sep 15, 2033				
	8901173	Jan 11, 2033	DP			
	9050302	Mar 15, 2033	U-1532			
	9050302*PED	Sep 15, 2033				
	9132107	Jan 11, 2033	DP			
	9486426	Mar 15, 2033	U-1532			
	9486426*PED	Sep 15, 2033				
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042 001	10617668	May 11, 2031	DP U-2441			
	10617668	May 11, 2031	DP U-2632			
	10617668	May 11, 2031	DP U-2794			
	10617668	May 11, 2031	DP U-2795			
	10617668	May 11, 2031	DP U-2796			
	10617668	May 11, 2031	DP U-2797			
	10617668	May 11, 2031	DP U-2798			
	10617668	May 11, 2031	DP U-2799			
	10617668	May 11, 2031	DP U-493			
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			
	8222219	Apr 11, 2025	U-2632			
	8222219	Apr 11, 2025	U-493			
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042 002	10617668	May 11, 2031	DP U-2441			
	10617668	May 11, 2031	DP U-2632			
	10617668	May 11, 2031	DP U-2794			
	10617668	May 11, 2031	DP U-2795			
	10617668	May 11, 2031	DP U-2796			
	10617668	May 11, 2031	DP U-2797			
	10617668	May 11, 2031	DP U-2798			
	10617668	May 11, 2031	DP U-2799			
	10617668	May 11, 2031	DP U-493			
	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025	U-2441			

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<u>CANAGLIFLOZIN - INVOKANA</u>						
N 204042	002	8222219	Apr 11, 2025		U-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		8513202	Dec 03, 2027	DS DP	U-493	
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353	001	11576894	Jul 06, 2030		DP	
		7943582	Feb 26, 2029	DS DP	U-2441	
		7943582	Feb 26, 2029	DS DP	U-2632	
		7943582	Feb 26, 2029	DS DP	U-493	
		7943788	Jul 14, 2027	DS DP		
		8222219	Apr 11, 2025		U-2441	
		8222219	Apr 11, 2025		U-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		8513202	Dec 03, 2027	DS DP	U-493	
		8785403	Jul 30, 2024		DP	
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353	002	11576894	Jul 06, 2030		DP	
		7943582	Feb 26, 2029	DS DP	U-2441	
		7943582	Feb 26, 2029	DS DP	U-2632	
		7943582	Feb 26, 2029	DS DP	U-493	
		7943788	Jul 14, 2027	DS DP		
		8222219	Apr 11, 2025		U-2441	
		8222219	Apr 11, 2025		U-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		8513202	Dec 03, 2027	DS DP	U-493	
		8785403	Jul 30, 2024		DP	
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353	003	11576894	Jul 06, 2030		DP	
		7943582	Feb 26, 2029	DS DP	U-2441	
		7943582	Feb 26, 2029	DS DP	U-2632	
		7943582	Feb 26, 2029	DS DP	U-493	
		7943788	Jul 14, 2027	DS DP		
		8222219	Apr 11, 2025		U-2441	
		8222219	Apr 11, 2025		U-2632	
		8222219	Apr 11, 2025		U-493	
		8513202	Dec 03, 2027	DS DP	U-2441	
		8513202	Dec 03, 2027	DS DP	U-2632	
		8513202	Dec 03, 2027	DS DP	U-493	
		8785403	Jul 30, 2024		DP	
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353	004	11576894	Jul 06, 2030		DP	
		7943582	Feb 26, 2029	DS DP	U-2441	
		7943582	Feb 26, 2029	DS DP	U-2632	

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<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u>						
N 204353 004	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 001	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 002	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 003	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			
	7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879 004	7943582	Feb 26, 2029	DS DP U-2441			
	7943582	Feb 26, 2029	DS DP U-2632			

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<u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u>						
N 205879	004 7943582	Feb 26, 2029	DS DP U-493			
	7943788	Jul 14, 2027	DS DP			
	8222219	Apr 11, 2025		U-2441		
	8222219	Apr 11, 2025		U-2632		
	8222219	Apr 11, 2025		U-493		
	8513202	Dec 03, 2027	DS DP U-2441			
	8513202	Dec 03, 2027	DS DP U-2632			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<u>CANGRELOR - KENGREAL</u>						
N 204958	001 10039780	Jul 10, 2035		U-2260		
	8680052	Mar 09, 2033		U-2979		
	9295687	Jul 10, 2035	DP			
	9427448	Nov 10, 2030		U-1926		
	9439921	Jul 10, 2035	DP			
	9700575	Jul 10, 2035	DP			
	9925265	May 13, 2029		U-2260		
<u>CANNABIDIOL - EPIDIOLEX</u>						
N 210365	001 10092525	Jun 17, 2035	U-2427		M-270	Oct 20, 2026
	10111840	Jun 17, 2035	U-2442		ODE-216	Sep 28, 2025
	10111840	Jun 17, 2035	U-2443		ODE-326	Jul 31, 2027
	10137095	Jun 17, 2035	U-2454		ODE-332	Jul 31, 2027
	10137095	Jun 17, 2035	U-2455			
	10603288	Jun 17, 2035	U-2780			
	10603288	Jun 17, 2035	U-2781			
	10603288	Jun 17, 2035	U-2782			
	10603288	Jun 17, 2035	U-2783			
	10709671	Jun 17, 2035	U-2862			
	10709673	Jun 17, 2035	DP			
	10709674	Jun 17, 2035	U-2780			
	10709674	Jun 17, 2035	U-2781			
	10849860	Jun 17, 2035	U-2427			
	10849860	Jun 17, 2035	U-2454			
	10918608	Oct 13, 2035	U-3071			
	10918608	Oct 13, 2035	U-3072			
	10918608	Oct 13, 2035	U-3073			
	10966939	Jun 17, 2035	DP U-2780			
	10966939	Jun 17, 2035	DP U-2781			
	11065209	Oct 13, 2035	U-3071			
	11096905	Oct 13, 2035	DS DP U-2780			
	11096905	Oct 13, 2035	DS DP U-2781			
	11154516	Jun 17, 2035	U-3235			
	11154516	Jun 17, 2035	U-3236			
	11160795	Mar 01, 2041	U-3233			
	11207292	Apr 26, 2039	DS U-3235			
	11207292	Apr 26, 2039	DS U-3236			
	11207292	Apr 26, 2039	DS U-3277			
	11311498	Jun 17, 2035	U-3375			
	11311498	Jun 17, 2035	U-3376			

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<u>CANNABIDIOL - EPIDIOLEX</u>						
N 210365 001	11357741	Jun 17, 2035	U-2862			
	11400055	Oct 13, 2035	U-3071			
	11406623	Mar 01, 2041	U-3233			
	11446258	Jun 17, 2035	U-2780			
	11446258	Jun 17, 2035	U-2781			
	11446258	Jun 17, 2035	U-3071			
	11633369	Jun 17, 2035	DP U-2780			
	11633369	Jun 17, 2035	DP U-2781			
	11633369	Jun 17, 2035	DP U-3071			
	11701330	Jun 17, 2035	U-2780			
	11701330	Jun 17, 2035	U-2781			
	11766411	Jun 17, 2035	U-2781			
	9949937	Jun 17, 2035	U-2421			
	9956183	Jun 17, 2035	U-2422			
	9956183	Jun 17, 2035	U-2423			
	9956184	Jun 17, 2035	U-2424			
	9956185	Jun 17, 2035	U-2425			
	9956186	Jun 17, 2035	U-2426			
<u>CANTHARIDIN - YCANTH</u>						
N 212905 001	11052064	May 28, 2035	DP U-3663			
	11052064	May 28, 2035	DP U-3665			
	11147790	Aug 22, 2038	DP U-3663			
	11147790	Aug 22, 2038	DP U-3664			
<u>CAPIVASERTIB - TRUOAP</u>						
N 218197 001	10039766	Apr 16, 2033	U-3762		NCE	Nov 16, 2028
	10059714	Oct 10, 2028	DS DP			
	10654855	Oct 10, 2028	U-3762			
	11760760	Oct 10, 2028	U-3762			
	8101623	Mar 10, 2030	DS DP U-3762			
	8809336	Oct 25, 2025	U-3762			
	9006430	Oct 25, 2025	DP			
	9487525	Apr 16, 2033	DS DP			
<u>CAPIVASERTIB - TRUOAP</u>						
N 218197 002	10039766	Apr 16, 2033	U-3762		NCE	Nov 16, 2028
	10059714	Oct 10, 2028	DS DP			
	10654855	Oct 10, 2028	U-3762			
	11760760	Oct 10, 2028	U-3762			
	8101623	Mar 10, 2030	DS DP U-3762			
	8809336	Oct 25, 2025	U-3762			
	9006430	Oct 25, 2025	DP			
	9487525	Apr 16, 2033	DS DP			
<u>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591 001	10596178	Jul 22, 2035	DS DP		NCE	May 06, 2025
	7767675	Nov 19, 2027	DS DP		ODE-291	May 06, 2027
	8420645	Jun 05, 2031	DS DP			
	8461330	Nov 19, 2027	DS DP			
	8901123	May 20, 2029	U-2813			

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<u>CAPMATINIB HYDROCHLORIDE - TABRECTA</u>						
N 213591 002	10596178	Jul 22, 2035	DS DP		NCE	May 06, 2025
	7767675	Nov 19, 2027	DS DP		ODE-291	May 06, 2027
	8420645	Jun 05, 2031	DS DP			
	8461330	Nov 19, 2027	DS DP			
	8901123	May 20, 2029		U-2813		
<u>CAPSAICIN - OUTENZA</u>						
N 022395 001	10034841	Sep 06, 2025	DP			
	8821920	Mar 26, 2030	DP			
	9226903	Dec 15, 2028	DP			
<u>CARBAMAZEPINE - EQUETRO</u>						
N 021710 001	6977253	May 19, 2024		U-693		
<u>CARBAMAZEPINE - EQUETRO</u>						
N 021710 002	6977253	May 19, 2024		U-693		
<u>CARBAMAZEPINE - EQUETRO</u>						
N 021710 003	6977253	May 19, 2024		U-693		
<u>CARBAMAZEPINE - CARNEXIV</u>						
N 206030 001	11529357	Jan 31, 2040	DP			
	7635773	Mar 13, 2029	DP			
	8410077	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
	9629797	Nov 10, 2028		U-2004		
	9629797	Nov 10, 2028		U-2005		
	9629797	Nov 10, 2028		U-2006		
	9750822	Mar 13, 2029	DP			
	9770407	Nov 10, 2028	DP			
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312 001	8377474	Dec 26, 2028	DP	U-1645		
	8377474	Dec 26, 2028	DP	U-219		
	8454998	Dec 26, 2028	DP	U-1645		
	8454998	Dec 26, 2028	DP	U-1646		
	8454998	Dec 26, 2028	DP	U-1647		
	8454998	Dec 26, 2028	DP	U-1649		
	8454998	Dec 26, 2028	DP	U-219		
	8557283	Dec 26, 2028	DP	U-1645		
	8557283	Dec 26, 2028	DP	U-219		
	9089607	Dec 26, 2028	DP	U-1645		
	9089607	Dec 26, 2028	DP	U-1720		
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP	U-219		
	9533046	Dec 26, 2028	DP	U-219		
	9901640	Dec 26, 2028	DP	U-219		
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312 002	8377474	Dec 26, 2028	DP	U-1645		
	8377474	Dec 26, 2028	DP	U-219		
	8454998	Dec 26, 2028	DP	U-1645		
	8454998	Dec 26, 2028	DP	U-1646		
	8454998	Dec 26, 2028	DP	U-1647		

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<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312 002	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
	9901640	Dec 26, 2028	DP U-219			
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312 003	8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
	9901640	Dec 26, 2028	DP U-219			
<u>CARBIDOPA; LEVODOPA - RYTARY</u>						
N 203312 004	8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
	9901640	Dec 26, 2028	DP U-219			
<u>CARBIDOPA; LEVODOPA - DHIVY</u>						
N 214869 001	11033521	Mar 28, 2039	DP U-219			
	11033521	Mar 28, 2039	DP U-3304			
	11033521	Mar 28, 2039	DP U-3305			
	11439613	Mar 28, 2039	U-3557			
	11819485	Mar 28, 2039	U-3557			

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<u>CARBINOXAMINE MALEATE - KARBINAL ER</u>						
N 022556	001 8062667	Mar 29, 2029	DP			
	9522191	Jun 15, 2027	DP			
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714	001 7232818	Apr 14, 2025	DS DP			
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025		U-1260		
	7491704	Apr 14, 2025		U-2319		
	7491704	Apr 14, 2025		U-2320		
	7491704	Apr 14, 2025		U-2947		
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025		U-1260		
	8129346	Apr 14, 2025		U-2319		
	8129346	Apr 14, 2025		U-2320		
	8129346	Apr 14, 2025		U-2947		
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025		U-1260		
	8207127	Apr 14, 2025		U-2319		
	8207127	Apr 14, 2025		U-2320		
	8207127	Apr 14, 2025		U-2947		
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	RE47954	Oct 21, 2029		U-3449		
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714	002 7232818	Apr 14, 2025	DS DP			
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025		U-1260		
	7491704	Apr 14, 2025		U-2319		
	7491704	Apr 14, 2025		U-2320		
	7491704	Apr 14, 2025		U-2947		
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025		U-1260		
	8129346	Apr 14, 2025		U-2319		
	8129346	Apr 14, 2025		U-2320		
	8129346	Apr 14, 2025		U-2947		
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025		U-1260		
	8207127	Apr 14, 2025		U-2319		
	8207127	Apr 14, 2025		U-2320		
	8207127	Apr 14, 2025		U-2947		
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	RE47954	Oct 21, 2029		U-3449		
<u>CARFILZOMIB - KYPROLIS</u>						
N 202714	003 7232818	Apr 14, 2025	DS DP			
	7417042	Jul 20, 2026	DS DP			
	7491704	Apr 14, 2025		U-2319		

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<u>CARFILZOMIB - KYPROLIS</u>						
N 202714 003	7491704	Apr 14, 2025	U-2320			
	7491704	Apr 14, 2025	U-2947			
	7737112	Dec 07, 2027	DP			
	8129346	Apr 14, 2025	U-2319			
	8129346	Apr 14, 2025	U-2320			
	8129346	Apr 14, 2025	U-2947			
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025	U-2319			
	8207127	Apr 14, 2025	U-2320			
	8207127	Apr 14, 2025	U-2947			
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	RE47954	Oct 21, 2029	U-3449			
<u>CARGLUMIC ACID - CARBAGLU</u>						
N 022562 001					ODE-345	Jan 22, 2028
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 001	7737142	Sep 17, 2029	DS DP U-1750		I-904	Dec 16, 2025
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7737142	Sep 17, 2029	DS DP U-2545			
	7737142	Sep 17, 2029	DS DP U-3503			
	7943621	Dec 16, 2028	DS DP			
	RE47350	Jul 16, 2029	U-1750			
	RE47350	Jul 16, 2029	U-2543			
	RE47350	Jul 16, 2029	U-2544			
	RE47350	Jul 16, 2029	U-2545			
	RE47350	Jul 16, 2029	U-3503			
	RE49110	Jul 16, 2029	U-2543			
	RE49110	Jul 16, 2029	U-2544			
	RE49110	Jul 16, 2029	U-2545			
	RE49110	Jul 16, 2029	U-3503			
	RE49302	Jul 16, 2029	U-2543			
	RE49302	Jul 16, 2029	U-2544			
	RE49302	Jul 16, 2029	U-2545			
	RE49302	Jul 16, 2029	U-3503			
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370 002	7737142	Sep 17, 2029	DS DP U-1750		I-904	Dec 16, 2025
	7737142	Sep 17, 2029	DS DP U-2543			
	7737142	Sep 17, 2029	DS DP U-2544			
	7737142	Sep 17, 2029	DS DP U-2545			
	7737142	Sep 17, 2029	DS DP U-3503			
	7943621	Dec 16, 2028	DS DP			
	RE49110	Jul 16, 2029	U-2543			
	RE49110	Jul 16, 2029	U-2544			
	RE49110	Jul 16, 2029	U-2545			
	RE49110	Jul 16, 2029	U-3503			
	RE49302	Jul 16, 2029	U-2543			

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<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370	002	RE49302	Jul 16, 2029	U-2544		
		RE49302	Jul 16, 2029	U-2545		
		RE49302	Jul 16, 2029	U-3503		
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370	003	7737142	Sep 17, 2029	DS DP U-1750	I-904	Dec 16, 2025
		7737142	Sep 17, 2029	DS DP U-2543		
		7737142	Sep 17, 2029	DS DP U-2544		
		7943621	Dec 16, 2028	DS DP		
		RE49110	Jul 16, 2029	U-2543		
		RE49110	Jul 16, 2029	U-2544		
		RE49110	Jul 16, 2029	U-2545		
		RE49302	Jul 16, 2029	U-2543		
		RE49302	Jul 16, 2029	U-2544		
		RE49302	Jul 16, 2029	U-2545		
		RE49302	Jul 16, 2029	U-3503		
<u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u>						
N 204370	004	7737142	Sep 17, 2029	DS DP U-1750	I-904	Dec 16, 2025
		7737142	Sep 17, 2029	DS DP U-2543		
		7737142	Sep 17, 2029	DS DP U-2544		
		7943621	Dec 16, 2028	DS DP		
		RE49110	Jul 16, 2029	U-2543		
		RE49110	Jul 16, 2029	U-2544		
		RE49110	Jul 16, 2029	U-2545		
		RE49302	Jul 16, 2029	U-2543		
		RE49302	Jul 16, 2029	U-2544		
		RE49302	Jul 16, 2029	U-2545		
		RE49302	Jul 16, 2029	U-3503		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	001	8101209	Sep 11, 2025	DP		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	002	8101209	Sep 11, 2025	DP		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	003	8101209	Sep 11, 2025	DP		
<u>CARVEDILOL PHOSPHATE - COREG CR</u>						
N 022012	004	8101209	Sep 11, 2025	DP		
<u>CASIMERSEN - AMONDYS 45</u>						
N 213026	001	10287586	Nov 12, 2030	DS DP	NCE	Feb 25, 2026
		10781450	Nov 12, 2030	U-3089	ODE-347	Feb 25, 2028
		9228187	Nov 12, 2030	DS DP		
		9447415	Jun 28, 2025	DS DP		
		9758783	Nov 12, 2030	U-3088		
		9758783	Nov 12, 2030	U-3089		
		RE48960	Jun 28, 2025	DS DP U-3087		
		RE48960	Jun 28, 2025	DS DP U-3088		
<u>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</u>						
N 206110	001	9636407	Dec 21, 2032	DP		

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<u>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</u>						
N 206110	002 9636407	Dec 21, 2032	DP			
<u>CEDAZURIDINE; DECITABINE - INQOVI</u>						
N 212576	001 8268800	Aug 22, 2030	DS U-2864		NCE	Jul 07, 2025
	8268800	Aug 22, 2030	DS U-2865		ODE-316	Jul 07, 2027
	8268800	Aug 22, 2030	DS U-2866			
	8268800	Aug 22, 2030	DS U-2867			
	8618075	Oct 16, 2028	U-2864			
	8618075	Oct 16, 2028	U-2867			
	9567363	Oct 16, 2028	DS			
<u>CEFIDEROCOL SULFATE TOSYLATE - FETROJA</u>						
N 209445	001 10004750	Sep 03, 2035	DS DP		NCE	Nov 14, 2024
	9238657	Nov 14, 2033	DS DP U-282		GAIN	Nov 14, 2029
	9238657	Nov 14, 2033	DS DP U-3470			
	9238657	Nov 14, 2033	DS DP U-3471			
	9949982	Sep 03, 2035	DP			
<u>CEFIXIME - SUPRAX</u>						
N 202091	001 9233112	Dec 14, 2028	DP U-1676			
<u>CEFTAROLINE FOSAMIL - TEFLARO</u>						
N 200327	001 8247400	Feb 10, 2031	DP U-282			
	9629861	Sep 21, 2030	DP			
<u>CEFTAROLINE FOSAMIL - TEFLARO</u>						
N 200327	002 8247400	Feb 10, 2031	DP U-282			
	9629861	Sep 21, 2030	DP			
<u>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</u>						
N 206829	001 10028963	Sep 07, 2032	U-2565		NCE	Dec 19, 2019
	10028963	Sep 07, 2032	U-2566		NPP	Apr 21, 2025
	10125149	Aug 14, 2035	DP		GAIN	Dec 19, 2024
	10376496	Sep 09, 2034	U-2610			
	10376496	Sep 09, 2034	U-2611			
	10420841	Mar 14, 2034	U-1672			
	10420841	Mar 14, 2034	U-2631			
	10420841	Mar 14, 2034	U-3360			
	10420841	Mar 14, 2034	U-3361			
	10933053	Sep 09, 2034	U-3090			
	10933053	Sep 09, 2034	U-3091			
	11278622	Mar 14, 2034	U-3335			
	11278622	Mar 14, 2034	U-3336			
	7129232	May 15, 2028	DS DP U-1676			
	7129232	May 15, 2028	DS DP U-3360			
	7129232	May 15, 2028	DS DP U-3361			
	7129232	May 15, 2028	DS DP U-36			
	8476425	Sep 27, 2032	DS			
	8685957	Sep 27, 2032	DS U-36			
	8906898	May 28, 2034	DS DP			
	8968753	Mar 14, 2034	U-1672			
	8968753	Mar 14, 2034	U-1673			
	8968753	Mar 14, 2034	U-3360			

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<u>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</u>						
N 206829	001 8968753	Mar 14, 2034	U-3361			
	9320740	Mar 14, 2034	DP			
	9724353	Sep 07, 2032	U-2565			
	9724353	Sep 07, 2032	U-2566			
	9872906	Mar 14, 2034	DP			
<u>CELECOXIB - ELYXYB</u>						
N 212157	001 10376527	May 27, 2036	DP U-2718			
	10722456	May 27, 2036	DP U-2718			
	10799517	May 27, 2036	DP U-2718			
	9572819	May 27, 2036	DP U-2718			
	9795620	May 27, 2036	DP U-2718			
	9949990	May 27, 2036	DP U-2718			
<u>CELECOXIB; TRAMADOL HYDROCHLORIDE - SEGLENTIS</u>						
N 213426	001 10238668	Apr 19, 2030	DS DP U-3244		NP	Oct 15, 2024
	10245276	Apr 19, 2030	DS DP			
	10548909	Apr 19, 2030	U-3244			
	11478488	Apr 19, 2030	U-3244			
	8598152	Apr 19, 2030	DS DP			
	8846744	Jun 03, 2031	DP			
	9012440	Apr 19, 2030	DS DP			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	001 11654133	Jun 16, 2039	U-3610		NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	002 11654133	Jun 16, 2039	U-3610		NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	003 11654133	Jun 16, 2039	U-3610		NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	004 11654133	Jun 16, 2039	U-3610		NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	005 11654133	Jun 16, 2039	U-3610		NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CENOBAMATE - XCOPRI</u>						
N 212839	006 11654133	Jun 16, 2039	U-3610		NCE	Mar 10, 2025
	7598279	Oct 30, 2032	DS			
<u>CERITINIB - ZYKADIA</u>						
N 205755	001 7893074	Apr 25, 2026	DS DP		ODE-145	May 26, 2024
	7964592	Apr 29, 2028	DS DP			
	8039479	Jun 29, 2030	DS DP			
	8377921	Nov 20, 2027	U-1179			
	8399450	Nov 20, 2027	DS DP			
	8703787	Feb 02, 2032	U-1179			
	9309229	Jan 18, 2032	DS DP			

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<u>CERITINIB - ZYKADIA</u>						
N 211225	001 7893074	Apr 25, 2026	DS DP		ODE*	May 26, 2024
	7964592	Apr 29, 2028	DS DP			
	8039479	Jun 29, 2030	DS DP			
	8377921	Nov 20, 2027		U-1179		
	8399450	Nov 20, 2027	DS DP			
	8703787	Feb 02, 2032		U-1179		
	9309229	Jan 18, 2032	DS DP			
<u>CETIRIZINE HYDROCHLORIDE - ZERVIATE</u>						
N 208694	001 8829005	Mar 15, 2030		U-1680		
	8829005*PED	Sep 15, 2030				
	9254286	Jul 09, 2032	DP			
	9254286*PED	Jan 09, 2033				
	9750684	Mar 15, 2030	DP			
	9993471	Mar 15, 2030		U-1680		
<u>CETIRIZINE HYDROCHLORIDE - OUZYTIR</u>						
N 211415	001 8263581	Feb 28, 2030		U-2635		
	8314083	Feb 28, 2030		U-2634		
	8513259	Feb 11, 2030		U-2636		
	9119771	Feb 11, 2030		U-2635		
	9180090	Feb 11, 2030		U-2635		
<u>CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE</u>						
N 021669	001 7066916	Feb 17, 2024		U-737		
	7427574	Apr 25, 2026	DP			
	7717889	Feb 27, 2025	DP	U-1022		
	7935093	Oct 02, 2027	DP	U-1022		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	002 6991394	Jan 31, 2024	DP			
	7422388	Apr 25, 2027	DP	U-1397		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	005 7422388	Apr 25, 2027	DP	U-1397		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
N 020832	006 6991394	Jan 31, 2024	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832	007 7422388	Apr 25, 2027	DP	U-1397		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - SOLUPREP S</u>						
N 208288	001				M-292	Jan 20, 2026
<u>CHLOROPROCAINE HYDROCHLORIDE - CLOROTEKAL</u>						
N 208791	001 8969412	Sep 05, 2026	DP	U-2609		
	9504666	Dec 11, 2033	DP			
<u>CHLOROPROCAINE HYDROCHLORIDE - IHEEZO</u>						
N 216227	001 10792271	Sep 15, 2038	DP	U-3457	NP	Sep 27, 2025
<u>CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE - TUXARIN ER</u>						
N 206323	001 9066942	Jan 03, 2032		U-1716		
	9107921	Jan 03, 2032	DP			

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<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - ADVIL ALLERGY SINUS</u>						
N 021441	001	7863287	Feb 28, 2027	DP		
<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - CHILDREN'S ADVIL ALLERGY SINUS</u>						
N 021587	001	10238640	May 25, 2024	DP		
<u>CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX - TUZISTRA XR</u>						
N 207768	001	8062667	Mar 29, 2029	DP		
		8790700	Mar 15, 2027	DP		
<u>CHOLINE FENOFIBRATE - TRILIPIX</u>						
N 022224	001	7259186	Jan 07, 2025	DS		
<u>CHOLINE FENOFIBRATE - TRILIPIX</u>						
N 022224	002	7259186	Jan 07, 2025	DS		
<u>CICLESONIDE - ALVESCO</u>						
N 021658	002	8371292	Feb 01, 2028		U-1355	
<u>CICLESONIDE - ALVESCO</u>						
N 021658	003	8371292	Feb 01, 2028		U-1355	
<u>CICLESONIDE - OMNARIS</u>						
N 022004	001	8371292	Feb 01, 2028		U-1356	
<u>CICLESONIDE - ZETONNA</u>						
N 202129	001	8371292	Feb 01, 2028		U-1357	
<u>CILASTATIN SODIUM; IMPENEM; RELEBACTAM - RECARBRIO</u>						
N 212819	001	8487093	Nov 19, 2029	DS DP	U-2586	NCE Jul 16, 2024
		8487093	Nov 19, 2029	DS DP	U-2587	GAIN Jul 16, 2029
		8487093	Nov 19, 2029	DS DP	U-2840	
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	001	7829595	Sep 22, 2026	DP	U-1098	
		9375405	Sep 22, 2026	DP		
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	002	7829595	Sep 22, 2026	DP	U-1098	
		9375405	Sep 22, 2026	DP		
<u>CINACALCET HYDROCHLORIDE - SENSIPAR</u>						
N 021688	003	7829595	Sep 22, 2026	DP	U-1098	
		9375405	Sep 22, 2026	DP		
<u>CIPROFLOXACIN - OTIPRIO</u>						
N 207986	001	11040004	Nov 12, 2037		U-2252	
		11246863	Nov 27, 2038	DP		
		11369566	Apr 21, 2029	DP		
		8318817	Apr 27, 2030		U-1792	
		9205048	Apr 21, 2029		U-1793	
		9220796	Jul 01, 2035	DP		
		9233068	Dec 11, 2029	DP		
		9603796	Apr 21, 2029	DS DP	U-2252	
<u>CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE - OTOVEL</u>						
N 208251	001	8932610	Mar 24, 2030	DP	U-1578	

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<u>CIPROFLOXACIN; DEXAMETHASONE - CIPRODEX</u>						
N 021537	001 8846650	Jun 04, 2025	DP U-1578			
<u>CITRIC ACID; LACTIC ACID; POTASSIUM BITARTRATE - PHEXXI</u>						
N 208352	001 10568855	Mar 15, 2033	U-1			
	11337989	Mar 15, 2033	U-1			
	11439610	Mar 15, 2033	DS DP			
	6706276	Mar 06, 2024	DP			
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - PREPOPIK</u>						
N 202535	001 8450338	Oct 10, 2028	DP			
	8481083	Oct 10, 2028	DP			
<u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - CLENPIO</u>						
N 209589	001 10624879	Jun 23, 2034	DP			
	11191753	Jun 23, 2034	U-3261			
	9827231	Jun 26, 2034	DP U-2162			
<u>CLADRIBINE - MAVENCLAD</u>						
N 022561	001 10849919	Nov 23, 2038	U-3411			
	7713947	Oct 16, 2026	U-2520			
	7888328	Apr 11, 2024	DP U-2521			
	8377903	May 31, 2026	U-2522			
	8785415	Apr 11, 2024	DP U-2523			
<u>CLASCOTERONE - WINLEVI</u>						
N 213433	001 10159682	Aug 14, 2028	U-2942		NCE	Aug 26, 2025
	11207332	Nov 20, 2028	DP U-3280			
	8143240	Jan 12, 2025	U-2942			
	8785427	Jul 25, 2030	DP			
	8865690	Jul 24, 2024	U-2942			
	9211295	Jul 24, 2024	DP			
	9433628	Feb 28, 2029	DP			
	9486458	Jul 24, 2028	U-2942			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	001 10010537	Oct 10, 2031	DP			
	11103490	Oct 10, 2031	DP			
	8658676	Oct 10, 2031	DP			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	002 10010537	Oct 10, 2031	DP			
	11103490	Oct 10, 2031	DP			
	8658676	Oct 10, 2031	DP			
<u>CLEVIDIPINE - CLEVIPREX</u>						
N 022156	003 10010537	Oct 10, 2031	DP			
	11103490	Oct 10, 2031	DP			
	8658676	Oct 10, 2031	DP			
<u>CLINDAMYCIN PHOSPHATE - CLINDESSE</u>						
N 050793	001 9789057	Dec 02, 2026	DP U-137			
<u>CLINDAMYCIN PHOSPHATE - EVOCLIN</u>						
N 050801	001 7141237	Feb 03, 2024	DP			
	7374747	Jan 23, 2024	DP U-921			

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<u>CLINDAMYCIN PHOSPHATE - XACIATO</u>						
N 215650	001 11129896	Sep 22, 2036	U-3293		NP	Dec 07, 2024
					GAIN	Dec 07, 2029
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	001 11541002	Jan 31, 2040	DP U-724			
	8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	002 11541002	Jan 31, 2040	DP U-724			
	8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			
<u>CLOBAZAM - SYMPAZAN</u>						
N 210833	003 11541002	Jan 31, 2040	DP U-724			
	8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			
<u>CLOBETASOL PROPIONATE - OLUX E</u>						
N 022013	001 8460641	Nov 05, 2028	DP U-1410			
	8962000	Aug 31, 2025	DP U-1410			
<u>CLOBETASOL PROPIONATE - IMPOYZ</u>						
N 209483	001 10064875	Aug 31, 2030	DP U-1408			
	10064875	Aug 31, 2030	DP U-1858			
	10064875	Aug 31, 2030	DP U-193			
	10064875	Aug 31, 2030	DP U-742			
	10064875	Aug 31, 2030	DP U-88			
	10588914	Aug 31, 2030	DP U-2771			
	9855334	Mar 11, 2035	DP			
	9956231	Aug 31, 2030	DP U-1408			
	9956231	Aug 31, 2030	DP U-1761			
	9956231	Aug 31, 2030	DP U-1858			
	9956231	Aug 31, 2030	DP U-193			
	9956231	Aug 31, 2030	DP U-742			
	9956231	Aug 31, 2030	DP U-88			
<u>CLONIDINE - NEXICLON XR</u>						
N 022500	001 8337890	Apr 17, 2027	DP			
	8623409	Sep 08, 2031	DP			
<u>CLONIDINE - NEXICLON XR</u>						
N 022500	002 8337890	Apr 17, 2027	DP			
	8623409	Sep 08, 2031	DP			
<u>CLOZAPINE - VERSACLOZ</u>						
N 203479	001 8057811	May 01, 2028	DP			
<u>COBICISTAT - TYBOST</u>						
N 203094	001 10039718	Oct 06, 2032	DP		ODE-260	Aug 22, 2026
	10039718*PED	Apr 06, 2033				
	8148374	Sep 03, 2029	DS DP U-1279			
	8148374*PED	Mar 03, 2030				

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<u>COBICISTAT; DARUNAVIR - PREZCOBIX</u>						
N 205395 001	10039718	Oct 06, 2032	DP			
	7700645	Dec 26, 2026	DS DP			
	7700645*PED	Jun 26, 2027				
	8148374	Sep 03, 2029	DS DP U-1279			
	8148374	Sep 03, 2029	DS DP U-2939			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>COBICISTAT; DARUNAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - SYMTUZA</u>						
N 210455 001	10039718	Oct 06, 2032	DP			
	10786518	Jul 19, 2038	U-2978			
	7390791	Apr 17, 2025	DS DP			
	7700645	Dec 26, 2026	DS DP			
	8148374	Sep 03, 2029	DS DP U-2353			
	8148374	Sep 03, 2029	DS DP U-2364			
	8148374	Sep 03, 2029	DS DP U-2365			
	8148374	Sep 03, 2029	DS DP U-2766			
	8148374	Sep 03, 2029	DS DP U-2767			
	8148374	Sep 03, 2029	DS DP U-2768			
	8518987	Feb 16, 2024	DS DP			
	8754065	Aug 15, 2032	DS DP U-2352			
	8754065	Aug 15, 2032	DS DP U-2765			
	9296769	Aug 15, 2032	DS DP U-2352			
	9296769	Aug 15, 2032	DS DP U-2765			
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</u>						
N 207561 001	10039718	Oct 06, 2032	DP			
	10039718*PED	Apr 06, 2033				
	7176220	Aug 27, 2026	DS DP U-257			
	7176220*PED	Feb 27, 2027				
	7390791	Apr 17, 2025	DS DP			
	7390791*PED	Oct 17, 2025				
	7635704	Oct 26, 2026	DS DP U-257			
	7635704*PED	Apr 26, 2027				
	8148374	Sep 03, 2029	DS DP U-1279			
	8148374*PED	Mar 03, 2030				
	8633219	Apr 30, 2030	DP U-257			
	8633219*PED	Oct 30, 2030				
	8754065	Aug 15, 2032	DS DP U-257			
	8754065*PED	Feb 15, 2033				
	8981103	Oct 26, 2026	DS DP			
	8981103*PED	Apr 26, 2027				
	9296769	Aug 15, 2032	DS DP U-257			
	9296769*PED	Feb 15, 2033				
	9891239	Sep 03, 2029	DP U-257			
	9891239*PED	Mar 03, 2030				
<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u>						
N 203100 001	10039718	Oct 06, 2032	DP			
	10039718*PED	Apr 06, 2033				
	7176220	Aug 27, 2026	DS DP U-257			

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<u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u>						
N 203100	001	7176220*PED				
		Feb 27, 2027				
		7635704	DS DP			U-257
		7635704*PED				
		Apr 26, 2027				
		8148374	DS DP			U-1279
		Sep 03, 2029				
		8592397				DP U-257
		Jan 13, 2024				
		8633219				DP U-257
		Apr 30, 2030				
		8633219*PED				
		Oct 30, 2030				
		8716264				DP U-257
		Jan 13, 2024				
		8981103	DS DP			
		Oct 26, 2026				
		8981103*PED				
		Apr 26, 2027				
		9457036				DP U-257
		Jan 13, 2024				
		9744181				DP U-257
		Jan 13, 2024				
		9891239				DP U-257
		Sep 03, 2029				
		9891239*PED				
		Mar 03, 2030				
<u>COBIMETINIB FUMARATE - COTELLIC</u>						
N 206192	001	10478400	DS DP		I-902	Oct 28, 2025
		10478400*PED			M-278	Jul 28, 2025
		Dec 29, 2036				
		10590102	DS DP		ODE-416	Oct 28, 2029
		Jun 30, 2036				
		10590102*PED			PED	Jan 28, 2026
		Dec 30, 2036				
		11087354				U-1776
		Jun 22, 2034				
		11087354*PED				
		Dec 22, 2034				
		11254649	DS DP			U-1776
		Jun 30, 2036				
		11254649*PED				
		Dec 30, 2036				
		11597699				U-3554
		Oct 05, 2026				
		7803839	DS DP			
		Nov 10, 2029				
		7803839*PED				
		May 10, 2030				
		8362002				U-1776
		Oct 05, 2026				
		8362002*PED				
		Apr 05, 2027				
<u>COCAINE HYDROCHLORIDE - NUMBRINO</u>						
N 209575	001	10016407				U-2225
		Feb 07, 2037				
		10016407				U-2226
		Feb 07, 2037				
		10016407				U-2329
		Feb 07, 2037				
		10149843				U-2478
		Feb 07, 2037				
		10149843				U-2479
		Feb 07, 2037				
		10231961				DP
		Feb 07, 2037				
		10413505				U-3014
		Feb 07, 2037				
		10420760				U-2478
		Feb 07, 2037				
		10857095				U-3014
		Feb 07, 2037				
		10894012				U-3014
		Feb 07, 2037				
		10933060				U-2478
		Feb 07, 2037				
		10933060				U-3014
		Feb 07, 2037				
		10973811				U-3680
		Feb 07, 2037				
		11040032				DP
		Feb 07, 2037				
		9867815				U-2225
		Feb 07, 2037				
		9867815				U-2226
		Feb 07, 2037				
		9867815				U-2227
		Feb 07, 2037				
		9867815				U-2329
		Feb 07, 2037				

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<u>COCAINE HYDROCHLORIDE - GOPRELTO</u>						
N 209963	001	10016407	Feb 07, 2037	U-2329		
		10149843	Feb 07, 2037	U-2478		
		10149843	Feb 07, 2037	U-2479		
		10231961	Feb 07, 2037	DP		
		10413505	Feb 07, 2037	U-2479		
		10420760	Feb 07, 2037	U-2478		
		10857095	Feb 07, 2037	U-3014		
		10894012	Feb 07, 2037	U-3014		
		10933060	Feb 07, 2037	U-3014		
		10973811	Feb 07, 2037	U-2226		
		10987347	Feb 07, 2037	U-2225		
		11040032	Feb 07, 2037	DP		
		9867815	Feb 07, 2037	U-2225		
		9867815	Feb 07, 2037	U-2226		
		9867815	Feb 07, 2037	U-2227		
<u>COLCHICINE - COLCHICINE</u>						
A 208678	001				PC	Apr 29, 2024
<u>COLCHICINE - COLCRYS</u>						
N 022352	001	7601758	Feb 10, 2029	U-1007		
		7619004	Dec 03, 2028	U-1020		
		7820681	Feb 17, 2029	U-1020		
		7906519	Feb 17, 2029	U-1116		
		7915269	Feb 17, 2029	U-1007		
		7935731	Dec 03, 2028	U-1116		
		7964647	Oct 06, 2028	U-1007		
		7964648	Oct 06, 2028	U-1161		
		7981938	Oct 06, 2028	U-1166		
		8093296	Oct 06, 2028	U-1007		
		8093297	Oct 06, 2028	U-1161		
		8093298	Oct 06, 2028	U-1116		
		8097655	Oct 06, 2028	U-1020		
		8415395	Oct 06, 2028	U-1007		
		8415396	Oct 06, 2028	U-1007		
		8440721	Feb 17, 2029	U-1007		
		8440722	Feb 17, 2029	U-1020		
<u>COLCHICINE - MITIGARE</u>						
N 204820	001	8927607	Aug 22, 2033	U-1020		
		9399036	Aug 22, 2033	U-1020		
		9555029	Aug 22, 2033	U-1020		
		9675613	Aug 22, 2033	U-1020		
		9789108	Aug 22, 2033	U-1020		
<u>COLCHICINE - GLOPERBA</u>						
N 210942	001	10226423	Dec 20, 2037	DP		
		10383820	Nov 22, 2036	DP	U-2814	
		10383821	Nov 22, 2036	DP		
		9907751	Nov 22, 2036	DP		

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<u>COLCHICINE - LODOCO</u>						
N 215727	001	10130585				
		Dec 31, 2034	U-3642			
		10206891				
		Jan 22, 2035	U-3641			
		10265281				
		Jan 22, 2035	U-3639			
		10842762				
		Jan 22, 2035	U-3640			
		11026899				
		Jan 22, 2035	U-3639			
		11026900				
		Jan 22, 2035	U-3639			
		11026901				
		Jan 22, 2035	U-3638			
		9744144				
		Jan 22, 2035	U-3643			
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 022362	001				M-232	Oct 20, 2024
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N 022362	002				M-232	Oct 20, 2024
<u>COPANLISIB DIHYDROCHLORIDE - ALIOOPA</u>						
N 209936	001	10383876		DS DP	ODE-155	Sep 14, 2024
		7511041		May 13, 2024	DS DP	
		9636344		Mar 29, 2032	U-2124	
		RE46856		Oct 22, 2029	DS DP U-2124	
<u>COPPER CU-64 DOTATATE - DETECTNET</u>						
N 213227	001	10159759			NCE	Sep 03, 2025
		10383961		Aug 23, 2032	U-2951	ODE-317
		11160888		Aug 23, 2032	U-2951	Sep 03, 2027
<u>CORTICOTROPIN - ACTHAR GEL</u>						
N 008372	008	11752199			U-3686	
		11752199		Feb 25, 2041	U-3687	
		11752199		Feb 25, 2041	U-3688	
<u>CRISABOROLE - EUCRISA</u>						
N 207695	001	8039451		DS DP	D-191	Apr 03, 2026
		8039451*PED		Dec 29, 2029		
		8168614		Jan 20, 2030	U-1932	
		8168614*PED		Jul 20, 2030		
		8501712		Feb 16, 2027	U-1932	
		8501712*PED		Aug 16, 2027		
		9682092		Feb 16, 2027	U-1932	
		9682092*PED		Aug 16, 2027		
<u>CRIZOTINIB - XALKORI</u>						
N 202570	001	7230098		DS	I-852	Jan 14, 2024
		7825137		May 12, 2027	U-3057	I-897
		7825137		May 12, 2027	U-3058	ODE-328
		7825137		May 12, 2027	U-3403	ODE-407
		7858643		Oct 08, 2029	DS DP	
		8217057		Nov 06, 2029	DS DP	
		8785632		Mar 01, 2025	DS	
<u>CRIZOTINIB - XALKORI</u>						
N 202570	002	7230098		DS	I-852	Jan 14, 2024
		7825137		May 12, 2027	U-3057	I-897
		7825137		May 12, 2027	U-3058	ODE-328

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<u>CRIZOTINIB - XALKORI</u>						
N 202570	002 7825137	May 12, 2027	U-3403		ODE-407	Jul 14, 2029
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS DP			
	8785632	Mar 01, 2025	DS			
<u>CRIZOTINIB - XALKORI</u>						
N 217581	001 7230098	Aug 26, 2025	DS		I-852	Jan 14, 2024
	7825137	May 12, 2027	U-3057		I-897	Jul 14, 2025
	7825137	May 12, 2027	U-3058			
	7825137	May 12, 2027	U-3403			
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS			
	8785632	Mar 01, 2025	DS DP			
<u>CRIZOTINIB - XALKORI</u>						
N 217581	002 7230098	Aug 26, 2025	DS		I-852	Jan 14, 2024
	7825137	May 12, 2027	U-3057		I-897	Jul 14, 2025
	7825137	May 12, 2027	U-3058			
	7825137	May 12, 2027	U-3403			
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS			
	8785632	Mar 01, 2025	DS DP			
<u>CRIZOTINIB - XALKORI</u>						
N 217581	003 7230098	Aug 26, 2025	DS		I-852	Jan 14, 2024
	7825137	May 12, 2027	U-3057		I-897	Jul 14, 2025
	7825137	May 12, 2027	U-3058			
	7825137	May 12, 2027	U-3403			
	7858643	Oct 08, 2029	DS DP			
	8217057	Nov 06, 2029	DS			
	8785632	Mar 01, 2025	DS DP			
<u>CROFELEMER - MYTESI</u>						
N 202292	001 8962680	Oct 31, 2031	U-1319			
	9585868	Oct 31, 2031	U-1319			
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE - TRALEMENT</u>						
N 209376	001 11786548	Jul 01, 2041	DP		NCE	Apr 30, 2024
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE - TRALEMENT</u>						
N 209376	002 11786548	Jul 01, 2041	DP		NCE	Apr 30, 2024
<u>CUPRIC SULFATE; MANGANESE SULFATE; SELENIOUS ACID; ZINC SULFATE - MULTRYS</u>						
N 209376	003 11786548	Jul 01, 2041	DP		NCE	Apr 30, 2024
<u>CYANOCOBALAMIN - NASCOBAL</u>						
N 021642	001 7229636	Aug 01, 2024	DP U-817			
	7404489	Mar 12, 2024	DP			
	7879349	Aug 01, 2024	DP U-1152			
	8003353	Aug 01, 2024	U-817			
	8940714	Feb 26, 2024	U-1152			
	9415007	Jul 28, 2024	U-1896			

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<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	001 9662342	Jun 26, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	002 9662342	Jun 26, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210735	003 9662342	Jun 26, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210852	001 10849916	Jul 13, 2035	DP			
	11382923	Dec 01, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210852	002 10849916	Jul 13, 2035	DP			
	11382923	Dec 01, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 210852	003 10849916	Jul 13, 2035	DP			
	11382923	Dec 01, 2035	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	001 10993952	Feb 15, 2036	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	002 10993952	Feb 15, 2036	DP			
<u>CYCLOPHOSPHAMIDE - CYCLOPHOSPHAMIDE</u>						
N 212501	003 10993952	Feb 15, 2036	DP			
<u>CYCLOSPORINE - RESTASIS MULTIDOSE</u>						
N 050790	002 8292129	Feb 25, 2031	DP			
	8561859	Apr 16, 2032	DP			
	8629111	Aug 27, 2024	DP			
	8633162	Aug 27, 2024		U-1479		
	8642556	Aug 27, 2024	DP			
	8648048	Aug 27, 2024		U-1483		
	8685930	Aug 27, 2024	DP			
	9248191	Aug 27, 2024		U-1479		
	9669974	May 11, 2034	DP			
	9676525	Feb 07, 2034	DP			
<u>CYCLOSPORINE - CEQUA</u>						
N 210913	001 10441630	Aug 23, 2033	DP			
	10918694	Feb 28, 2037	DP			
	8980839	Aug 23, 2033	DP	U-1483		
	9937225	Aug 23, 2033	DP	U-1483		
<u>CYCLOSPORINE - VERKAZIA</u>						
N 214965	001 11612658	Jan 27, 2026		U-3560	NP	Jun 23, 2024
	7973081	Jan 27, 2026	DP		ODE-358	Jun 23, 2028
	8298568	Nov 03, 2027	DP			
	8524779	Jan 27, 2026	DP			
	9132071	Jun 02, 2029	DP			
	9220694	Jan 27, 2026	DP			
	9956289	Jan 27, 2026	DP			

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<u>CYCLOSPORINE - VEVYE</u>						
N 217469 001	10813976	Sep 22, 2037	DP		NP	May 30, 2026
	11154513	Nov 20, 2038	DP U-1900			
	11413323	Oct 11, 2039	U-3627			
	8614178	Dec 13, 2030	DP			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389 001	10143665	Aug 16, 2036	U-1399		ODE-162	Dec 22, 2024
	10143665*PED	Feb 16, 2037			PED	Jun 22, 2025
	10328037	Aug 16, 2036	U-1399			
	10328037*PED	Feb 16, 2037				
	10548859	Aug 16, 2036	U-1399			
	10548859*PED	Feb 16, 2037				
	10905662	Aug 16, 2036	U-1399			
	10905662*PED	Feb 16, 2037				
	8026284	Sep 22, 2027	U-1399			
	8026284*PED	Mar 22, 2028				
	9173851	Jun 17, 2034	DP			
	9173851*PED	Dec 17, 2034				
	9192590	Jan 26, 2027	U-1399			
	9192590*PED	Jul 26, 2027				
	9198882	Jan 26, 2027	U-1399			
	9198882*PED	Jul 26, 2027				
	9233077	Jun 17, 2034	DP			
	9233077*PED	Dec 17, 2034				
	9925156	Jan 26, 2027	DS DP U-1399			
	9925156*PED	Jul 26, 2027				
	9925157	Jan 26, 2027	DS DP U-1399			
	9925157*PED	Jul 26, 2027				
	9925158	Jan 26, 2027	DS DP U-1399			
	9925158*PED	Jul 26, 2027				
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389 002	10143665	Aug 16, 2036	U-1399		ODE-162	Dec 22, 2024
	10143665*PED	Feb 16, 2037			PED	Jun 22, 2025
	10328037	Aug 16, 2036	U-1399			
	10328037*PED	Feb 16, 2037				
	10548859	Aug 16, 2036	U-1399			
	10548859*PED	Feb 16, 2037				
	10905662	Aug 16, 2036	U-1399			
	10905662*PED	Feb 16, 2037				
	8026284	Sep 22, 2027	U-1399			
	8026284*PED	Mar 22, 2028				
	9173851	Jun 17, 2034	DP			
	9173851*PED	Dec 17, 2034				
	9192590	Jan 26, 2027	U-1399			
	9192590*PED	Jul 26, 2027				
	9198882	Jan 26, 2027	U-1399			
	9198882*PED	Jul 26, 2027				
	9233077	Jun 17, 2034	DP			
	9233077*PED	Dec 17, 2034				
	9925156	Jan 26, 2027	DS DP U-1399			

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<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 203389	002	9925156*PED	Jul 26, 2027			
		9925157	Jan 26, 2027	DS DP	U-1399	
		9925157*PED	Jul 26, 2027			
		9925158	Jan 26, 2027	DS DP	U-1399	
		9925158*PED	Jul 26, 2027			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	001	10143665	Aug 16, 2036	U-1399	ODE*	Dec 22, 2024
		10143665*PED	Feb 16, 2037		PED	Jun 22, 2025
		10328037	Aug 16, 2036	U-1399		
		10328037*PED	Feb 16, 2037			
		10548859	Aug 16, 2036	U-1399		
		10548859*PED	Feb 16, 2037			
		10905662	Aug 16, 2036	U-1399		
		10905662*PED	Feb 16, 2037			
		8026284	Sep 22, 2027	U-1399		
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027	U-1399		
		9192590*PED	Jul 26, 2027			
		9198882	Jan 26, 2027	U-1399		
		9198882*PED	Jul 26, 2027			
		9233077	Jun 17, 2034	DP		
		9233077*PED	Dec 17, 2034			
		9925156	Jan 26, 2027	DP	U-1399	
		9925156*PED	Jul 26, 2027			
		9925157	Jan 26, 2027	DP	U-1399	
		9925157*PED	Jul 26, 2027			
		9925158	Jan 26, 2027	DP	U-1399	
		9925158*PED	Jul 26, 2027			
<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	002	10143665	Aug 16, 2036	U-1399	ODE*	Dec 22, 2024
		10143665*PED	Feb 16, 2037		PED	Jun 22, 2025
		10328037	Aug 16, 2036	U-1399		
		10328037*PED	Feb 16, 2037			
		10548859	Aug 16, 2036	U-1399		
		10548859*PED	Feb 16, 2037			
		10905662	Aug 16, 2036	U-1399		
		10905662*PED	Feb 16, 2037			
		8026284	Sep 22, 2027	U-1399		
		8026284*PED	Mar 22, 2028			
		9173851	Jun 17, 2034	DP		
		9173851*PED	Dec 17, 2034			
		9192590	Jan 26, 2027	U-1399		
		9192590*PED	Jul 26, 2027			
		9198882	Jan 26, 2027	U-1399		
		9198882*PED	Jul 26, 2027			
		9233077	Jun 17, 2034	DP		
		9233077*PED	Dec 17, 2034			

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<u>CYSTEAMINE BITARTRATE - PROCYSBI</u>						
N 213491	002 9925156	Jan 26, 2027	DP U-1399			
	9925156*PED	Jul 26, 2027				
	9925157	Jan 26, 2027	DP U-1399			
	9925157*PED	Jul 26, 2027				
	9925158	Jan 26, 2027	DP U-1399			
	9925158*PED	Jul 26, 2027				
<u>CYSTEINE HYDROCHLORIDE - ELCYS</u>						
N 210660	001 10478453	Jan 15, 2039	DP U-2752			
	10583155	Jan 15, 2039	DP U-2752			
	10653719	Jan 15, 2039	DP			
	10905713	Jan 15, 2039	DP			
	10905714	Jan 15, 2039	DP			
	10912795	Jan 15, 2039	DP			
	10918662	Jan 15, 2039	DP			
	10933089	Jan 15, 2039	DP			
	11510941	Jan 15, 2039	DP			
	11510942	Jan 15, 2039	DP U-2752			
	11642370	Jan 15, 2039	DP			
	11648262	Jan 15, 2039	DP			
	11679125	Jan 15, 2039	DP			
	11684636	Jan 15, 2039	DP			
	11826383	Jan 15, 2039	DP			
<u>CYSTEINE HYDROCHLORIDE - NOURESS</u>						
N 212535	001 10478453	Jan 15, 2039	DP U-2752			
	10493051	Mar 15, 2039	DP			
	10543186	Mar 15, 2039	U-2722			
	10583155	Jan 15, 2039	DP U-2752			
	10653719	Jan 15, 2039	DP			
	10702490	Mar 15, 2039	DP			
	10905713	Jan 15, 2039	DP			
	10905714	Jan 15, 2039	DP			
	10912795	Jan 15, 2039	DP			
	10918662	Jan 15, 2039	DP			
	10933089	Jan 15, 2039	DP			
	11045438	Mar 15, 2039	DP			
	11510941	Mar 02, 2039	DP			
	11510942	Jan 15, 2039	DP U-2752			
<u>CYTARABINE; DAUNORUBICIN - VYXEOS</u>						
N 209401	001 10028912	Oct 15, 2032	DP U-3149		NPP	Mar 30, 2024
	10028912	Oct 15, 2032	DP U-3150		ODE-287	Aug 03, 2024
	10166184	Oct 15, 2032	DP U-3149		ODE-350	Mar 30, 2028
	10835492	Oct 15, 2032	U-3150			
	7850990	Jan 23, 2027	DP U-3147			
	8022279	Sep 14, 2027	DP U-3147			
	8092828	Apr 01, 2029	U-3147			
	8431806	Apr 22, 2025	DP U-3147			
	8518437	Jun 07, 2026	DP			
	9271931	Jan 23, 2027	DP			

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<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512 001	7932273	Sep 07, 2025	DS DP		I-862	Jun 21, 2024
	9034822	Jan 20, 2031	U-1759		PED	Dec 21, 2024
	9034822*PED	Jul 20, 2031				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512 002	7932273	Sep 07, 2025	DS DP		I-862	Jun 21, 2024
	9034822	Jan 20, 2031	U-1759		PED	Dec 21, 2024
	9034822*PED	Jul 20, 2031				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 022512 003	7866474	Aug 31, 2027	DP	Y	I-862	Jun 21, 2024
	7866474*PED	Mar 02, 2028			PED	Dec 21, 2024
	7932273	Sep 07, 2025	DS DP			
	7932273*PED	Mar 07, 2026				
	9034822	Jan 20, 2031	U-1759			
	9034822*PED	Jul 20, 2031				
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358 001	7932273	Sep 07, 2025	DS DP		NP	Jun 21, 2024
	7932273*PED	Mar 07, 2026			PED	Dec 21, 2024
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358 002	7932273	Sep 07, 2025	DS DP		NP	Jun 21, 2024
	7932273*PED	Mar 07, 2026			PED	Dec 21, 2024
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358 003	7932273	Sep 07, 2025	DS DP		NP	Jun 21, 2024
	7932273*PED	Mar 07, 2026			PED	Dec 21, 2024
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358 004	7932273	Sep 07, 2025	DS DP		NP	Jun 21, 2024
	7932273*PED	Mar 07, 2026			PED	Dec 21, 2024
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358 005	7932273	Sep 07, 2025	DS DP		NP	Jun 21, 2024
	7932273*PED	Mar 07, 2026			PED	Dec 21, 2024
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>						
N 214358 006	7932273	Sep 07, 2025	DS DP		NP	Jun 21, 2024
	7932273*PED	Mar 07, 2026			PED	Dec 21, 2024
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 001	10869869	Aug 30, 2033	U-3185		I-894	Jun 22, 2025
	10869869*PED	Mar 02, 2034			I-908	Mar 16, 2026
	7994185	Jan 20, 2030	DS DP U-1406		ODE-147	Jun 22, 2024
	7994185	Jan 20, 2030	DS DP U-2031		ODE-182	Apr 30, 2025
	7994185	Jan 20, 2030	DS DP U-2032		ODE-183	May 04, 2025
	7994185	Jan 20, 2030	DS DP U-2296		ODE-428	Mar 16, 2030
	7994185*PED	Jul 20, 2030			PED	Dec 22, 2024
	8415345	Jan 20, 2030	DS DP U-1406		PED	Oct 30, 2025
	8415345	Jan 20, 2030	DS DP U-2031		PED	Nov 04, 2025
	8415345	Jan 20, 2030	DS DP U-2032		PED	Dec 22, 2025
	8415345	Jan 20, 2030	DS DP U-2296			
	8415345*PED	Jul 20, 2030				
	8703781	Oct 15, 2030	DS DP U-1713			

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<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 001	8703781	Oct 15, 2030	DS DP U-2032			
	8703781	Oct 15, 2030	DS DP U-2296			
	8703781	Oct 15, 2030	DS DP U-2298			
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-2026			
	8835443	Jun 10, 2025	U-2027			
	8835443	Jun 10, 2025	U-2296			
	8835443	Jun 10, 2025	U-2298			
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030	U-2027			
	8952018*PED	Apr 15, 2031				
	9233956	May 04, 2029	U-1811			
	9233956	May 04, 2029	U-2031			
	9233956	May 04, 2029	U-2032			
	9233956	May 04, 2029	U-2296			
	9233956*PED	Nov 04, 2029				
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 202806 002	10869869	Aug 30, 2033	U-3185		I-894	Jun 22, 2025
	10869869*PED	Mar 02, 2034			I-908	Mar 16, 2026
	7994185	Jan 20, 2030	DS DP U-1406		ODE-147	Jun 22, 2024
	7994185	Jan 20, 2030	DS DP U-2031		ODE-182	Apr 30, 2025
	7994185	Jan 20, 2030	DS DP U-2032		ODE-183	May 04, 2025
	7994185	Jan 20, 2030	DS DP U-2296		ODE-428	Mar 16, 2030
	7994185*PED	Jul 20, 2030			PED	Dec 22, 2024
	8415345	Jan 20, 2030	DS DP U-1406		PED	Oct 30, 2025
	8415345	Jan 20, 2030	DS DP U-2031		PED	Nov 04, 2025
	8415345	Jan 20, 2030	DS DP U-2032		PED	Dec 22, 2025
	8415345	Jan 20, 2030	DS DP U-2296			
	8415345*PED	Jul 20, 2030				
	8703781	Oct 15, 2030	DS DP U-1713			
	8703781	Oct 15, 2030	DS DP U-2032			
	8703781	Oct 15, 2030	DS DP U-2296			
	8703781	Oct 15, 2030	DS DP U-2298			
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-2026			
	8835443	Jun 10, 2025	U-2027			
	8835443	Jun 10, 2025	U-2296			
	8835443	Jun 10, 2025	U-2298			
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030	U-2027			
	8952018*PED	Apr 15, 2031				
	9233956	May 04, 2029	U-1811			
	9233956	May 04, 2029	U-2031			
	9233956	May 04, 2029	U-2032			
	9233956	May 04, 2029	U-2296			
	9233956*PED	Nov 04, 2029				
<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 217514 001	11504333	Jun 29, 2038	DP		NP	Mar 16, 2026
	7994185	Jan 20, 2030	DS DP		ODE-428	Mar 16, 2030

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<u>DABRAFENIB MESYLATE - TAFINLAR</u>						
N 217514	001 8415345	Jan 20, 2030	DS DP			
	8703781	Oct 15, 2030	DS DP U-3565			
	8835443	Jun 10, 2025	U-3565			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843	001 8329159	Jul 24, 2029	DS			
	8629171	Jun 13, 2031	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1725			
	8900566	Aug 08, 2027	U-1724			
	8900566	Aug 08, 2027	U-1725			
	9421192	Aug 08, 2027	DS U-1724			
	9421192	Aug 08, 2027	DS U-1725			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843	002 8329159	Jul 24, 2029	DS			
	8629171	Jun 13, 2031	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1724			
	8642025	Aug 11, 2027	DS DP U-1725			
	8900566	Aug 08, 2027	U-1724			
	8900566	Aug 08, 2027	U-1725			
	9421192	Aug 08, 2027	DS U-1724			
	9421192	Aug 08, 2027	DS U-1725			
<u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u>						
N 206843	003 9421192	Aug 08, 2027	DS U-1724			
	9421192	Aug 08, 2027	DS U-1725			
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288	001 10596162	Feb 02, 2026	U-3338		ODE-206	Sep 27, 2025
	10603314	Feb 02, 2026	U-3337		ODE-213	Sep 27, 2025
	7772243	Aug 26, 2028	DS DP			
	8623883	May 05, 2025	U-1403			
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288	002 10596162	Feb 02, 2026	U-3338		ODE-206	Sep 27, 2025
	10603314	Feb 02, 2026	U-3337		ODE-213	Sep 27, 2025
	7772243	Aug 26, 2028	DS DP			
	8623883	May 05, 2025	U-1403			
<u>DACOMITINIB - VIZIMPRO</u>						
N 211288	003 10596162	Feb 02, 2026	U-3338		ODE-206	Sep 27, 2025
	10603314	Feb 02, 2026	U-3337		ODE-213	Sep 27, 2025
	7772243	Aug 26, 2028	DS DP			
	8623883	May 05, 2025	U-1403			
<u>DALBAVANCIN HYDROCHLORIDE - DALVANCE</u>						
N 021883	001 6900175	May 23, 2028	U-3499		NCE	May 23, 2019
					NPP	Jul 22, 2024
					GAIN	May 23, 2024
<u>DANTROLENE SODIUM - RYANODEX</u>						
N 205579	001 7758890	Jun 30, 2025	DP			

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<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293 001	6515117	Oct 04, 2025	DS DP U-2139		I-857	Apr 30, 2024
	6515117	Oct 04, 2025	DS DP U-493		M-298	May 08, 2026
	7456254	Jun 30, 2025	DP U-2139			
	7851502	Aug 19, 2028	DP			
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028	DP			
	8329648	Aug 18, 2026	U-2139			
	8329648	Aug 18, 2026	U-2212			
	8329648	Aug 18, 2026	U-2213			
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8431685	Apr 13, 2025	DP U-2139			
	8461105	Apr 13, 2025	DP U-2139			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	8716251	Mar 21, 2028	DP			
	8721615	Jan 18, 2030	DP	Y		
	8906851	Aug 18, 2026	U-2139			
	9238076	Apr 15, 2024	DP U-2139			
<u>DAPAGLIFLOZIN - FARXIGA</u>						
N 202293 002	10973836	Mar 09, 2040	U-3127		I-857	Apr 30, 2024
	11826376	Jul 18, 2039	U-3766		M-298	May 08, 2026
	6515117	Oct 04, 2025	DS DP U-2139			
	6515117	Oct 04, 2025	DS DP U-493			
	7456254	Jun 30, 2025	DP U-2139			
	7851502	Aug 19, 2028	DP			
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028	DP			
	8329648	Aug 18, 2026	U-2139			
	8329648	Aug 18, 2026	U-2212			
	8329648	Aug 18, 2026	U-2213			
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8431685	Apr 13, 2025	DP U-2139			
	8461105	Apr 13, 2025	DP U-2139			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	8716251	Mar 21, 2028	DP			
	8721615	Jan 18, 2030	DP	Y		
	8906851	Aug 18, 2026	U-2139			
	9238076	Apr 15, 2024	DP U-2139			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 001	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9616028	Nov 12, 2030	DP			

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<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 002	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 003	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 004	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u>						
N 205649 005	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - OTERNMET XR</u>						
N 210874 001	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - OTERNMET XR</u>						
N 210874 002	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - OTERNMET XR</u>						
N 210874 003	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8628799	Jul 13, 2025	DP			
	8716251	Mar 21, 2028	DP			
	9616028	Nov 12, 2030	DP			
<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - OTERNMET XR</u>						
N 210874 004	6515117	Oct 04, 2025	DS DP U-493			
	7919598	Dec 16, 2029	DS			

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<u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - QTERNMET XR</u>						
N 210874	004	8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
		9616028	Nov 12, 2030	DP		
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091	001	6515117	Oct 04, 2025	DS DP U-493		
		7919598	Dec 16, 2029	DS		
		8221786	Mar 21, 2028	DP		
		8361972	Mar 21, 2028	U-1976		
		8361972	Mar 21, 2028	U-1977		
		8361972	Mar 21, 2028	U-493		
		8501698	Jun 20, 2027	DP U-1976		
		8501698	Jun 20, 2027	DP U-1977		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
<u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u>						
N 209091	002	6515117	Oct 04, 2025	DS DP U-493		
		7919598	Dec 16, 2029	DS		
		8221786	Mar 21, 2028	DP		
		8361972	Mar 21, 2028	U-493		
		8501698	Jun 20, 2027	DP U-493		
		8628799	Jul 13, 2025	DP		
		8716251	Mar 21, 2028	DP		
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951	001	11117871	Mar 13, 2038	DS DP U-3535	NCE	Feb 01, 2028
		11643397	Jun 22, 2027	DS U-3535		
		11649217	Mar 13, 2038	DP U-3535		
		8324208	Dec 11, 2028	DS DP		
		8557834	Jun 22, 2027	U-1238		
		8815884	Jun 22, 2027	DP U-1238		
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951	002	11117871	Mar 13, 2038	DS DP U-3535	NCE	Feb 01, 2028
		11643397	Jun 22, 2027	DS U-3535		
		11649217	Mar 13, 2038	DP U-3535		
		8324208	Dec 11, 2028	DS DP		
		8557834	Jun 22, 2027	U-1238		
		8815884	Jun 22, 2027	DP U-1238		
<u>DAPRODUSTAT - JESDUVROO</u>						
N 216951	003	11117871	Mar 13, 2038	DS DP U-3535	NCE	Feb 01, 2028
		11643397	Jun 22, 2027	DS U-3535		
		11649217	Mar 13, 2038	DP U-3535		
		8324208	Dec 11, 2028	DS DP		
		8557834	Jun 22, 2027	U-1238		
		8815884	Jun 22, 2027	DP U-1238		

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<u>DAPRODUSTAT - JESDUVROQ</u>						
N 216951	004	11117871	Mar 13, 2038	DS DP U-3535	NCE	Feb 01, 2028
		11643397	Jun 22, 2027	DS U-3535		
		11649217	Mar 13, 2038	DP U-3535		
		8324208	Dec 11, 2028	DS DP		
		8557834	Jun 22, 2027	U-1238		
		8815884	Jun 22, 2027	DP U-1238		
<u>DAPRODUSTAT - JESDUVROQ</u>						
N 216951	005	11117871	Mar 13, 2038	DS DP U-3535	NCE	Feb 01, 2028
		11643397	Jun 22, 2027	DS U-3535		
		11649217	Mar 13, 2038	DP U-3535		
		8324208	Dec 11, 2028	DS DP		
		8557834	Jun 22, 2027	U-1238		
		8815884	Jun 22, 2027	DP U-1238		
<u>DAPSONE - ACZONE</u>						
N 207154	001	11273132	Nov 18, 2033	DP		
		9161926	Nov 18, 2033	DP		
		9517219	Nov 18, 2033	U-1033		
<u>DAPTOMYCIN - CUBICIN</u>						
N 021572	002	8003673	Sep 04, 2028	U-1180		
<u>DAPTOMYCIN - CUBICIN RF</u>						
N 021572	003	9138456	Nov 23, 2030	DP		
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 210282	001	10357535	Sep 11, 2033	DP U-3176		
		9655946	Sep 11, 2033	DP U-3175		
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 210282	002	10357535	Sep 11, 2033	DP U-3176		
		9655946	Sep 11, 2033	DP U-3175		
<u>DAPTOMYCIN - DAPZURA RT</u>						
N 213645	001	11173189	Mar 11, 2041	DP U-3294		
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 217415	001	11759497	Aug 28, 2038	DP		
<u>DAPTOMYCIN - DAPTOMYCIN</u>						
N 217415	002	11759497	Aug 28, 2038	DP		
<u>DARIDOREXANT HYDROCHLORIDE - QUVIVIQ</u>						
N 214985	001	10023560	Dec 02, 2034	U-620	NCE	Apr 07, 2027
		9732075	Jun 12, 2033	DS DP U-620		
		9790208	Dec 02, 2034	DS DP		
<u>DARIDOREXANT HYDROCHLORIDE - QUVIVIQ</u>						
N 214985	002	10023560	Dec 02, 2034	U-620	NCE	Apr 07, 2027
		9732075	Jun 12, 2033	DS DP U-620		
		9790208	Dec 02, 2034	DS DP		
<u>DAROLUTAMIDE - NUBEQA</u>						
N 212099	001	10010530	Jan 28, 2036	DS	I-900	Aug 05, 2025
		10383853	Jan 28, 2036	DS	NCE	Jul 30, 2024
		10711013	Oct 27, 2030	DS DP		

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<u>DAROLUTAMIDE - NUBEQA</u>						
N 212099 001	10835515	Jan 28, 2036	DP U-2605			
	11046713	Oct 27, 2030	DS			
	11168058	Feb 27, 2038	DS DP			
	8975254	Mar 25, 2033	DS DP U-2605			
	9657003	Oct 27, 2030	DS DP U-2605			
<u>DARUNAVIR - PREZISTA</u>						
N 021976 001	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 002	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 003	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 004	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 005	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 021976 006	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DARUNAVIR - PREZISTA</u>						
N 202895 001	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
<u>DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619 001	10201542	Oct 18, 2033	DP U-1753			
	8188104	May 17, 2029	DS DP U-1636			
	8268349	Aug 25, 2024	DP			
	8399015	Aug 25, 2024	DP			
	8420596	Apr 10, 2031	DS DP			
	8466159	Sep 04, 2032	U-1637			
	8492386	Sep 04, 2032	U-1840			
	8501238	Dec 19, 2028	DS DP U-1636			
	8642538	Sep 10, 2029	DS DP U-1638			
	8680106	Sep 04, 2032	U-1637			
	8685984	Sep 04, 2032	U-1840			
	8686026	Jun 09, 2031	DP			

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<u>DASABUVIR SODIUM; OMBITASVIR, PARITAPREVIR, RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619	001	8691938	Apr 13, 2032	DS DP		
		9006387	Jun 10, 2030		U-1687	
		9044480	Apr 10, 2031		U-1638	
		9139536	Nov 09, 2028		U-1753	
		9629841	Oct 18, 2033	DP	U-1753	
<u>DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA XR</u>						
N 208624	001	10105365	Jan 02, 2035	DP	U-1889	
		10201541	May 17, 2032	DP		
		10201584	May 17, 2032		U-1889	
		8188104	May 17, 2029	DS DP	U-1636	
		8268349	Aug 25, 2024	DP		
		8399015	Aug 25, 2024	DP		
		8420596	Apr 10, 2031	DS DP		
		8466159	Sep 04, 2032		U-1637	
		8492386	Sep 04, 2032		U-1840	
		8501238	Sep 17, 2028	DS DP	U-1636	
		8642538	Sep 10, 2029	DS DP	U-1638	
		8680106	Sep 04, 2032		U-1637	
		8685984	Sep 04, 2032		U-1840	
		8686026	Jun 09, 2031	DP		
		8691938	Apr 13, 2032	DS DP		
		9006387	Jun 10, 2030		U-1687	
		9044480	Apr 10, 2031		U-1638	
		9139536	Nov 09, 2028		U-1753	
		9333204	Jan 02, 2035	DP	U-1889	
		9744170	Jan 02, 2035	DP	U-1889	
<u>DASATINIB - SPRYCEL</u>						
N 021986	001	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	002	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	003	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986	004	7491725	Mar 28, 2026	DS DP	ODE-164	Nov 09, 2024
		7491725*PED	Sep 28, 2026		ODE-225	Dec 21, 2025
		8680103	Feb 04, 2025	DP	PED	May 09, 2025
		8680103*PED	Aug 04, 2025		PED	Jun 21, 2026

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<u>DASATINIB - SPRYCEL</u>						
N 021986 005	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - SPRYCEL</u>						
N 021986 006	7491725	Mar 28, 2026	DS DP		ODE-164	Nov 09, 2024
	7491725*PED	Sep 28, 2026			ODE-225	Dec 21, 2025
	8680103	Feb 04, 2025	DP		PED	May 09, 2025
	8680103*PED	Aug 04, 2025			PED	Jun 21, 2026
<u>DASATINIB - PHYRAGO</u>						
N 216099 001	11202778	Jan 22, 2041		U-3770		
	11202778	Jan 22, 2041		U-3771		
	11202778	Jan 22, 2041		U-3772		
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041		U-3767		
	11324745	Jan 22, 2041		U-3768		
	11324745	Jan 22, 2041		U-3769		
<u>DASATINIB - PHYRAGO</u>						
N 216099 002	11202778	Jan 22, 2041		U-3770		
	11202778	Jan 22, 2041		U-3771		
	11202778	Jan 22, 2041		U-3772		
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041		U-3767		
	11324745	Jan 22, 2041		U-3768		
	11324745	Jan 22, 2041		U-3769		
<u>DASATINIB - PHYRAGO</u>						
N 216099 003	11202778	Jan 22, 2041		U-3770		
	11202778	Jan 22, 2041		U-3771		
	11202778	Jan 22, 2041		U-3772		
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041		U-3767		
	11324745	Jan 22, 2041		U-3768		
	11324745	Jan 22, 2041		U-3769		
<u>DASATINIB - PHYRAGO</u>						
N 216099 004	11202778	Jan 22, 2041		U-3770		
	11202778	Jan 22, 2041		U-3771		
	11202778	Jan 22, 2041		U-3772		
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041		U-3767		
	11324745	Jan 22, 2041		U-3768		
	11324745	Jan 22, 2041		U-3769		
<u>DASATINIB - PHYRAGO</u>						
N 216099 005	11202778	Jan 22, 2041		U-3770		
	11202778	Jan 22, 2041		U-3771		
	11202778	Jan 22, 2041		U-3772		
	11298356	Jan 22, 2041	DP			
	11324745	Jan 22, 2041		U-3767		

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<u>DASATINIB - PHYRAGO</u>						
N 216099	005	11324745	Jan 22, 2041	U-3768		
		11324745	Jan 22, 2041	U-3769		
<u>DASATINIB - PHYRAGO</u>						
N 216099	006	11202778	Jan 22, 2041	U-3770		
		11202778	Jan 22, 2041	U-3771		
		11202778	Jan 22, 2041	U-3772		
		11298356	Jan 22, 2041	DP		
		11324745	Jan 22, 2041	U-3767		
		11324745	Jan 22, 2041	U-3768		
		11324745	Jan 22, 2041	U-3769		
<u>DASIGLUCAGON HYDROCHLORIDE - ZEGALOGUE</u>						
N 214231	001	10442847	Feb 03, 2035	DS DP	NCE	Mar 22, 2026
		11795204	Jan 06, 2034	U-3752		
<u>DASIGLUCAGON HYDROCHLORIDE - ZEGALOGUE (AUTOINJECTOR)</u>						
N 214231	002	10442847	Feb 03, 2035	DS DP	NCE	Mar 22, 2026
		11795204	Jan 06, 2034	U-3752		
<u>DEFERASIROX - JADENU</u>						
N 206910	001	9283209	Nov 21, 2034	DS DP		
<u>DEFERASIROX - JADENU</u>						
N 206910	002	9283209	Nov 21, 2034	DS DP		
<u>DEFERASIROX - JADENU</u>						
N 206910	003	9283209	Nov 21, 2034	DS DP		
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	001				I-859	Apr 30, 2024
					ODE-417	Apr 30, 2028
					ODE-420	Apr 30, 2028
					ODE-421	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 021825	002				I-859	Apr 30, 2024
					ODE-417	Apr 30, 2028
					ODE-420	Apr 30, 2028
					ODE-421	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	001	8703156	Oct 26, 2029	DP U-3083	I-859	Apr 30, 2024
					ODE-417	Apr 30, 2028
					ODE-418	Apr 30, 2028
					ODE-419	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 208030	002	8703156	Oct 26, 2029	DP U-3083	I-859	Apr 30, 2024
					ODE-417	Apr 30, 2028
					ODE-418	Apr 30, 2028
					ODE-419	Apr 30, 2028
<u>DEFERIPRONE - FERRIPROX</u>						
N 212269	001	10780055	Oct 25, 2038	DP U-3083	I-859	Apr 30, 2024
		10940115	Oct 25, 2038	DP U-3083	ODE-417	Apr 30, 2028

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<u>DEFERIPRONE - FERRIPROX</u>						
N 212269	001 10940116	Oct 25, 2038	DP		ODE-420	Apr 30, 2028
	11357731	Oct 25, 2038	U-3083		ODE-421	Apr 30, 2028
	11458103	Oct 25, 2038	DP			
	11723874	Oct 25, 2038	U-3083			
<u>DEFIBROTIDE SODIUM - DEFITELIO</u>						
N 208114	001 11085043	Jun 22, 2032	DP			
	11236328	Jun 22, 2032	U-3301			
	11746348	Jun 22, 2032	DP			
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	001				ODE-130	Feb 09, 2024
					ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	002				ODE-130	Feb 09, 2024
					ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	003				ODE-130	Feb 09, 2024
					ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208684	004				ODE-130	Feb 09, 2024
					ODE-252	Jun 07, 2026
<u>DEFLAZACORT - EMFLAZA</u>						
N 208685	001				ODE-130	Feb 09, 2024
					ODE-252	Jun 07, 2026
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	001 10695398	Apr 27, 2032	U-1895			
	10729739	Feb 10, 2029	U-1978			
	10973870	Feb 10, 2029	U-1978			
	11766468	Apr 27, 2032	U-1978			
	11826397	Apr 27, 2032	U-1978			
	9415085	Apr 27, 2032	U-1895			
	9579359	Feb 10, 2029	U-1978			
<u>DEGARELIX ACETATE - FIRMAGON</u>						
N 022201	002 10695398	Apr 27, 2032	U-1895			
	10729739	Feb 10, 2029	U-1978			
	10973870	Feb 10, 2029	U-1978			
	11766468	Apr 27, 2032	U-1978			
	11826397	Apr 27, 2032	U-1978			
	9415085	Apr 27, 2032	U-1895			
	9579359	Feb 10, 2029	U-1978			
<u>DELAFLOXACIN MEGLUMINE - BAXDELA</u>						
N 208610	001 7728143	Jun 19, 2031	DS		NCE	Jun 19, 2022
	8252813	Oct 02, 2026	DP U-2028		GAIN	Jun 19, 2027
	8273892	Aug 06, 2026	DS			
	8648093	Oct 07, 2025	DP U-2028			
	8871938	Sep 23, 2029	DS			
	8969569	Oct 07, 2025	DP U-2028			

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<u>DELAFLORACIN MEGLUMINE - BAXDELA</u>						
N 208610	001 9539250	Oct 07, 2025	DS DP U-2028			
	RE46617	Dec 28, 2029	DS			
<u>DELAFLORACIN MEGLUMINE - BAXDELA</u>						
N 208611	001 7635773	Mar 13, 2029	DP		NCE	Jun 19, 2022
	7728143	Jun 19, 2031	DS		GAIN	Jun 19, 2027
	8252813	Oct 02, 2026	DP U-2028			
	8273892	Aug 06, 2026	DS			
	8410077	Mar 13, 2029	DP			
	8648093	Oct 07, 2025	DP U-2028			
	8871938	Sep 23, 2029	DS			
	9200088	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
	9539250	Oct 07, 2025	DS DP U-2028			
	9750822	Mar 13, 2029	DP			
	RE46617	Dec 28, 2029	DS			
<u>DEOXYCHOLIC ACID - KYBELLA</u>						
N 206333	001 10500214	Mar 02, 2030	DP			
	7622130	Dec 10, 2027	U-1690			
	7754230	Dec 10, 2027	U-1690			
	8101593	Mar 02, 2030	DP			
	8242294	May 16, 2028	DS			
	8298556	Aug 03, 2025	U-1690			
	8367649	Mar 02, 2030	DP			
	8461140	Feb 21, 2028	DP			
	8546367	Feb 21, 2028	DP U-1690			
	8653058	Mar 02, 2030	DP			
	8846066	Feb 08, 2025	U-1690			
	8883770	Feb 21, 2028	DP			
	9522155	Feb 21, 2028	DP U-1940			
	9636349	Feb 21, 2028	U-1940			
	9949986	Feb 21, 2028	U-1940			
<u>DESMOPRESSIN ACETATE - NOCDURNA</u>						
N 022517	001 11020448	May 21, 2029	U-2327			
	7560429	Feb 02, 2024	DP U-2326			
	9974826	Apr 13, 2030	U-2326			
<u>DESMOPRESSIN ACETATE - NOCDURNA</u>						
N 022517	002 10137167	May 21, 2029	U-2327			
	7560429	Feb 02, 2024	DP U-2326			
	9974826	Apr 13, 2030	U-2327			
<u>DESMOPRESSIN ACETATE - NOCTIVA</u>						
N 201656	001 11419914	Jun 15, 2030	U-3431			
	7799761	Sep 26, 2024	DP			
	9539302	Jun 15, 2030	DP			
<u>DESMOPRESSIN ACETATE - NOCTIVA</u>						
N 201656	002 11419914	Jun 15, 2030	U-3431			
	9539302	Jun 15, 2030	DP			

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<u>DESONIDE - VERDESO</u>						
N 021978	001 8460641	Aug 13, 2027	DP U-1412			
	8962000	Aug 31, 2025	DP U-1412			
	9492384	Aug 31, 2025	DP U-1412			
<u>DESOXIMETASONE - TOPICORT</u>						
N 204141	001 8277780	Sep 01, 2028	DP U-1408			
	8715624	May 26, 2026	DP U-1408			
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>						
N 021992	001 8269040	Jul 05, 2027	DS			
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>						
N 021992	002 8269040	Jul 05, 2027	DS			
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>						
N 021992	003 8269040	Jul 05, 2027	DS			
<u>DEUCRAVACITINIB - SOTYKTU</u>						
N 214958	001 10000480	Nov 07, 2033	DS DP		NCE	Sep 09, 2027
	11021475	Nov 07, 2033	U-3434			
	RE47929	Nov 07, 2033	DS DP U-3434			
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082	001 10959996	Mar 07, 2036	U-3055		M-54	Jun 24, 2024
	10959996*PED	Sep 07, 2036			ODE-134	Apr 03, 2024
	11179386	Mar 15, 2038	DP U-1995		PED	Oct 03, 2024
	11179386	Mar 15, 2038	DP U-3055		PED	Dec 24, 2024
	11179386*PED	Sep 15, 2038				
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11666566	Sep 18, 2033	DP			
	11666566*PED	Mar 18, 2034				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9233959	Sep 18, 2033	DP			
	9233959*PED	Mar 18, 2034				
	9296739	Sep 18, 2033	DP			
	9296739*PED	Mar 18, 2034				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
	9814708	Sep 18, 2033	DP			
	9814708*PED	Mar 18, 2034				

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<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 002	10959996	Mar 07, 2036	U-3055		M-54	Jun 24, 2024
	10959996*PED	Sep 07, 2036			ODE-134	Apr 03, 2024
	11179386	Mar 15, 2038	DP U-1995		PED	Oct 03, 2024
	11179386	Mar 15, 2038	DP U-3055		PED	Dec 24, 2024
	11179386*PED	Sep 15, 2038				
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11666566	Sep 18, 2033	DP			
	11666566*PED	Mar 18, 2034				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9233959	Sep 18, 2033	DP			
	9233959*PED	Mar 18, 2034				
	9296739	Sep 18, 2033	DP			
	9296739*PED	Mar 18, 2034				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
	9814708	Sep 18, 2033	DP			
	9814708*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082 003	10959996	Mar 07, 2036	U-3055		M-54	Jun 24, 2024
	10959996*PED	Sep 07, 2036			ODE-134	Apr 03, 2024
	11179386	Mar 15, 2038	DP U-1995		PED	Oct 03, 2024
	11179386	Mar 15, 2038	DP U-3055		PED	Dec 24, 2024
	11179386*PED	Sep 15, 2038				
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11666566	Sep 18, 2033	DP			
	11666566*PED	Mar 18, 2034				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	8524733	Apr 03, 2031	DS DP			

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<u>DEUTETRABENAZINE - AUSTEDO</u>						
N 208082	003 8524733*PED	Oct 03, 2031				
	9233959	Sep 18, 2033	DP			
	9233959*PED	Mar 18, 2034				
	9296739	Sep 18, 2033	DP			
	9296739*PED	Mar 18, 2034				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
	9814708	Sep 18, 2033	DP			
	9814708*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354	001 10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354	002 10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				

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<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 002	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEUTETRABENAZINE - AUSTEDO XR</u>						
N 216354 003	10959996	Mar 07, 2036	U-3055			
	10959996*PED	Sep 07, 2036				
	11179386	Mar 15, 2038	DP U-1995			
	11179386	Mar 15, 2038	DP U-3055			
	11179386*PED	Sep 15, 2038				
	11311488	Jun 10, 2041	DP U-1995			
	11311488	Jun 10, 2041	DP U-3055			
	11357772	Mar 07, 2036	U-1995			
	11357772	Mar 07, 2036	U-3055			
	11357772*PED	Sep 07, 2036				
	11446291	Mar 07, 2036	U-1995			
	11446291*PED	Sep 07, 2036				
	11564917	Mar 07, 2036	U-1995			
	11564917*PED	Sep 07, 2036				
	11648244	Mar 07, 2036	U-1995			
	11648244*PED	Sep 07, 2036				
	11813232	Mar 15, 2038	DP			
	11813232*PED	Sep 15, 2038				
	8524733	Apr 03, 2031	DS DP			
	8524733*PED	Oct 03, 2031				
	9550780	Sep 18, 2033	DS DP U-1995			
	9550780	Sep 18, 2033	DS DP U-3055			
	9550780*PED	Mar 18, 2034				
<u>DEXAMETHASONE - DEXAMETHASONE</u>						
A 217695 003					CGT	Apr 29, 2024
<u>DEXAMETHASONE - DEXTENZA</u>						
N 208742 001	11458041	Nov 16, 2037	U-1680		I-876	Oct 07, 2024
	11458041	Nov 16, 2037	U-3455			
	8409606	May 14, 2030	DP			
	8563027	Feb 12, 2030	U-2487			
	9254267	Sep 11, 2024	DP			
<u>DEXAMETHASONE - DEXYCU KIT</u>						
N 208912 001	10022502	Jun 22, 2034	U-2340			
	10028965	May 23, 2034	U-2340			
	10159683	May 23, 2034	DP			
	10799642	May 11, 2032	DP			
	11097061	Jun 23, 2039	DP U-3418			

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<u>DEXAMETHASONE - HEMADY</u>						
N 211379	001 10537585	Dec 18, 2037	DP		ODE-271	Oct 03, 2026
	11304961	Dec 18, 2037	DP U-3344			
<u>DEXAMETHASONE; TOBRAMYCIN - TOBRADEX ST</u>						
N 050818	001 7795316	Aug 03, 2028	DP U-1082			
	8101582	Dec 19, 2027	DP U-1082			
	8450287	Dec 19, 2027	DP			
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N 022287	001 7790755	Aug 02, 2026	DP			
	8105626	Sep 27, 2026	DP			
	8173158	Mar 17, 2030	U-949			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-951			
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8722084*PED	Apr 15, 2024				
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028	DP			
	9011926	Feb 24, 2026	DP			
	9233103	Mar 05, 2032	U-1805			
	9238029	Jan 17, 2026	DP			
<u>DEXLANSOPRAZOLE - DEXILANT</u>						
N 022287	002 7790755	Aug 02, 2026	DP			
	8105626	Sep 27, 2026	DP			
	8173158	Mar 17, 2030	U-949			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-951			
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8722084*PED	Apr 15, 2024				
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028	DP			
	9011926	Feb 24, 2026	DP			
	9233103	Mar 05, 2032	U-1805			
	9238029	Jan 17, 2026	DP			
<u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u>						
N 208056	001 8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028	DP			
	8871273*PED	Jul 11, 2028				
	9011926	Feb 24, 2026	DP			
	9238029	Jan 17, 2026	DP			
	9241910	Mar 10, 2029	DP			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038	001				NPP	Dec 16, 2025

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<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038 002	10016396	Jan 04, 2032	DP		NPP	Dec 16, 2025
	8242158	Jan 04, 2032	DP			
	8242158*PED	Jul 04, 2032				
	8338470	Jan 04, 2032	DP			
	8338470*PED	Jul 04, 2032				
	8455527	Jan 04, 2032		U-421		
	8455527*PED	Jul 04, 2032				
	8648106	Jan 04, 2032	DP			
	8648106*PED	Jul 04, 2032				
	9320712	Jan 04, 2032	DP			
	9320712*PED	Jul 04, 2032				
	9616049	Jan 04, 2032	DP			
	9616049*PED	Jul 04, 2032				
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038 003	10016396	Jan 04, 2032	DP		NPP	Dec 16, 2025
	8242158	Jan 04, 2032	DP			
	8242158*PED	Jul 04, 2032				
	8338470	Jan 04, 2032	DP			
	8338470*PED	Jul 04, 2032				
	8455527	Jan 04, 2032		U-421		
	8455527*PED	Jul 04, 2032				
	8648106	Jan 04, 2032	DP			
	8648106*PED	Jul 04, 2032				
	9320712	Jan 04, 2032	DP			
	9320712*PED	Jul 04, 2032				
	9616049	Jan 04, 2032	DP			
	9616049*PED	Jul 04, 2032				
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038 004	8242158	Jan 04, 2032	DP		NPP	Dec 16, 2025
	8242158*PED	Jul 04, 2032				
	8338470	Jan 04, 2032	DP			
	8338470*PED	Jul 04, 2032				
	8455527	Jan 04, 2032		U-421		
	8455527*PED	Jul 04, 2032				
	8648106	Jan 04, 2032	DP			
	8648106*PED	Jul 04, 2032				
	9320712	Jan 04, 2032	DP			
	9320712*PED	Jul 04, 2032				
	9616049	Jan 04, 2032	DP			
	9616049*PED	Jul 04, 2032				
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>						
N 021038 005					NPP	Dec 16, 2025
<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
N 206628 003	9649296	Apr 20, 2036	DP			
	9717796	Apr 20, 2036	DP			
<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
N 206628 004	9649296	Apr 20, 2036	DP			
	9717796	Apr 20, 2036	DP			

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<u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u>						
N 206628	004	9649296	Apr 20, 2036	DP		
		9717796	Apr 20, 2036	DP		
<u>DEXMEDETOMIDINE HYDROCHLORIDE - IGALMI</u>						
N 215390	001	10792246	Jun 26, 2039	DP U-3350	NP	Apr 05, 2025
		11478422	Jun 26, 2039	DP		
		11497711	Jun 26, 2039	DP U-3645		
		11517524	Jun 26, 2039	DP U-3645		
		11786508	Dec 29, 2037	U-3698		
		11806334	Jan 12, 2043	U-3725		
		11839604	Dec 29, 2037	U-3756		
<u>DEXMEDETOMIDINE HYDROCHLORIDE - IGALMI</u>						
N 215390	002	10792246	Jun 26, 2039	DP U-3350	NP	Apr 05, 2025
		11478422	Jun 26, 2039	DP		
		11497711	Jun 26, 2039	DP U-3645		
		11517524	Jun 26, 2039	DP U-3645		
		11786508	Dec 29, 2037	U-3698		
		11806334	Jan 12, 2043	U-3725		
		11839604	Dec 29, 2037	U-3756		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994	001	10584112	Dec 09, 2037	DS DP	NCE	May 07, 2026
		10584113	Dec 09, 2037	DP		
		10759778	Dec 09, 2037	DP		
		10858341	Dec 09, 2037	U-3094		
		10954213	Dec 09, 2037	U-3094		
		9079928	Jul 27, 2032	DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994	002	10584112	Dec 09, 2037	DS DP	NCE	May 07, 2026
		10584113	Dec 09, 2037	DP		
		10759778	Dec 09, 2037	DP		
		10858341	Dec 09, 2037	U-3094		
		10954213	Dec 09, 2037	U-3094		
		9079928	Jul 27, 2032	DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; SERDEXMETHYLPHENIDATE CHLORIDE - AZSTARYS</u>						
N 212994	003	10584112	Dec 09, 2037	DS DP	NCE	May 07, 2026
		10584113	Dec 09, 2037	DP		
		10759778	Dec 09, 2037	DP		
		10858341	Dec 09, 2037	U-3094		
		10954213	Dec 09, 2037	U-3094		
		9079928	Jul 27, 2032	DP		
<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401	001	11559501	Jan 06, 2042	DP	NP	Mar 22, 2025
		8591941	Oct 07, 2025	DP		
		8632802	Oct 07, 2025	DP		
		9034370	Oct 07, 2025	DP		
		9456993	Oct 24, 2033	DP U-3340		
		9474722	Oct 24, 2033	DP		

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<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401 002	11559501	Jan 06, 2042	DP		NP	Mar 22, 2025
	8591941	Oct 07, 2025	DP			
	8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9456993	Oct 24, 2033	DP U-3340			
	9474722	Oct 24, 2033	DP			
<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401 003	11559501	Jan 06, 2042	DP		NP	Mar 22, 2025
	8591941	Oct 07, 2025	DP			
	8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9456993	Oct 24, 2033	DP U-3340			
	9474722	Oct 24, 2033	DP			
<u>DEXTROAMPHETAMINE - XELSTRYM</u>						
N 215401 004	11559501	Jan 06, 2042	DP		NP	Mar 22, 2025
	8591941	Oct 07, 2025	DP			
	8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9456993	Oct 24, 2033	DP U-3340			
	9474722	Oct 24, 2033	DP			
<u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u>						
N 021879 001	7659282	Aug 13, 2026		U-1093		
<u>DIAZEPAM - VALTOCO</u>						
N 211635 001	10265402	May 11, 2025	DP		ODE-279	Jan 10, 2027
	11241414	Mar 27, 2029	DP			
	11793786	Mar 27, 2029	DP			
	8895546	Mar 27, 2029	DP			
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 002	10265402	May 11, 2025	DP		ODE-279	Jan 10, 2027
	11241414	Mar 27, 2029	DP			
	11793786	Mar 27, 2029	DP			
	8895546	Mar 27, 2029	DP			
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
<u>DIAZEPAM - VALTOCO</u>						
N 211635 003	10265402	May 11, 2025	DP		ODE-279	Jan 10, 2027
	11241414	Mar 27, 2029	DP			
	11793786	Mar 27, 2029	DP			
	8895546	Mar 27, 2029	DP			
	8927497	Jul 21, 2025	DP U-2727			
	9642913	May 11, 2025	DP			
<u>DICLOFENAC - ZORVOLEX</u>						
N 204592 001	8679544	Apr 23, 2030	DP			
	8999387	Apr 23, 2030		U-55		
	9017721	Apr 23, 2030	DP			

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<u>DICLOFENAC - ZORVOLEX</u>						
N 204592	001	9173854	Apr 23, 2030	DP		
		9180095	Apr 23, 2030	U-55		
		9180096	Apr 23, 2030	DP		
		9186328	Apr 23, 2030	U-55		
<u>DICLOFENAC - ZORVOLEX</u>						
N 204592	002	8679544	Apr 23, 2030	DP		
		8999387	Apr 23, 2030	U-55		
		9017721	Apr 23, 2030	DP		
		9173854	Apr 23, 2030	DP		
		9180095	Apr 23, 2030	U-55		
		9180096	Apr 23, 2030	DP		
		9186328	Apr 23, 2030	U-55		
<u>DICLOFENAC EPOLAMINE - LICART</u>						
N 206976	001	11344520	Feb 20, 2035	U-3393		
		11351133	Feb 20, 2035	U-3393		
<u>DICLOFENAC POTASSIUM - CAMBIA</u>						
N 022165	001	7759394	Jun 16, 2026	DS DP U-436		
		8097651	Jun 16, 2026	DS DP U-436		
		8927604	Jun 16, 2026	U-436		
		9827197	Jun 16, 2026	DP		
<u>DICLOFENAC POTASSIUM - ZIPSOR</u>						
N 022202	001	7662858	Feb 24, 2029	U-1035	NPP	May 25, 2024
		7884095	Feb 24, 2029	U-1111		
		7939518	Feb 24, 2029	U-980		
		8110606	Feb 24, 2029	U-980		
		8623920	Feb 24, 2029	U-1482		
		9561200	Feb 24, 2029	U-1482		
<u>DICLOFENAC SODIUM - PENNSAID</u>						
N 020947	001	8217078	Jul 10, 2029	U-1248		
		8546450	Aug 09, 2030	U-1435		
		8546450	Aug 09, 2030	U-1436		
		8618164	Jul 10, 2029	U-1477		
		8741956	Jul 10, 2029	U-1435		
<u>DICLOFENAC SODIUM - DYLOJECT</u>						
N 022396	001	8946292	Mar 22, 2027	U-1659		
<u>DICLOFENAC SODIUM - PENNSAID</u>						
N 204623	001	8217078	Jul 10, 2029	U-1477		
		8252838	Apr 21, 2028	DP U-1489		
		8546450	Aug 09, 2030	U-1435		
		8546450	Aug 09, 2030	U-1436		
		8563613	Oct 17, 2027	DP U-1488		
		8618164	Jul 10, 2029	U-1477		
		8741956	Jul 10, 2029	U-1435		
		8871809	Oct 17, 2027	U-1614		
		9066913	Oct 17, 2027	DP U-1488		
		9101591	Oct 17, 2027	DP U-1488		
		9132110	Oct 17, 2027	U-1488		

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<u>DICLOFENAC SODIUM - PENNSAID</u>						
N 204623	001 9168304	Oct 17, 2027	DP			
	9168305	Oct 17, 2027	U-1488			
	9220784	Oct 17, 2027	U-1488			
	9339551	Oct 17, 2027	U-1488			
	9339552	Oct 17, 2027	DP U-1488			
	9370501	Jul 10, 2029	U-1614			
	9375412	Jul 10, 2029	U-1614			
	9415029	Jul 10, 2029	U-1614			
	9539335	Oct 17, 2027	U-1614			
<u>DIENOGEST; DIENOGEST; DIENOGEST; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE - NATAZIA</u>						
N 022252	001 8071577	May 13, 2026	DP U-1			
	8153616	Jan 30, 2028	U-1240			
<u>DIFELIKEFALIN ACETATE - KORSUVA</u>						
N 214916	001 10017536	Nov 12, 2027	DS U-3204		NCE	Aug 23, 2026
	10138270	Nov 12, 2027	U-3204			
	10793596	Nov 12, 2027	DS DP U-3204			
	7402564	Nov 12, 2027	DS DP U-3204			
	7713937	Nov 12, 2027	DS DP U-3204			
	7727963	Nov 12, 2027	DS DP U-3204			
	8217007	Nov 12, 2027	U-3204			
	8236766	Nov 12, 2027	U-3204			
	8486894	Nov 12, 2027	U-3204			
	8536131	Nov 12, 2027	DS DP U-3204			
	9334305	Nov 12, 2027	U-3204			
	9359399	Nov 12, 2027	U-3204			
<u>DIHYDROERGOTAMINE MESYLATE - TRUDHESA</u>						
N 213436	001 10507295	Dec 25, 2032	DP		NP	Sep 02, 2024
	10940278	Jan 23, 2033	DP			
	11185497	Jan 04, 2039	U-3218			
	11266799	Nov 05, 2036	DP			
	9550036	Sep 05, 2032	DP			
	9919117	Mar 17, 2033	DP U-3218			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	001 10391160	Mar 13, 2035	U-3148			
	10555993	Mar 13, 2035	U-3148			
	10959972	Nov 16, 2035	U-1384			
	10994003	Mar 13, 2035	U-3148			
	11007166	Nov 16, 2035	U-1384			
	11007167	Nov 16, 2035	U-1384			
	11129806	Nov 16, 2035	U-1384			
	11246850	Nov 16, 2035	U-1384			
	8399514	Feb 07, 2028	U-1384			
<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	002 10391160	Mar 13, 2035	U-3148			
	10555993	Mar 13, 2035	U-3148			
	10959972	Nov 16, 2035	U-1384			
	10994003	Mar 13, 2035	U-3148			

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<u>DIMETHYL FUMARATE - TECFIDERA</u>						
N 204063	002	11007166				
		Nov 16, 2035				U-1384
		11007167				U-1384
		Nov 16, 2035				U-1384
		11129806				U-1384
		Nov 16, 2035				U-1384
		11246850				U-1384
		Nov 16, 2035				U-1384
		8399514				U-1384
		Feb 07, 2028				U-1384
<u>DIROXIMEL FUMARATE - VUMERITY</u>						
N 211855	001	10080733				
		Sep 20, 2033	DS DP			U-1384
		8669281				DS DP
		Sep 20, 2033				
		9090558				U-1384
		Sep 20, 2033				
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	001	10842770				
		Aug 07, 2031				DP U-2998
		8940786				DP U-1789
		Sep 30, 2033				
		9308195				DP
		Sep 30, 2033				
		9763880				U-2558
		Sep 30, 2033				
		9763880				U-2559
		Sep 30, 2033				
		9763880				U-2560
		Sep 30, 2033				
		9763880				U-2561
		Sep 30, 2033				
		9763880				U-2562
		Sep 30, 2033				
		9763880				U-2563
		Sep 30, 2033				
		9763880				U-2564
		Sep 30, 2033				
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	002	10842770				
		Aug 07, 2031				DP U-2998
		8940786				DP U-1789
		Sep 30, 2033				
		9308195				DP
		Sep 30, 2033				
		9763880				U-2558
		Sep 30, 2033				
		9763880				U-2559
		Sep 30, 2033				
		9763880				U-2560
		Sep 30, 2033				
		9763880				U-2561
		Sep 30, 2033				
		9763880				U-2562
		Sep 30, 2033				
		9763880				U-2563
		Sep 30, 2033				
		9763880				U-2564
		Sep 30, 2033				
<u>DOCETAXEL - DOCETAXEL</u>						
N 205934	003	10842770				
		Aug 07, 2031				DP U-2998
		8940786				DP U-1789
		Sep 30, 2033				
		9308195				DP
		Sep 30, 2033				
		9763880				U-2558
		Sep 30, 2033				
		9763880				U-2559
		Sep 30, 2033				
		9763880				U-2560
		Sep 30, 2033				
		9763880				U-2561
		Sep 30, 2033				
		9763880				U-2562
		Sep 30, 2033				
		9763880				U-2563
		Sep 30, 2033				
		9763880				U-2564
		Sep 30, 2033				
<u>DOCETAXEL - DOCETAXEL</u>						
N 215813	001	10398785				
		Mar 14, 2036				DP
<u>DOCETAXEL - DOCETAXEL</u>						
N 215813	002	10398785				
		Mar 14, 2036				DP

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<u>DOCETAXEL - DOCETAXEL</u>						
N 215813	003 10398785	Mar 14, 2036	DP			
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790	001 8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790	002 8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM - TIVICAY</u>						
N 204790	003 8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM - TIVICAY PD</u>						
N 213983	001 8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM; LAMIVUDINE - DOVATO</u>						
N 211994	001 11234985	Jan 24, 2031	DP U-257			
	8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE - JULUCA</u>						
N 210192	001 10426780	Jan 24, 2031	DS DP U-257			
	7125879	Apr 21, 2025	DS DP U-257			
	8129385	Oct 05, 2027	DS DP			
	8129385*PED	Apr 05, 2028				
	9242986	Dec 08, 2029	DS DP			
	9242986*PED	Jun 08, 2030				
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>						
N 022568	001 8481565	Oct 04, 2026	DP			
<u>DONEPEZIL HYDROCHLORIDE - ADLARITY</u>						
N 212304	001 10016372	Jul 26, 2037	U-3334		NP	Mar 11, 2025
	10300025	Jul 26, 2037	DP			
	10307379	Jul 26, 2037	DP			
	10835499	May 20, 2038	DP U-3334			
	10966936	Aug 11, 2038	DP U-3334			
	11103463	Jul 26, 2037	U-3334			
	11648214	Sep 23, 2037	DP U-3334			
	11679086	May 26, 2037	U-3334			
	9993466	Jul 26, 2037	DP			

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<u>DONEPEZIL HYDROCHLORIDE - ADLARITY</u>						
N 212304 002	10016372	Jul 26, 2037	U-3334		NP	Mar 11, 2025
	10300025	Jul 26, 2037	DP			
	10307379	Jul 26, 2037	DP			
	10835499	May 20, 2038	DP U-3334			
	10966936	Aug 11, 2038	DP U-3334			
	11103463	Jul 26, 2037	U-3334			
	11648214	Sep 23, 2037	DP U-3334			
	11679086	May 26, 2037	U-3334			
	9993466	Jul 26, 2037	DP			
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 001	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 002	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				

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<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 002	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 003	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439 004	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				

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<u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u>						
N 206439	004	8293794	Nov 22, 2025	DP		
		8329752	Nov 22, 2025	DP		
		8329752*PED	May 22, 2026			
		8338485	Nov 22, 2025	DP		
		8338486	Nov 22, 2025	U-1641		
		8362085	Nov 22, 2025	U-1641		
		8362085*PED	May 22, 2026			
		8580858	Nov 22, 2025	U-1641		
		8598233	Nov 22, 2025	DP		
		8598233*PED	May 22, 2026			
<u>DORAVIRINE - PIFELTRO</u>						
N 210806	001	8486975	Oct 07, 2031	DS DP U-2394	NPP	Jan 27, 2025
		8486975	Oct 07, 2031	DS DP U-2630		
		8486975	Oct 07, 2031	DS DP U-3308		
<u>DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE - DELSTRIGO</u>						
N 210807	001	10603282	Nov 29, 2036	DP	NPP	Jan 27, 2025
		10842751	Nov 29, 2036	DP		
		8486975	Oct 07, 2031	DS DP U-2395		
		8486975	Oct 07, 2031	DS DP U-2629		
		8486975	Oct 07, 2031	DS DP U-3307		
<u>DOXEPIN HYDROCHLORIDE - SILENOR</u>						
N 022036	001	10238620	May 18, 2027	U-620		
		10548871	Apr 11, 2028	U-620		
		10653660	Jul 20, 2027	U-620		
		10653662	May 18, 2027	U-620		
		11096920	Apr 11, 2028	U-620		
		11110074	Jul 20, 2027	U-620		
		11234954	Jan 18, 2028	U-620		
		7915307	Aug 24, 2027	U-620		
		8513299	Sep 07, 2030	U-620		
		9107898	May 01, 2028	U-620		
		9486437	May 18, 2027	U-620		
		9532971	Jun 01, 2029	DP		
		9572814	Jul 20, 2027	U-620		
		9861607	May 18, 2027	U-620		
		9907780	Apr 11, 2028	DP		
<u>DOXEPIN HYDROCHLORIDE - SILENOR</u>						
N 022036	002	10238620	May 18, 2027	U-620		
		10548871	Apr 11, 2028	U-620		
		10653660	Jul 20, 2027	U-620		
		10653662	May 18, 2027	U-620		
		11096920	Apr 11, 2028	U-620		
		11110074	Jul 20, 2027	U-620		
		11234954	Jan 18, 2028	U-620		
		7915307	Aug 24, 2027	U-620		
		8513299	Sep 07, 2030	U-620		
		9107898	May 01, 2028	U-620		
		9486437	May 18, 2027	U-620		

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<u>DOXEPIN HYDROCHLORIDE - SILENOR</u>						
N 022036	002 9532971	Jun 01, 2029	DP			
	9572814	Jul 20, 2027	U-620			
	9861607	May 18, 2027	U-620			
	9907780	Apr 11, 2028	DP			
<u>DOXYCYCLINE - ORACEA</u>						
N 050805	001 7749532	Dec 19, 2027	DP U-1063			
	8206740	Dec 24, 2025	DP U-925			
	8394405	Apr 07, 2024	DP U-925			
	8394406	Apr 07, 2024	DP U-925			
	8470364	Apr 07, 2024	DP U-925			
	8709478	Apr 07, 2024	U-1063			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	001 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	002 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	003 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	004 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	005 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX</u>						
N 050795	006 8715724	Feb 03, 2028	DP			
<u>DOXYCYCLINE HYCLATE - DORYX MPC</u>						
N 050795	007 8715724	Feb 03, 2028	DP			
	9295652	Oct 23, 2034	DP U-918			
	9446057	Dec 23, 2034	DP U-918			
	9511031	Oct 23, 2034	DP			
<u>DOXYCYCLINE HYCLATE - DORYX MPC</u>						
N 050795	008 8715724	Feb 03, 2028	DP			
	9295652	Oct 23, 2034	DP U-918			
	9446057	Dec 23, 2034	DP U-918			
	9511031	Oct 23, 2034	DP			
<u>DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE - BONJESTA</u>						
N 209661	001 9089489	Feb 18, 2033	DP U-1382			
	9375404	Feb 18, 2033	DP U-1382			
	9526703	Feb 18, 2033	DP U-1382			
	9937132	Feb 18, 2033	DP U-1382			
<u>DRONABINOL - SYNDROS</u>						
N 205525	001 10265293	Aug 06, 2028	DS DP			
	11253472	Aug 06, 2028	DS DP			
	8222292	Aug 06, 2028	DS DP			
	9345771	Aug 06, 2028	DS DP			

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<u>DRONEDARONE HYDROCHLORIDE - MULTAQ</u>						
N 022425	001 8410167	Apr 16, 2029	U-1387			
	8410167	Apr 16, 2029	U-1388			
	8602215	Jun 30, 2031	U-1473			
	9107900	Apr 16, 2029	U-1726			
	9107900	Apr 16, 2029	U-1728			
<u>DROSPIRENONE - SLYND</u>						
N 211367	001 10179140	Jun 28, 2031	U-2553			
	10603281	Jun 28, 2031	U-2553			
	10849857	Jun 28, 2031	DP U-2553			
	10987364	Jun 28, 2031	DP			
	11123299	Jun 28, 2031	DP			
	11291632	Jun 28, 2031	DP			
	11291633	Jun 28, 2031	DP			
	11351122	Jun 28, 2031	DP			
	11413249	Jun 28, 2031	U-2553			
	11439598	Jun 28, 2031	DP			
	11478487	Jun 28, 2031	DP			
	11504334	Jun 28, 2031	DP			
	9603860	Jun 28, 2031	U-2553			
<u>DROSPIRENONE - DROSPIRENONE</u>						
N 216285	001 10179140	Jun 28, 2031	U-2553		NP	Jun 29, 2025
	10603281	Jun 28, 2031	U-2553			
	10849857	Jun 28, 2031	DP U-2553			
	10987364	Jun 28, 2031	DP			
	11123299	Jun 28, 2031	DP			
	11291632	Jun 28, 2031	DP			
	11291633	Jun 28, 2031	DP			
	11351122	Jun 28, 2031	DP			
	11413249	Jun 28, 2031	U-2553			
	11439598	Jun 28, 2031	DP			
	11478487	Jun 28, 2031	DP			
	11504334	Jun 28, 2031	DP			
	9603860	Jun 28, 2031	U-2553			
<u>DROSPIRENONE; ESTETROL - NEXTSTELLIS</u>						
N 214154	001 11793760	Jun 17, 2036	DP		NCE	Apr 15, 2026
	7732430	Mar 02, 2025	DP U-3152			
<u>DROSPIRENONE; ESTRADIOL - ANGELIO</u>						
N 021355	001 8906890	Oct 22, 2031	DP			
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - BEYAZ</u>						
N 022532	001 11617751	Jul 17, 2030	DP U-1			
	11617751	Jul 17, 2030	DP U-3572			
	11617751	Jul 17, 2030	DP U-3573			
	11617751	Jul 17, 2030	DP U-3574			
	8617597	Feb 08, 2030	DP			
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - SAFYRAL</u>						
N 022574	001 11617751	Jul 17, 2030	DP U-1			
	11617751	Jul 17, 2030	DP U-3572			

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<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - SAFYRAL</u>						
N 022574	001 8617597	Feb 08, 2030	DP			
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	001 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	11202772	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	002 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	11202772	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	003 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	11202772	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<u>DULOXETINE HYDROCHLORIDE - DRIZALMA SPRINKLE</u>						
N 212516	004 10413525	Apr 13, 2037	DP			
	10959982	Apr 13, 2037	DP			
	11202772	Apr 13, 2037	DP			
	9839626	Apr 13, 2037	DP			
<u>DURLOBACTAM SODIUM; DURLOBACTAM SODIUM; SULBACTAM SODIUM - XACDURO (COPACKAGED)</u>						
N 216974	001 10376499	Nov 17, 2035	DP U-2840		NCE	May 23, 2028
	9309245	Apr 02, 2033	DS		GAIN	May 23, 2033
	9623014	Apr 02, 2033		U-2840		
	9968593	Nov 17, 2035	DP U-2840			
<u>DUVELISIB - COPIKTRA</u>						
N 211155	001 11312718	Jan 10, 2032	DP		ODE-208	Sep 24, 2025
	8193182	Feb 13, 2030	DS			
	9216982	Jan 05, 2029		U-2412		
	9216982	Jan 05, 2029		U-2413		
	9840505	Jan 10, 2032		U-2412		
	9840505	Jan 10, 2032		U-2413		
	RE46621	May 17, 2032	DS DP			
<u>DUVELISIB - COPIKTRA</u>						
N 211155	002 11312718	Jan 10, 2032	DP		ODE-208	Sep 24, 2025
	8193182	Feb 13, 2030	DS			
	9216982	Jan 05, 2029		U-2412		
	9216982	Jan 05, 2029		U-2413		
	9840505	Jan 10, 2032		U-2412		
	9840505	Jan 10, 2032		U-2413		
	RE46621	May 17, 2032	DS DP			
<u>ECONAZOLE NITRATE - ECOZA</u>						
N 205175	001 10071054	Aug 08, 2031	DP			

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<u>EDARAVONE - RADICAVA</u>						
N 209176	001				ODE-144	May 05, 2024
<u>EDARAVONE - RADICAVA</u>						
N 209176	002				ODE*	May 05, 2024
<u>EDARAVONE - RADICAVA ORS</u>						
N 215446	001	10987341	Nov 01, 2039	DP	NP	May 12, 2025
		11241416	Nov 01, 2039	DP		
		11478450	Nov 01, 2039	U-3468		
		11826352	Nov 01, 2039	DP		
<u>EDETATE CALCIUM DISODIUM - EDETATE CALCIUM DISODIUM</u>						
A 216435	001				CGT	Jan 08, 2024
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	001	7365205	Apr 18, 2027	DS		
		9149532	Mar 28, 2028	DP		
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	002	7365205	Apr 18, 2027	DS		
		9149532	Mar 28, 2028	DP		
<u>EDOXABAN TOSYLATE - SAVAYSA</u>						
N 206316	003	7365205	Apr 18, 2027	DS		
		9149532	Mar 28, 2028	DP		
<u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u>						
N 021937	001	8592397	Jan 13, 2024	DP U-1170		
		8592397	Jan 13, 2024	DP U-750		
		8598185	Apr 28, 2029	DP		
		8716264	Jan 13, 2024	DP U-257		
		9018192	Jun 13, 2026	U-1170		
		9018192	Jun 13, 2026	U-750		
		9457036	Jan 13, 2024	DP U-257		
		9545414	Jun 13, 2026	DP U-1170		
		9545414	Jun 13, 2026	DP U-750		
		9744181	Jan 13, 2024	DP U-257		
<u>EFINACONAZOLE - JUBLIA</u>						
N 203567	001	10105444	Jul 08, 2030	DP		
		10342875	Oct 02, 2034	DP U-2720		
		10478601	Apr 25, 2035	DP U-2721		
		10512640	Jan 03, 2028	U-1969		
		10828293	Oct 02, 2034	U-2720		
		10828369	Jan 03, 2028	DP		
		10864274	Oct 02, 2034	U-2720		
		11213519	Jan 03, 2028	U-2720		
		11654139	Oct 02, 2034	U-1969		
		7214506	Feb 22, 2026	U-281		
		8039494	Jul 08, 2030	U-281		
		8486978	Oct 24, 2030	DP		
		9302009	Oct 24, 2030	DP		
		9566272	Jan 03, 2028	U-1969		
		9662394	Oct 02, 2034	DP		

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<u>EFINACONAZOLE - JUBLIA</u>						
N 203567	001 9861698	Jul 08, 2030	DP			
	9877955	Jan 03, 2028	U-1969			
<u>ELACESTRANT DIHYDROCHLORIDE - ORSERDU</u>						
N 217639	001 10071066	Oct 10, 2034	U-3524		NCE	Jan 27, 2028
	10385008	Jan 05, 2038	DS DP			
	10420734	Oct 10, 2034	U-3524			
	10745343	Jan 05, 2038	U-3523			
	11779552	Oct 10, 2034	U-3524			
	11819480	Nov 29, 2036	U-3523			
	7612114	Aug 18, 2026	DS DP U-3523			
	8399520	Dec 25, 2024	DS DP U-3523			
<u>ELACESTRANT DIHYDROCHLORIDE - ORSERDU</u>						
N 217639	002 10071066	Oct 10, 2034	U-3524		NCE	Jan 27, 2028
	10385008	Jan 05, 2038	DS DP			
	10420734	Oct 10, 2034	U-3524			
	10745343	Jan 05, 2038	U-3523			
	11779552	Oct 10, 2034	U-3524			
	11819480	Nov 29, 2036	U-3523			
	7612114	Aug 18, 2026	DS DP U-3523			
	8399520	Dec 25, 2024	DS DP U-3523			
<u>ELAGOLIX SODIUM - ORLISSA</u>						
N 210450	001 10537572	Sep 01, 2036	U-2735			
	10682351	Sep 01, 2036	U-2850			
	11542239	Jul 23, 2039	DS DP			
	11690845	Aug 27, 2040	U-3654			
	11690854	Apr 19, 2038	U-3653			
	11707464	Mar 14, 2034	U-3672			
	7056927	Sep 10, 2024	DS DP			
	7176211	Jul 06, 2024	U-2360			
	7419983	Jul 06, 2024	DS DP U-2360			
<u>ELAGOLIX SODIUM - ORLISSA</u>						
N 210450	002 11344551	Mar 14, 2034	U-3388			
	11344551	Mar 14, 2034	U-3389			
	11542239	Jul 23, 2039	DS DP			
	11690845	Aug 27, 2040	U-3654			
	7056927	Sep 10, 2024	DS DP			
	7176211	Jul 06, 2024	U-2360			
	7419983	Jul 06, 2024	DS DP U-2360			
<u>ELAGOLIX SODIUM, ESTRADIOL, NORETHINDRONE ACETATE; ELAGOLIX SODIUM - ORIAHNN (COPACKAGED)</u>						
N 213388	001 10881659	Mar 14, 2034	U-2842			
	11045470	Mar 14, 2034	U-2842			
	11459305	Nov 07, 2028	DP U-2842			
	11542239	Jul 23, 2039	DS DP			
	11690845	Aug 27, 2040	U-3655			
	7056927	Sep 10, 2024	DS DP			
	7419983	Jul 06, 2024	DS DP			

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<u>ELBASVIR; GRAZOPREXIVIR - ZEPATIER</u>						
N 208261	001 7973040	Jul 24, 2029	DS DP U-1813		NPP	Dec 09, 2024
	8871759	May 04, 2031	DS DP U-1813			
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273	001 10022352	Apr 09, 2027	DP U-2651		NCE	Oct 21, 2024
	10022352	Apr 09, 2027	DP U-3156		NPP	Jun 08, 2024
	10081621	Mar 25, 2031	DP U-2652		ODE-275	Oct 21, 2026
	10081621	Mar 25, 2031	DP U-3032		ODE-323	Dec 21, 2027
	10081621	Mar 25, 2031	DP U-3157		ODE-357	Jun 08, 2028
	10239867	Apr 09, 2027	DS DP U-2653			
	10239867	Apr 09, 2027	DS DP U-3033			
	10239867	Apr 09, 2027	DS DP U-3158			
	10646481	Aug 13, 2029	DP			
	10758534	Oct 06, 2035	DS DP U-2645			
	10758534	Oct 06, 2035	DS DP U-3028			
	10758534	Oct 06, 2035	DS DP U-3144			
	10793547	Dec 08, 2037	DS DP U-2645			
	10793547	Dec 08, 2037	DS DP U-3028			
	10793547	Dec 08, 2037	DS DP U-3144			
	11179367	Dec 08, 2037	DP U-3253			
	11426407	Oct 06, 2035	DS DP U-3425			
	11453655	Dec 08, 2037	DS DP			
	11517564	Dec 08, 2037	DP U-3498			
	11564916	Aug 13, 2029	U-3525			
	11578062	Mar 25, 2031	DP U-3544			
	11639347	Apr 09, 2027	DS DP U-3587			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-2645			
	8324242	Aug 05, 2027	U-3028			
	8324242	Aug 05, 2027	U-3144			
	8354427	Jul 06, 2026	U-2646			
	8354427	Jul 06, 2026	U-3029			
	8354427	Jul 06, 2026	U-3145			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-2645			
	8415387	Nov 12, 2027	U-3028			
	8415387	Nov 12, 2027	U-3144			
	8598181	May 01, 2027	U-2645			
	8598181	May 01, 2027	U-3028			
	8598181	May 01, 2027	U-3144			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-2648			
	8629162	Jun 24, 2025	U-3030			
	8629162	Jun 24, 2025	U-3146			
	8754224	Dec 28, 2026	DS DP			
	9012496	Jul 15, 2033	U-2649			
	9012496	Jul 15, 2033	U-3154			
	9670163	Dec 28, 2026	DP U-2650			

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<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273	001	9670163	Dec 28, 2026	DP U-3031		
		9670163	Dec 28, 2026	DP U-3155		
		9931334	Dec 28, 2026	DP U-2650		
		9931334	Dec 28, 2026	DP U-3031		
		9931334	Dec 28, 2026	DP U-3155		
		9974781	Apr 09, 2027	DP U-2645		
		9974781	Apr 09, 2027	DP U-3028		
		9974781	Apr 09, 2027	DP U-3144		
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 212273	002	10022352	Apr 09, 2027	DP U-3156	NCE	Oct 21, 2024
		10081621	Mar 25, 2031	DP U-3157	NPP	Jun 08, 2024
		10239867	Apr 09, 2027	DS DP U-3158	ODE-357	Jun 08, 2028
		10646481	Aug 13, 2029	DP		
		10758534	Oct 06, 2035	DS DP U-3144		
		10793547	Dec 08, 2037	DS DP U-3144		
		11179367	Dec 08, 2037	DP U-3253		
		11426407	Oct 06, 2035	DS DP U-3425		
		11453655	Dec 08, 2037	DS DP		
		11517564	Dec 08, 2037	DP U-3498		
		11564916	Aug 13, 2029	U-3525		
		11578062	Mar 25, 2031	DP U-3544		
		11639347	Apr 09, 2027	DS DP U-3587		
		7495103	May 20, 2027	DS DP		
		7645789	May 01, 2027	DS DP		
		7776905	Jun 03, 2027	DS DP		
		8324242	Aug 05, 2027	U-3144		
		8354427	Jul 06, 2026	U-3145		
		8410274	Dec 28, 2026	DP		
		8415387	Nov 12, 2027	U-3144		
		8598181	May 01, 2027	U-3144		
		8623905	May 01, 2027	DS DP		
		8629162	Jun 24, 2025	U-3146		
		8754224	Dec 28, 2026	DS DP		
		9012496	Jul 15, 2033	U-3154		
		9670163	Dec 28, 2026	DP U-3155		
		9931334	Dec 28, 2026	DP U-3155		
		9974781	Apr 09, 2027	DP U-3144		
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 217660	001	10022352	Apr 09, 2027	DP U-3596	NCE	Oct 21, 2024
		10081621	Mar 25, 2031	DP U-3600	NP	Apr 26, 2026
		10239867	Apr 09, 2027	DS DP U-3590	ODE-433	Apr 26, 2030
		10272046	Feb 27, 2033	DP U-3599		
		10646481	Aug 13, 2029	DP		
		10758534	Oct 06, 2035	DS DP U-3589		
		10793547	Dec 08, 2037	DS DP U-3588		
		11147770	Feb 27, 2033	DP U-3598		
		11179367	Dec 08, 2037	DP U-3597		
		11426407	Oct 06, 2035	DS DP U-3595		
		11453655	Dec 08, 2037	DS DP		

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<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 217660 001	11517564	Dec 08, 2037	DP U-3586			
	11564916	Aug 13, 2029	U-3585			
	11578062	Mar 25, 2031	DP U-3584			
	11639347	Apr 09, 2027	DS DP U-3583			
	11752106	Feb 27, 2033	DP U-3696			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-3589			
	8354427	Jul 06, 2026	U-3593			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-3589			
	8598181	May 01, 2027	U-3589			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-3592			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-3591			
	9931334	Dec 28, 2026	DP U-3591			
	9974781	Apr 09, 2027	DP U-3589			
<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 217660 002	10022352	Apr 09, 2027	DP U-3596		NCE	Oct 21, 2024
	10081621	Mar 25, 2031	DP U-3600		NP	Apr 26, 2026
	10239867	Apr 09, 2027	DS DP U-3590		ODE-433	Apr 26, 2030
	10272046	Feb 27, 2033	DP U-3599			
	10646481	Aug 13, 2029	DP			
	10758534	Oct 06, 2035	DS DP U-3589			
	10793547	Dec 08, 2037	DS DP U-3588			
	11147770	Feb 27, 2033	DP U-3598			
	11179367	Dec 08, 2037	DP U-3597			
	11426407	Oct 06, 2035	DS DP U-3595			
	11453655	Dec 08, 2037	DS DP			
	11517564	Dec 08, 2037	DP U-3586			
	11564916	Aug 13, 2029	U-3585			
	11578062	Mar 25, 2031	DP U-3584			
	11639347	Apr 09, 2027	DS DP U-3583			
	11752106	Feb 27, 2033	DP U-3696			
	7495103	May 20, 2027	DS DP			
	7645789	May 01, 2027	DS DP			
	7776905	Jun 03, 2027	DS DP			
	8324242	Aug 05, 2027	U-3589			
	8354427	Jul 06, 2026	U-3593			
	8410274	Dec 28, 2026	DP			
	8415387	Nov 12, 2027	U-3589			
	8598181	May 01, 2027	U-3589			
	8623905	May 01, 2027	DS DP			
	8629162	Jun 24, 2025	U-3592			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			

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<u>ELEXACAFTOR, IVACAFTOR, TEZACAFTOR; IVACAFTOR - TRIKAFTA (COPACKAGED)</u>						
N 217660 002	9670163	Dec 28, 2026	DP U-3591			
	9931334	Dec 28, 2026	DP U-3591			
	9974781	Apr 09, 2027	DP U-3589			
<u>ELIGLUSTAT TARTRATE - CERDELGA</u>						
N 205494 001	10888544	Dec 13, 2038	U-3040			
	10888544	Dec 13, 2038	U-3041			
	10888547	Jan 31, 2031	U-3042			
	10888547	Jan 31, 2031	U-3043			
	11458119	Nov 24, 2030	DS DP			
	7196205	Jun 26, 2026	DS			
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 001	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP U-1306			
	8052993	Aug 01, 2027	DP U-1575			
	8052993	Aug 01, 2027	DP U-2451			
	8052993	Aug 01, 2027	DP U-930			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 002	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1306			
	8052994	Aug 01, 2027	DP U-2451			
	8052994	Aug 01, 2027	DP U-930			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			

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<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 002	8828430	Aug 01, 2027	DP U-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 003	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1306			
	8062665	Aug 01, 2027	DP U-2451			
	8062665	Aug 01, 2027	DP U-930			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 004	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-2451			
	7547719	Jul 13, 2025	DS DP U-2452			
	7547719*PED	Jan 13, 2026				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1306			
	8071129	Aug 01, 2027	DP U-2451			
	8071129	Aug 01, 2027	DP U-930			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-2451			
	8828430*PED	Feb 01, 2028				
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
N 022291 005	7547719	Jul 13, 2025	DS DP U-1306		ODE-210	Nov 16, 2025
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				

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<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027	001	7547719	Jul 13, 2025	DS DP U-1306	ODE*	Nov 16, 2025
		7547719	Jul 13, 2025	DS DP U-1575		
		7547719	Jul 13, 2025	DS DP U-1736		
		7547719	Jul 13, 2025	DS DP U-2452		
		7547719*PED	Jan 13, 2026			
<u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u>						
N 207027	002	7547719	Jul 13, 2025	DS DP U-1306	ODE*	Nov 16, 2025
		7547719	Jul 13, 2025	DS DP U-1575		
		7547719	Jul 13, 2025	DS DP U-1736		
		7547719	Jul 13, 2025	DS DP U-2452		
		7547719*PED	Jan 13, 2026			
<u>ELUXADOLINE - VIBERZI</u>						
N 206940	001	10188632	Mar 14, 2033	DP		
		10213415	Mar 14, 2025	DS U-2152		
		11007179	Mar 14, 2033	DP		
		11090291	Mar 14, 2033	DP		
		11160792	Mar 14, 2033	DP		
		11229627	Mar 14, 2033	DP		
		11311516	Mar 14, 2033	DP		
		7741356	May 27, 2029	DS DP		
		7786158	Mar 14, 2025	DS		
		8344011	Mar 14, 2025	U-1709		
		8609709	Mar 14, 2025	DS		
		8691860	Jul 07, 2028	DS U-1709		
		8772325	Mar 14, 2025	U-1709		
		9115091	Jul 07, 2028	DS DP U-1738		
		9205076	Mar 14, 2025	U-1709		
		9364489	Jul 07, 2028	U-1709		
		9675587	Mar 14, 2033	DP		
		9700542	Mar 14, 2025	DP		
		9789125	Jul 07, 2028	DP U-1709		
		9789125	Jul 07, 2028	DP U-2152		
<u>ELUXADOLINE - VIBERZI</u>						
N 206940	002	10188632	Mar 14, 2033	DP		
		10213415	Mar 14, 2025	DS U-2152		
		11007179	Mar 14, 2033	DP		
		11090291	Mar 14, 2033	DP		
		11160792	Mar 14, 2033	DP		
		11229627	Mar 14, 2033	DP		
		11311516	Mar 14, 2033	DP		
		11484527	Mar 14, 2033	U-3475		
		7741356	May 27, 2029	DS DP		
		7786158	Mar 14, 2025	DS		
		8344011	Mar 14, 2025	U-1709		
		8609709	Mar 14, 2025	DS		
		8691860	Jul 07, 2028	DS U-1709		
		8772325	Mar 14, 2025	U-1709		
		9115091	Jul 07, 2028	DS DP U-1738		

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<u>ELUXADOLINE - VIBERZI</u>						
N 206940 002	9205076	Mar 14, 2025	U-1709			
	9364489	Jul 07, 2028	U-1709			
	9675587	Mar 14, 2033	DP			
	9700542	Mar 14, 2025	DP			
	9789125	Jul 07, 2028	DP U-1709			
	9789125	Jul 07, 2028	DP U-2152			
<u>ELVITEGRAVIR - VITEKTA</u>						
N 203093 001	7176220	Aug 27, 2026	DS DP U-257			
	7176220*PED	Feb 27, 2027				
	7635704	Oct 26, 2026	DS DP U-257			
	7635704*PED	Apr 26, 2027				
	8981103	Oct 26, 2026	DS DP			
	8981103*PED	Apr 26, 2027				
<u>ELVITEGRAVIR - VITEKTA</u>						
N 203093 002	7176220	Aug 27, 2026	DS DP U-257			
	7176220*PED	Feb 27, 2027				
	7635704	Oct 26, 2026	DS DP U-257			
	7635704*PED	Apr 26, 2027				
	8981103	Oct 26, 2026	DS DP			
	8981103*PED	Apr 26, 2027				
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629 001	10258637	Apr 03, 2034	U-2290		I-869	Aug 18, 2024
	10258637*PED	Oct 03, 2034			I-922	Sep 21, 2026
	11090323	Apr 03, 2034	U-3191		M-82	Feb 24, 2025
	11090323*PED	Oct 03, 2034			NPP	Jun 20, 2026
	11666590	Apr 03, 2034	U-3691		PED	Aug 24, 2025
	11813275	Apr 03, 2034	U-3759		PED	Dec 20, 2026
	11813275	Apr 03, 2034	U-3760			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	U-1651			
	8551957*PED	Apr 14, 2030				
	9949997	May 17, 2034	U-2292			
	9949997	May 17, 2034	U-3199			
	9949997	May 17, 2034	U-3325			
	9949997*PED	Nov 17, 2034				
	9949998	Jun 11, 2034	U-2290			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629 002	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	11090323	Apr 03, 2034	U-3191			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN - JARDIANCE</u>						
N 204629 002	11833166	Apr 03, 2034	U-3777			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	U-1651			
	8551957*PED	Apr 14, 2030				
	9949997	May 17, 2034	U-2292			
	9949997*PED	Nov 17, 2034				
	9949998	Jun 11, 2034	U-2290			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 001	10258637	Apr 03, 2034	U-2290			
	10258637*PED	Oct 03, 2034				
	11033552	May 04, 2027	DP			
	11033552*PED	Nov 04, 2027				
	11090323	Apr 03, 2034	U-3191			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-1651			
	8551957*PED	Apr 14, 2030				
	8673927	May 04, 2027	U-1652		Y	
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9173859	May 04, 2027	DP U-1772		Y	
	9949998	Jun 11, 2034	U-2290			
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 002	10258637	Apr 03, 2034	U-2290			
	10258637*PED	Oct 03, 2034				
	11033552	May 04, 2027	DP			
	11033552*PED	Nov 04, 2027				
	11090323	Apr 03, 2034	U-3191			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u>						
N 206073 002	8551957	Oct 14, 2029	DP U-1651			
	8551957*PED	Apr 14, 2030				
	8673927	May 04, 2027	U-1652	Y		
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9173859	May 04, 2027	DP U-1772	Y		
	9949998	Jun 11, 2034	U-2290			
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 001	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11564886	Mar 07, 2032	DP U-3531			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-2730			
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 002	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11564886	Mar 07, 2032	DP U-3531			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			

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<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 002	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-2730			
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 003	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11564886	Mar 07, 2032	DP U-3531			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029	DP U-2730			
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034	U-2731			
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 004	10022379	Apr 02, 2029	DP U-2732			
	10258637	Apr 03, 2034	U-2731			
	10258637*PED	Oct 03, 2034				
	10406172	Jun 15, 2030	DP U-2733			
	10596120	Mar 07, 2032	DP U-2776			
	10596120	Mar 07, 2032	DP U-2790			
	11090323	Apr 03, 2034	U-3192			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7407955	May 02, 2025	DS DP			

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<u>EMPAGLIFLOZIN; LINAGLIPTIN; METFORMIN HYDROCHLORIDE - TRIJARDY XR</u>						
N 212614 004	7407955*PED	Nov 02, 2025				
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	8551957	Oct 14, 2029		DP U-2730		
	8551957*PED	Apr 14, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9415016	Apr 02, 2029	DP			
	9949998	Jun 11, 2034		U-2731		
	9949998*PED	Dec 11, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 001	10258637	Apr 03, 2034		U-2290	NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034		U-3193		
	11090323*PED	Oct 03, 2034				
	11813275	Apr 03, 2034		U-3759		
	11833166	Apr 03, 2034		U-3776		
	11833166	Apr 03, 2034		U-3777		
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034		U-3532		
	9949997	May 17, 2034		U-3533		
	9949997*PED	Nov 17, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 002	10258637	Apr 03, 2034		U-2290	NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034		U-3193		
	11090323*PED	Oct 03, 2034				
	11813275	Apr 03, 2034		U-3759		
	11833166	Apr 03, 2034		U-3776		
	11833166	Apr 03, 2034		U-3777		
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034		U-3532		
	9949997	May 17, 2034		U-3533		
	9949997*PED	Nov 17, 2034				

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<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 003	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u>						
N 206111 004	10258637	Apr 03, 2034	U-2290		NPP	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10610489	Sep 30, 2030	DP			
	10610489*PED	Mar 30, 2031				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658 001	10258637	Apr 03, 2034	U-2290		M-296	Jun 20, 2026
	10258637*PED	Oct 03, 2034			PED	Dec 20, 2026
	10596120	Mar 07, 2032	DP U-2775			
	10596120	Mar 07, 2032	DP U-2792			
	10596120*PED	Sep 07, 2032				
	11090323	Apr 03, 2034	U-3193			
	11090323*PED	Oct 03, 2034				
	11813275	Apr 03, 2034	U-3759			
	11833166	Apr 03, 2034	U-3776			
	11833166	Apr 03, 2034	U-3777			
	7579449	Aug 01, 2028	DS			
	7579449*PED	Feb 01, 2029				
	7713938	Apr 15, 2027	DS DP			
	7713938*PED	Oct 15, 2027				
	9949997	May 17, 2034	U-3532			
	9949997	May 17, 2034	U-3533			
	9949997*PED	Nov 17, 2034				

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<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	001	9949998	Jun 11, 2034	U-2290		
		9949998*PED	Dec 11, 2034			
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	002	10258637	Apr 03, 2034	U-2290	M-296	Jun 20, 2026
		10258637*PED	Oct 03, 2034		PED	Dec 20, 2026
		10596120	Mar 07, 2032	DP U-2775		
		10596120	Mar 07, 2032	DP U-2792		
		10596120*PED	Sep 07, 2032			
		11090323	Apr 03, 2034	U-3193		
		11090323*PED	Oct 03, 2034			
		11813275	Apr 03, 2034	U-3759		
		11833166	Apr 03, 2034	U-3776		
		11833166	Apr 03, 2034	U-3777		
		7579449	Aug 01, 2028	DS		
		7579449*PED	Feb 01, 2029			
		7713938	Apr 15, 2027	DS DP		
		7713938*PED	Oct 15, 2027			
		9949997	May 17, 2034	U-3532		
		9949997	May 17, 2034	U-3533		
		9949997*PED	Nov 17, 2034			
		9949998	Jun 11, 2034	U-2290		
		9949998*PED	Dec 11, 2034			
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	003	10258637	Apr 03, 2034	U-2290	M-296	Jun 20, 2026
		10258637*PED	Oct 03, 2034		PED	Dec 20, 2026
		10596120	Mar 07, 2032	DP U-2775		
		10596120	Mar 07, 2032	DP U-2792		
		10596120*PED	Sep 07, 2032			
		11090323	Apr 03, 2034	U-3193		
		11090323*PED	Oct 03, 2034			
		11833166	Apr 03, 2034	U-3776		
		11833166	Apr 03, 2034	U-3777		
		7579449	Aug 01, 2028	DS		
		7579449*PED	Feb 01, 2029			
		7713938	Apr 15, 2027	DS DP		
		7713938*PED	Oct 15, 2027			
		9949997	May 17, 2034	U-3532		
		9949997	May 17, 2034	U-3533		
		9949997*PED	Nov 17, 2034			
		9949998	Jun 11, 2034	U-2290		
		9949998*PED	Dec 11, 2034			
<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	004	10258637	Apr 03, 2034	U-2290	M-296	Jun 20, 2026
		10258637*PED	Oct 03, 2034		PED	Dec 20, 2026
		10596120	Mar 07, 2032	DP U-2775		
		10596120	Mar 07, 2032	DP U-2792		
		10596120*PED	Sep 07, 2032			
		11090323	Apr 03, 2034	U-3193		

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<u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u>						
N 208658	004	11090323*PED	Oct 03, 2034			
		11833166	Apr 03, 2034	U-3776		
		11833166	Apr 03, 2034	U-3777		
		7579449	Aug 01, 2028	DS		
		7579449*PED	Feb 01, 2029			
		7713938	Apr 15, 2027	DS DP		
		7713938*PED	Oct 15, 2027			
		9949997	May 17, 2034	U-3532		
		9949997	May 17, 2034	U-3533		
		9949997*PED	Nov 17, 2034			
		9949998	Jun 11, 2034	U-2290		
		9949998*PED	Dec 11, 2034			
<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY</u>						
N 208351	001	7125879	Apr 21, 2025	DS DP U-257		
		7390791	Apr 17, 2025	DS DP		
		7390791*PED	Oct 17, 2025			
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
<u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA</u>						
N 202123	001	10857102	Jan 14, 2033	DP		
		7125879	Apr 21, 2025	DS DP U-257		
		8592397	Jan 13, 2024	DP U-257		
		8716264	Jan 13, 2024	DP U-257		
		8841310	Dec 09, 2025	DP U-257		
		9457036	Jan 13, 2024	DP U-257		
		9744181	Jan 13, 2024	DP U-257		
<u>EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCOVY</u>						
N 208215	001	7390791	Apr 17, 2025	DS DP	ODE-284	Sep 28, 2024
		7390791*PED	Oct 17, 2025		ODE-285	Sep 28, 2024
		8754065	Aug 15, 2032	DS DP U-1259		
		8754065	Aug 15, 2032	DS DP U-1663		
		8754065	Aug 15, 2032	DS DP U-257		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-1259		
		9296769	Aug 15, 2032	DS DP U-1663		
		9296769	Aug 15, 2032	DS DP U-257		
		9296769*PED	Feb 15, 2033			
<u>EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCOVY</u>						
N 208215	002	7390791	Apr 17, 2025	DS DP		
		7390791*PED	Oct 17, 2025			
		8754065	Aug 15, 2032	DS DP U-1663		
		8754065*PED	Feb 15, 2033			
		9296769	Aug 15, 2032	DS DP U-1663		
		9296769*PED	Feb 15, 2033			

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<u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u>						
N 021752	001 8592397	Jan 13, 2024	DP U-1170			
	8592397	Jan 13, 2024	DP U-248			
	8592397	Jan 13, 2024	DP U-541			
	8716264	Jan 13, 2024	DP U-257			
	9457036	Jan 13, 2024	DP U-257			
	9744181	Jan 13, 2024	DP U-257			
<u>ENALAPRIL MALEATE - EPANED KIT</u>						
N 204308	001 8568747	Nov 06, 2032	DP			
	8778366	Nov 06, 2032	U-1723			
	8778366	Nov 06, 2032	U-185			
	8778366	Nov 06, 2032	U-1892			
	8778366	Nov 06, 2032	U-3			
	8778366	Nov 06, 2032	U-71			
	9855214	Nov 06, 2032	DP			
	9968553	Nov 06, 2032	U-1723			
	9968553	Nov 06, 2032	U-185			
	9968553	Nov 06, 2032	U-1892			
	9968553	Nov 06, 2032	U-3			
	9968553	Nov 06, 2032	U-71			
<u>ENALAPRIL MALEATE - EPANED</u>						
N 208686	001 10039745	Mar 25, 2036	DP			
	10154987	Mar 25, 2036	U-1723			
	10154987	Mar 25, 2036	U-185			
	10154987	Mar 25, 2036	U-1892			
	10154987	Mar 25, 2036	U-3			
	10154987	Mar 25, 2036	U-71			
	10772868	Mar 25, 2036	DP			
	10786482	Mar 25, 2036	DP			
	11040023	Mar 25, 2036	DP			
	11141405	Mar 25, 2036	DP			
	11173141	Mar 25, 2036	DP			
	9669008	Mar 25, 2036	DP			
	9808442	Mar 25, 2036	U-1723			
	9808442	Mar 25, 2036	U-185			
	9808442	Mar 25, 2036	U-1892			
	9808442	Mar 25, 2036	U-3			
	9808442	Mar 25, 2036	U-71			
<u>ENASIDENIB MESYLATE - IDHIFA</u>						
N 209606	001 10093654	Aug 01, 2034	DS DP U-2087		ODE-151	Aug 01, 2024
	10294215	Jan 07, 2033	DP U-2087			
	10610125	Jun 21, 2030	U-2087			
	9512107	Jan 07, 2033	DS DP U-2087			
	9732062	Sep 16, 2034	DS			
	9738625	Aug 01, 2034	DS			
<u>ENASIDENIB MESYLATE - IDHIFA</u>						
N 209606	002 10093654	Aug 01, 2034	DS DP U-2087		ODE-151	Aug 01, 2024
	10294215	Jan 07, 2033	DP U-2087			
	10610125	Jun 21, 2030	U-2087			

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<u>ENASIDENIB MESYLATE - IDHIFA</u>						
N 209606 002	9512107	Jan 07, 2033	DS DP U-2087			
	9732062	Sep 16, 2034	DS			
	9738625	Aug 01, 2034	DS			
<u>ENCORAFENIB - BRAFTOVI</u>						
N 210496 001	10005761	Aug 27, 2030	U-2335		I-928	Oct 11, 2026
	8501758	Mar 04, 2031	DS DP		ODE-194	Jun 27, 2025
	8541575	Feb 26, 2030	DS DP U-2335		ODE-445	Oct 11, 2030
	8946250	Jul 23, 2029	DS DP			
	9314464	Jul 04, 2031	U-2336			
	9387208	Nov 21, 2032	DP			
	9593099	Aug 27, 2030	DS			
	9593100	Aug 27, 2030	DP			
	9763941	Nov 21, 2032	U-2335			
	9850229	Aug 27, 2030	U-2337			
	9850230	Aug 27, 2030	U-2334			
<u>ENCORAFENIB - BRAFTOVI</u>						
N 210496 002	10005761	Aug 27, 2030	U-2335		I-928	Oct 11, 2026
	10005761	Aug 27, 2030	U-2802		ODE-194	Jun 27, 2025
	10005761	Aug 27, 2030	U-2803		ODE-445	Oct 11, 2030
	10005761	Aug 27, 2030	U-3738			
	10258622	Nov 21, 2032	U-2802			
	8501758	Aug 27, 2030	DS DP			
	8541575	Feb 26, 2030	DS DP U-2335			
	8541575	Feb 26, 2030	DS DP U-2802			
	8541575	Feb 26, 2030	DS DP U-2803			
	8541575	Feb 26, 2030	DS DP U-3738			
	8946250	Jul 23, 2029	DS DP			
	9314464	Jul 04, 2031	U-2336			
	9314464	Jul 04, 2031	U-2802			
	9314464	Jul 04, 2031	U-2803			
	9314464	Jul 04, 2031	U-3738			
	9387208	Nov 21, 2032	DP			
	9474754	Aug 05, 2033	U-2802			
	9593099	Aug 27, 2030	DS			
	9593100	Aug 27, 2030	DP			
	9763941	Nov 21, 2032	U-2335			
	9850229	Aug 27, 2030	U-2337			
	9850230	Aug 27, 2030	U-2334			
	9850230	Aug 27, 2030	U-2802			
	9850230	Aug 27, 2030	U-2803			
	9850230	Aug 27, 2030	U-3738			
	RE49556	Feb 27, 2030	DS DP			
<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 001	10231965	Feb 17, 2035	U-2617		NCE	Aug 15, 2024
	10231965	Feb 17, 2035	U-2618		NPP	Oct 20, 2026
	10398693	Jul 18, 2038	DP		ODE-265	Aug 15, 2026
	10561651	Feb 19, 2035	U-2745		ODE-313	Aug 15, 2026
	10738037	May 18, 2037	DS DP U-2946		ODE-448	Oct 20, 2030

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<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 001	11091469	May 18, 2037	U-2617			
	11091469	May 18, 2037	U-2618			
	11253515	Jul 18, 2038	DP			
	8299057	Mar 01, 2029	DS DP			
	8673893	Jul 08, 2028	U-2617			
	8673893	Jul 08, 2028	U-2618			
	9029356	Jul 08, 2028	DS DP			
	9085558	Jul 08, 2028	DP			
	9085565	May 22, 2033	DS DP			
	9255087	Jul 08, 2028	U-2617			
	9255087	Jul 08, 2028	U-2618			
	9616059	Jul 08, 2028	U-2618			
	9649306	May 22, 2033	U-2617			
	9649306	May 22, 2033	U-2618			
<u>ENTRECTINIB - ROZLYTREK</u>						
N 212725 002	10231965	Feb 17, 2035	U-2617		NCE	Aug 15, 2024
	10231965	Feb 17, 2035	U-2618		NPP	Oct 20, 2026
	10398693	Jul 18, 2038	DP		ODE-265	Aug 15, 2026
	10561651	Feb 19, 2035	U-2745		ODE-313	Aug 15, 2026
	10738037	May 18, 2037	DS DP U-2946		ODE-448	Oct 20, 2030
	11091469	May 18, 2037	U-2617			
	11091469	May 18, 2037	U-2618			
	11253515	Jul 18, 2038	DP			
	8299057	Mar 01, 2029	DS DP			
	8673893	Jul 08, 2028	U-2617			
	8673893	Jul 08, 2028	U-2618			
	9029356	Jul 08, 2028	DS DP			
	9085558	Jul 08, 2028	DP			
	9085565	May 22, 2033	DS DP			
	9255087	Jul 08, 2028	U-2617			
	9255087	Jul 08, 2028	U-2618			
	9616059	Jul 08, 2028	U-2618			
	9649306	May 22, 2033	U-2617			
	9649306	May 22, 2033	U-2618			
<u>ENTRECTINIB - ROZLYTREK</u>						
N 218550 001					NCE	Aug 15, 2024
					NP	Oct 20, 2026
					ODE-448	Oct 20, 2030
<u>ENZALUTAMIDE - XTANDI</u>						
N 203415 001	7709517	Aug 13, 2027	DS DP		I-926	Nov 17, 2026
	8183274	Aug 24, 2026	U-1281			
	8183274	Aug 24, 2026	U-1588			
	8183274	Aug 24, 2026	U-2345			
	8183274	Aug 24, 2026	U-2708			
	8183274	Aug 24, 2026	U-3763			
	9126941	May 15, 2026	U-1588			
	9126941	May 15, 2026	U-2345			
	9126941	May 15, 2026	U-2708			

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<u>ENZALUTAMIDE - XTANDI</u>						
N 203415	001 9126941	May 15, 2026	U-3763			
<u>ENZALUTAMIDE - XTANDI</u>						
N 213674	001 11839689	Sep 11, 2033	DP		I-926	Nov 17, 2026
	7709517	Aug 13, 2027	DS DP			
	8183274	Aug 24, 2026	U-2345			
	8183274	Aug 24, 2026	U-2708			
	8183274	Aug 24, 2026	U-3763			
	9126941	May 15, 2026	U-2345			
	9126941	May 15, 2026	U-2708			
	9126941	May 15, 2026	U-3763			
<u>ENZALUTAMIDE - XTANDI</u>						
N 213674	002 11839689	Sep 11, 2033	DP		I-926	Nov 17, 2026
	7709517	Aug 13, 2027	DS DP			
	8183274	Aug 24, 2026	U-2345			
	8183274	Aug 24, 2026	U-2708			
	8183274	Aug 24, 2026	U-3763			
	9126941	May 15, 2026	U-2345			
	9126941	May 15, 2026	U-2708			
	9126941	May 15, 2026	U-3763			
<u>EPHEDRINE SULFATE - EMERPHED</u>						
N 213407	001 11090278	May 16, 2040	U-3183			
	11241400	May 16, 2040	U-3183			
	11464752	May 16, 2040	DP			
	11478436	May 16, 2040	U-3183			
	11571398	May 16, 2040	U-3183			
<u>EPHEDRINE SULFATE - EMERPHED</u>						
N 213407	002 11464752	May 16, 2040	DP			
	11571398	May 16, 2040	U-3183			
<u>EPHEDRINE SULFATE - EPHEDRINE SULFATE</u>						
N 213994	001 10869845	Jan 22, 2040	DP			
<u>EPHEDRINE SULFATE - EPHEDRINE SULFATE</u>						
N 213994	002 10869845	Jan 22, 2040	DP			
<u>EPINEPHRINE - EPIPEN</u>						
N 019430	001 7449012	Sep 11, 2025	DP			
	7794432	Sep 11, 2025	DP			
	8048035	Sep 11, 2025	DP			
	8870827	Sep 11, 2025	DP			
	9586010	Sep 11, 2025	DP			
<u>EPINEPHRINE - EPIPEN JR.</u>						
N 019430	002 7449012	Sep 11, 2025	DP			
	7794432	Sep 11, 2025	DP			
	8048035	Sep 11, 2025	DP			
	8870827	Sep 11, 2025	DP			
	9586010	Sep 11, 2025	DP			

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<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	001	10314977	Nov 23, 2024	DP		
		10335549	Apr 30, 2025	DP		
		10737028	Nov 23, 2024	DP		
		11590286	Dec 12, 2026	DP		
		7918823	Nov 23, 2024	DP		
		7947017	Mar 12, 2028	DP		
		8021344	Nov 02, 2029	DP		
		8206360	Feb 27, 2027	DP		
		8226610	Apr 10, 2029	DP		
		8231573	Nov 25, 2028	DP		
		8313466	Nov 23, 2024	DP		
		8361029	Nov 23, 2024	DP		
		8425462	Nov 23, 2024	DP		
		8608698	Nov 23, 2024	DP		
		9056170	Nov 23, 2024	DP		
		9149579	Jul 19, 2025	U-1758		
		9259539	Feb 01, 2026	DP		
		9278182	Feb 01, 2026	DP		
		9724471	May 23, 2027	DP	U-2092	
		9737669	Nov 23, 2024	DP		
<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	002	10314977	Nov 23, 2024	DP		
		10335549	Apr 30, 2025	DP		
		10688244	Dec 21, 2037	DP	U-2980	
		10737028	Nov 23, 2024	DP		
		10842938	Dec 21, 2037	DP	U-2980	
		11590286	Dec 12, 2026	DP		
		11771830	Dec 21, 2037	DP	U-2980	
		7918823	Nov 23, 2024	DP		
		7947017	Mar 12, 2028	DP		
		8021344	Nov 02, 2029	DP		
		8206360	Feb 27, 2027	DP		
		8226610	Apr 10, 2029	DP		
		8231573	Nov 25, 2028	DP		
		8313466	Nov 23, 2024	DP		
		8361029	Nov 23, 2024	DP		
		8425462	Nov 23, 2024	DP		
		8608698	Nov 23, 2024	DP		
		9056170	Nov 23, 2024	DP		
		9149579	Jul 19, 2025	U-1758		
		9259539	Feb 01, 2026	DP		
		9278182	Feb 01, 2026	DP		
		9724471	May 23, 2027	DP	U-2092	
		9737669	Nov 23, 2024	DP		
<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	003	10314977	Nov 23, 2024	DP		
		10335549	Apr 30, 2025	DP		
		10688244	Dec 21, 2037	DP	U-2980	
		10737028	Nov 23, 2024	DP		

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<u>EPINEPHRINE - AUVI-Q</u>						
N 201739	003 10842938	Dec 21, 2037	DP U-2980			
	11590286	Dec 12, 2026	DP			
	11771830	Dec 21, 2037	DP U-2980			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	9056170	Nov 23, 2024	DP			
	9149579	Jul 19, 2025		U-1758		
	9259539	Feb 01, 2026	DP			
	9278182	Feb 01, 2026	DP			
	9724471	May 23, 2027	DP U-2092			
	9737669	Nov 23, 2024	DP			
	9833573	Nov 23, 2024		U-2172		
<u>EPINEPHRINE - ADRENALIN</u>						
N 204200	001 9119876	Mar 13, 2035	DP			
	9295657	Mar 13, 2035		U-1829		
<u>EPINEPHRINE - ADRENALIN</u>						
N 204640	001 10130592	Mar 13, 2035	DP			
	9119876	Mar 13, 2035	DP			
	9295657	Mar 13, 2035		U-1829		
<u>EPINEPHRINE - EPINEPHRINE</u>						
N 205029	001 10004700	Aug 14, 2034	DP U-2325		Y	
	10039728	Aug 14, 2034		U-1828		Y
	9283197	Aug 15, 2034	DP U-1828			Y
	9283197	Aug 15, 2034	DP U-1829			Y
	9283197	Aug 15, 2034	DP U-1830			Y
<u>EPINEPHRINE - PRIMATENE MIST</u>						
N 205920	001 8367734	Jan 26, 2026	DP			
<u>EPINEPHRINE - SYMJEPI</u>						
N 207534	001 11141540	Oct 20, 2036	DP U-3379			
<u>EPINEPHRINE - SYMJEPI</u>						
N 207534	002 11141540	Oct 20, 2036	DP U-3379			
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	001 10653646	Mar 21, 2039	DP			
	11083698	Mar 21, 2039		U-3567		
	11207280	Mar 21, 2039	DP			
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	002 10653646	Mar 21, 2039	DP			
	11083698	Mar 21, 2039		U-3567		

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<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	002 11207280	Mar 21, 2039	DP			
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	003 10653646	Mar 21, 2039	DP			
	11083698	Mar 21, 2039		U-3567		
	11207280	Mar 21, 2039	DP			
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	004 10653646	Mar 21, 2039	DP			
	11083698	Mar 21, 2039		U-3567		
	11207280	Mar 21, 2039	DP			
<u>EPINEPHRINE - ADRENALIN</u>						
N 215875	005 10653646	Mar 21, 2039	DP			
	11083698	Mar 21, 2039		U-3567		
	11207280	Mar 21, 2039	DP			
<u>EPOPROSTENOL SODIUM - VELETRI</u>						
N 022260	001 8318802	Mar 15, 2027	DP			
	8598227	Feb 02, 2027				
<u>EPOPROSTENOL SODIUM - VELETRI</u>						
N 022260	002 8318802	Mar 15, 2027	DP			
	8598227	Feb 02, 2027				
<u>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	001 10961190	Oct 19, 2037	DS		NCE	Aug 27, 2023
	11578044	Oct 19, 2037	DS		GAIN	Aug 27, 2028
	8796245	Aug 07, 2029		U-2380		
	8906887	Dec 28, 2030	DP			
<u>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u>						
N 211109	002 10961190	Oct 19, 2037	DS		NCE	Aug 27, 2023
	11578044	Oct 19, 2037	DS		GAIN	Aug 27, 2028
	8796245	Aug 07, 2029		U-2380		
	8906887	Dec 28, 2030	DP			
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	001 10898482	Feb 09, 2036	DP	U-2518	NCE	Apr 12, 2024
	10898482	Feb 09, 2036	DP	U-3065		
	10898482	Feb 09, 2036	DP	U-3066		
	10898482	Feb 09, 2036	DP	U-3067		
	11077106	Feb 02, 2038		U-3196		
	11684620	Feb 09, 2036		U-2518		
	11684620	Feb 09, 2036		U-3065		
	11684620	Feb 09, 2036		U-3066		
	11684620	Feb 09, 2036		U-3067		
	8895601	Apr 12, 2033	DS	DP		
	9464071	Apr 28, 2031		U-2518		
	9902714	Mar 26, 2035	DP			
<u>ERDAFITINIB - BALVERSA</u>						
N 212018	002 10898482	Feb 09, 2036	DP	U-2518	NCE	Apr 12, 2024
	10898482	Feb 09, 2036	DP	U-3065		
	10898482	Feb 09, 2036	DP	U-3066		

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<u>ERDAFITINIB - BALVERSA</u>						
N 212018 002	10898482	Feb 09, 2036	DP U-3067			
	11077106	Feb 02, 2038	U-3196			
	11684620	Feb 09, 2036	U-2518			
	11684620	Feb 09, 2036	U-3065			
	11684620	Feb 09, 2036	U-3066			
	11684620	Feb 09, 2036	U-3067			
	8895601	Apr 12, 2033	DS DP			
	9464071	Apr 28, 2031	U-2518			
	9902714	Mar 26, 2035	DP			
<u>ERDAFITINIB - BALVERSA</u>						
N 212018 003	10898482	Feb 09, 2036	DP U-2518		NCE	Apr 12, 2024
	10898482	Feb 09, 2036	DP U-3065			
	10898482	Feb 09, 2036	DP U-3066			
	10898482	Feb 09, 2036	DP U-3067			
	11077106	Feb 02, 2038	U-3196			
	11684620	Feb 09, 2036	U-2518			
	11684620	Feb 09, 2036	U-3065			
	11684620	Feb 09, 2036	U-3066			
	11684620	Feb 09, 2036	U-3067			
	8895601	Apr 12, 2033	DS DP			
	9464071	Apr 28, 2031	U-2518			
	9902714	Mar 26, 2035	DP			
<u>ERIBULIN MESYLATE - HALAVEN</u>						
N 201532 001	6214865*PED	Jan 20, 2024			M-280	Sep 13, 2025
	RE46965	Jan 08, 2027	DP		PED	Mar 13, 2026
	RE46965*PED	Jul 08, 2027				
<u>ERTUGLIFLOZIN - STEGLATRO</u>						
N 209803 001	8080580	Jul 13, 2030	DS DP U-2214		M-275	Sep 17, 2024
<u>ERTUGLIFLOZIN - STEGLATRO</u>						
N 209803 002	8080580	Jul 13, 2030	DS DP U-2214		M-275	Sep 17, 2024
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806 001	8080580	Jul 13, 2030	DS DP U-2214		M-275	Sep 17, 2024
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806 002	8080580	Jul 13, 2030	DS DP U-2214		M-275	Sep 17, 2024
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806 003	8080580	Jul 13, 2030	DS DP U-2214		M-275	Sep 17, 2024
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUOMET</u>						
N 209806 004	8080580	Jul 13, 2030	DS DP U-2214		M-275	Sep 17, 2024
	9308204	Oct 21, 2030	DP			
	9439902	Oct 21, 2030	U-2214			

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<u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u>						
N 209805 001	7326708	Nov 24, 2026	DS DP U-2214		M-275	Sep 17, 2024
	7326708*PED	May 24, 2027				
	8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439901	Oct 21, 2030	U-2214			
<u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u>						
N 209805 002	7326708	Nov 24, 2026	DS DP U-2214		M-275	Sep 17, 2024
	7326708*PED	May 24, 2027				
	8080580	Jul 13, 2030	DS DP U-2214			
	9308204	Oct 21, 2030	DP			
	9439901	Oct 21, 2030	U-2214			
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 001					NPP	May 12, 2026
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 002					NPP	May 12, 2026
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021323 003					NPP	May 12, 2026
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
N 021365 001					NPP	May 12, 2026
<u>ESKETAMINE HYDROCHLORIDE - SPRAVATO</u>						
N 211243 001	10869844	Sep 10, 2035	U-3034		NCE*	Mar 05, 2024
	10869844	Sep 10, 2035	U-3035			
	10869844	Sep 10, 2035	U-3036			
	11173134	Sep 10, 2035	U-3257			
	11173134	Sep 10, 2035	U-3536			
	11311500	Sep 10, 2035	U-3034			
	11311500	Sep 10, 2035	U-3035			
	11311500	Sep 10, 2035	U-3036			
	11446260	Mar 14, 2034	U-3444			
	11446260	Mar 14, 2034	U-3445			
	11446260	Mar 14, 2034	U-3446			
	8785500	Mar 05, 2033	U-2502			
	9592207	Mar 20, 2027	U-2502			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 001	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			

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<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 001	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 002	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 003	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416 004	10675287	May 06, 2025	U-2041			
	10675287	May 06, 2025	U-2831			
	10695354	May 06, 2025	U-2501			
	10695354	May 06, 2025	U-2831			
	10702536	May 06, 2025	U-2501			
	10912781	Oct 23, 2028	DP			
	11364247	May 06, 2025	U-2501			
	11364247	May 06, 2025	U-2831			
	8372431	Apr 17, 2030	DP			
	9206135	Apr 21, 2026	DS			
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			

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<u>ESLICARBAZEPINE ACETATE - APTIOM</u>						
N 022416	004	9750747	Aug 24, 2032	U-2121		
		9763954	Sep 13, 2028	U-2123		
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER</u>						
N 205703	001	8829054	Mar 15, 2033	DP		
		8835505	Mar 15, 2033	DP		
<u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER</u>						
N 205703	002	8829054	Mar 15, 2033	DP		
		8835505	Mar 15, 2033	DP		
<u>ESOMEPRAZOLE MAGNESIUM - ESOMEPRAZOLE MAGNESIUM</u>						
N 214278	001	10076494	Dec 08, 2036	DP		
		10835488	Dec 08, 2036	DP		
<u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u>						
N 022511	001	8945621	Oct 17, 2031	U-1661		
		9220698	Mar 10, 2031	U-1781		
		9393208	Sep 03, 2029	U-1781		
<u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u>						
N 022511	002	8945621	Oct 17, 2031	U-1661		
		9393208	Sep 03, 2029	U-1781		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	001	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028		U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	002	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028		U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	003	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028		U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	004	8231906	Jul 04, 2030	DS DP		
		9730900	Jul 10, 2028		U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - MINIVELLE</u>						
N 203752	005	8231906	Jul 04, 2030	DS DP		
		9724310	Jul 10, 2028	DS DP		
		9730900	Jul 10, 2028		DP U-2086	
		9833419	Jul 10, 2028	DP		
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	001	10258630	Dec 20, 2033	U-2316		
		10258630	Dec 20, 2033	U-2317		
		10398708	Dec 20, 2033	U-2317		
		10398708	Dec 20, 2033	U-2614		
		10471072	Jun 18, 2033	U-2316		

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<u>ESTRADIOL - IMVEXXY</u>						
N 208564	001	10471072				
		Jun 18, 2033	U-2317			
		10537581		DP		U-2316
		Nov 21, 2032	DP			U-2317
		10537581		DP		U-2317
		Nov 21, 2032	DP			U-2317
		10568891				U-2316
		Jun 18, 2033	U-2316			
		10568891				U-2317
		Jun 18, 2033	U-2317			
		10668082				U-2316
		Jun 18, 2033	U-2316			
		10668082				U-2317
		Jun 18, 2033	U-2317			
		10806697		DP		
		Nov 21, 2032	DP			
		10835487				U-2316
		Nov 21, 2032	U-2316			
		10835487				U-2317
		Nov 21, 2032	U-2317			
		10888516				U-2316
		Jun 18, 2033	U-2316			
		10888516				U-2317
		Jun 18, 2033	U-2317			
		11065197		DP		
		Jun 18, 2033	DP			
		11116717		DP		
		Jun 18, 2033	DP			
		11123283		DP		
		Jun 18, 2033	DP			
		11241445				U-2316
		Nov 21, 2032	U-2316			
		11241445				U-2317
		Nov 21, 2032	U-2317			
		11246875				U-2316
		Nov 21, 2032	U-2316			
		11246875				U-2317
		Nov 21, 2032	U-2317			
		11266661				U-2316
		Feb 02, 2034	U-2316			
		11266661				U-2317
		Feb 02, 2034	U-2317			
		11304959		DP		
		Nov 21, 2032	DP			
		11351182				U-2316
		Nov 21, 2032	U-2316			
		11351182				U-2317
		Nov 21, 2032	U-2317			
		11497709				U-2316
		Nov 21, 2032	U-2316			
		11497709				U-2317
		Nov 21, 2032	U-2317			
		9180091		DP		U-2316
		Dec 20, 2033	DP			U-2317
		9180091		DP		U-2317
		Dec 20, 2033	DP			U-2317
		9289382		DP		
		Nov 21, 2032	DP			
<u>ESTRADIOL - IMVEXXY</u>						
N 208564	002	10258630				
		Dec 20, 2033	U-2316			
		10258630				U-2317
		Dec 20, 2033	U-2317			
		10398708				U-2317
		Dec 20, 2033	U-2317			
		10398708				U-2614
		Dec 20, 2033	U-2614			
		10471072				U-2316
		Jun 18, 2033	U-2316			
		10471072				U-2317
		Jun 18, 2033	U-2317			
		10537581		DP		U-2316
		Nov 21, 2032	DP			U-2317
		10537581		DP		U-2317
		Nov 21, 2032	DP			U-2317
		10568891				U-2316
		Jun 18, 2033	U-2316			
		10568891				U-2317
		Jun 18, 2033	U-2317			
		10668082				U-2316
		Jun 18, 2033	U-2316			
		10668082				U-2317
		Jun 18, 2033	U-2317			
		10806697		DP		
		Nov 21, 2032	DP			
		10835487				U-2316
		Nov 21, 2032	U-2316			
		10835487				U-2317
		Nov 21, 2032	U-2317			
		10888516				U-2316
		Jun 18, 2033	U-2316			
		10888516				U-2317
		Jun 18, 2033	U-2317			
		11065197		DP		
		Jun 18, 2033	DP			
		11116717		DP		
		Jun 18, 2033	DP			

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<u>ESTRADIOL - IMVEXXY</u>						
N 208564 002	11123283	Jun 18, 2033	DP			
	11241445	Nov 21, 2032	U-2316			
	11241445	Nov 21, 2032	U-2317			
	11246875	Nov 21, 2032	U-2316			
	11246875	Nov 21, 2032	U-2317			
	11266661	Feb 02, 2034	U-2316			
	11266661	Feb 02, 2034	U-2317			
	11304959	Nov 21, 2032	DP			
	11351182	Nov 21, 2032	U-2316			
	11351182	Nov 21, 2032	U-2317			
	11497709	Nov 21, 2032	U-2316			
	11497709	Nov 21, 2032	U-2317			
	9180091	Dec 20, 2033	DP U-2316			
	9180091	Dec 20, 2033	DP U-2317			
	9289382	Nov 21, 2032	DP			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633 001	7572779	Oct 02, 2025	U-904			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633 002	7572779	Oct 02, 2025	U-904			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N 021633 003	7572779	Oct 02, 2025	U-904			
<u>ESTRADIOL; NORETHINDRONE ACETATE; RELUGOLIX - MYFEMBREE</u>						
N 214846 001	11033551	Sep 29, 2037	U-3129		I-898	Aug 05, 2025
	11793812	May 03, 2038	U-2360		M-289	Jan 27, 2026
	11795178	Sep 27, 2033	DS DP		NCE	Dec 18, 2025
	7300935	Jan 28, 2025	DS		NP	May 26, 2024
	8058280	Jan 28, 2025	DS			
	9346822	Feb 17, 2024	U-3129			
<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132 001	10052386	Nov 21, 2032	DP			
	10206932	Nov 21, 2032	U-2439			
	10639375	Nov 21, 2032	DP			
	10675288	Nov 21, 2032	U-2439			
	10806740	Nov 21, 2032	DP U-2439			
	11033626	Nov 21, 2032	DP U-2439			
	11103513	Nov 21, 2032	U-2439			
	11103516	Nov 21, 2032	DP			
	11110099	Nov 21, 2032	DP			
	11529360	Nov 21, 2032	DP			
	11793819	Nov 21, 2032	U-2439			
	8633178	Nov 21, 2032	DP			
	8846648	Nov 21, 2032	U-2439			
	8846649	Nov 21, 2032	DP U-2439			
	8987237	Nov 21, 2032	DP			
	8993548	Nov 21, 2032	DP			
	8993549	Nov 21, 2032	DP			
	9006222	Nov 21, 2032	DP U-2439			
	9114145	Nov 21, 2032	U-2439			

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<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	001	9114146	Nov 21, 2032	DP	U-2439	
		9301920	Nov 21, 2032	DP	U-2439	
<u>ESTRADIOL; PROGESTERONE - BIJUVA</u>						
N 210132	002	10052386	Nov 21, 2032	DP		
		10206932	Nov 21, 2032		U-2439	
		10675288	Nov 21, 2032		U-2439	
		11033626	Nov 21, 2032	DP	U-2439	
		11166963	Nov 21, 2032	DP		
		11793819	Nov 21, 2032		U-2439	
		8633178	Nov 21, 2032	DP		
		8846648	Nov 21, 2032		U-2439	
		8846649	Nov 21, 2032	DP	U-2439	
		8933059	Nov 21, 2032	DP	U-2439	
		8987237	Nov 21, 2032	DP		
		8993548	Nov 21, 2032	DP		
		8993549	Nov 21, 2032	DP		
		9114145	Nov 21, 2032		U-2439	
		9114146	Nov 21, 2032	DP	U-2439	
		9301920	Nov 21, 2032	DP	U-2439	
<u>ETELALCETIDE - PARSABIV</u>						
N 208325	001	10344765	Jun 27, 2034	DP		
		11162500	Jun 27, 2034	DP		
		8377880	Jul 29, 2030	DS DP		
		8999932	Feb 07, 2031	DS DP	U-2014	
		9278995	Jul 29, 2030	DS		
		9701712	Jul 29, 2030	DS DP	U-2014	
		9820938	Jun 27, 2034	DP		
<u>ETELALCETIDE - PARSABIV</u>						
N 208325	002	10344765	Jun 27, 2034	DP		
		11162500	Jun 27, 2034	DP		
		8377880	Jul 29, 2030	DS DP		
		8999932	Feb 07, 2031	DS DP	U-2014	
		9278995	Jul 29, 2030	DS		
		9701712	Jul 29, 2030	DS DP	U-2014	
		9820938	Jun 27, 2034	DP		
<u>ETELALCETIDE - PARSABIV</u>						
N 208325	003	10344765	Jun 27, 2034	DP		
		11162500	Jun 27, 2034	DP		
		8377880	Jul 29, 2030	DS DP		
		8999932	Feb 07, 2031	DS DP	U-2014	
		9278995	Jul 29, 2030	DS		
		9701712	Jul 29, 2030	DS DP	U-2014	
		9820938	Jun 27, 2034	DP		
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	001	10337003	Mar 14, 2034		U-1918	
		10364431	Mar 14, 2034		U-1918	
		10364431	Mar 14, 2034		U-1919	
		10781451	Jun 28, 2025	DS DP		

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<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	001	9018368	Jun 28, 2025	DS DP		
		9243245	Oct 27, 2028	DS	U-2097	
		9243245	Oct 27, 2028	DS	U-2098	
		9506058	Mar 14, 2034		U-1918	
		9506058	Mar 14, 2034		U-1919	
		RE47751	Jun 28, 2025		U-1918	
		RE47751	Jun 28, 2025		U-2664	
		RE47751	Jun 28, 2025		U-2673	
		RE47751	Jun 28, 2025		U-2674	
		RE47769	Feb 02, 2029	DP		
		RE48468	Oct 27, 2028		U-2097	
<u>ETEPLIRSEN - EXONDYS 51</u>						
N 206488	002	10337003	Mar 14, 2034		U-1918	
		10364431	Mar 14, 2034		U-1918	
		10364431	Mar 14, 2034		U-1919	
		10781451	Jun 28, 2025	DS DP		
		9018368	Jun 28, 2025	DS DP		
		9243245	Oct 27, 2028	DS	U-2097	
		9243245	Oct 27, 2028	DS	U-2098	
		9506058	Mar 14, 2034		U-1918	
		9506058	Mar 14, 2034		U-1919	
		RE47751	Jun 28, 2025		U-1918	
		RE47751	Jun 28, 2025		U-2664	
		RE47751	Jun 28, 2025		U-2673	
		RE47751	Jun 28, 2025		U-2674	
		RE47769	Feb 02, 2029	DP		
		RE48468	Oct 27, 2028		U-2097	
<u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u>						
N 021840	001	7320969	Jan 30, 2024		U-828	
		7855190	Dec 05, 2028		U-1	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - LOSEASONIQUE</u>						
N 022262	001	7855190	Dec 05, 2028		U-1	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - TWIRLA</u>						
N 204017	001	8246978	Aug 26, 2028	DP		
		8747888	Jul 10, 2028	DP		
		9050348	Jul 10, 2028	DP		
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - QUARTETTE</u>						
N 204061	001	8415332	Mar 11, 2029	DP		
		8450299	Oct 07, 2025		U-1	
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - BALCOLTRA</u>						
N 208612	001	7838042	Jun 01, 2027	DS	U-3251	
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO LOESTRIN FE</u>						
N 022501	001	7704984	Feb 02, 2029		U-1090	
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO MINASTRIN FE</u>						
N 204654	001	7704984	Feb 02, 2029		U-1	

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<u>ETHINYL ESTRADIOL; SEGESTERONE ACETATE - ANNOVERA</u>						
N 209627	001 10632066	Feb 01, 2039	U-2786			
	10632066	Feb 01, 2039	U-2787			
	10765628	Feb 01, 2039	U-2786			
	10765628	Feb 01, 2039	U-2787			
	10780047	Feb 01, 2039	U-2786			
	10780047	Feb 01, 2039	U-2787			
	10918649	Jun 21, 2039	DP			
	10925882	Jun 21, 2039	DP			
	10940157	Jun 21, 2039	DP			
	11529308	Jun 21, 2039	DP			
<u>ETONOGESTREL - IMPLANON</u>						
N 021529	001 9757552	Jul 28, 2030	DP U-1			
<u>ETONOGESTREL - NEXPLANON</u>						
N 021529	002 10821277	May 31, 2027	DP			
	8722037	Sep 28, 2027	DP			
	8888745	Aug 28, 2026	DP			
	9757552	Jul 28, 2030	DP U-1			
<u>ETRASIMOD ARGININE - VELSIPITY</u>						
N 216956	001 10301262	Jun 21, 2036	DS DP		NCE	Oct 12, 2028
	10676435	Jun 21, 2036	U-3731			
	11007175	Jan 06, 2036	U-3730			
	11091435	Jun 21, 2036	DS DP			
	8580841	Mar 05, 2030	DS DP			
	9126932	Jul 22, 2029	U-3732			
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	001 8410131	Nov 01, 2025	U-1368			
	8410131*PED	May 01, 2026				
	9006224	Jul 01, 2028	U-1681			
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	002 8410131	Nov 01, 2025	U-1368			
	8410131*PED	May 01, 2026				
	9006224	Jul 01, 2028	U-1681			
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	003 8410131	Nov 01, 2025	U-1368			
	8410131*PED	May 01, 2026				
	9006224	Jul 01, 2028	U-1681			
<u>EVEROLIMUS - AFINITOR</u>						
N 022334	004 8410131	Nov 01, 2025	U-1368			
	8410131*PED	May 01, 2026				
	9006224	Jul 01, 2028	U-1681			
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985	001				ODE-169	Apr 10, 2025
<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985	002				ODE-169	Apr 10, 2025

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<u>EVEROLIMUS - AFINITOR DISPERZ</u>						
N 203985	003				ODE-169	Apr 10, 2025
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N 021773	001				M-232	Nov 04, 2024
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N 021773	002				M-232	Nov 04, 2024
<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200	001	6515117	Oct 04, 2025	DS DP U-2588	NPP	Jul 22, 2024
		6515117*PED	Apr 04, 2026		PED	Jan 22, 2025
		7456254	Jun 30, 2025	DP U-2588		
		7456254	Jun 30, 2025	DP U-2589		
		7456254	Jun 30, 2025	DP U-2590		
		7456254	Jun 30, 2025	DP U-3188		
		7456254	Jun 30, 2025	DP U-3189		
		7456254	Jun 30, 2025	DP U-3190		
		7456254*PED	Dec 30, 2025			
		7563871	Apr 15, 2024	DP		
		7612176	Apr 13, 2025	DP U-2588		
		7612176	Apr 13, 2025	DP U-2589		
		7612176	Apr 13, 2025	DP U-2590		
		7612176	Apr 13, 2025	DP U-3188		
		7612176	Apr 13, 2025	DP U-3189		
		7612176	Apr 13, 2025	DP U-3190		
		7612176*PED	Oct 13, 2025			
		8329648	Aug 18, 2026	U-2588		
		8329648	Aug 18, 2026	U-2589		
		8329648	Aug 18, 2026	U-2590		
		8329648	Aug 18, 2026	U-2593		
		8329648	Aug 18, 2026	U-2594		
		8329648	Aug 18, 2026	U-2595		
		8329648	Aug 18, 2026	U-2596		
		8329648	Aug 18, 2026	U-3188		
		8329648	Aug 18, 2026	U-3189		
		8329648	Aug 18, 2026	U-3190		
		8329648*PED	Feb 18, 2027			
		8361972	Mar 21, 2028	U-2588		
		8361972*PED	Sep 21, 2028			
		8431685	Apr 13, 2025	DP U-2588		
		8431685	Apr 13, 2025	DP U-2589		
		8431685	Apr 13, 2025	DP U-2590		
		8431685	Apr 13, 2025	DP U-3188		
		8431685	Apr 13, 2025	DP U-3189		
		8431685	Apr 13, 2025	DP U-3190		
		8431685*PED	Oct 13, 2025			
		8461105	Apr 13, 2025	DP U-2588		
		8461105	Apr 13, 2025	DP U-2589		
		8461105	Apr 13, 2025	DP U-2590		
		8461105	Apr 13, 2025	DP U-3188		
		8461105	Apr 13, 2025	DP U-3189		

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<u>EXENATIDE SYNTHETIC - BYDUREON</u>						
N 022200 001	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP U-2588			
	8501698*PED	Dec 20, 2027				
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	8906851	Aug 18, 2026	U-3188			
	8906851	Aug 18, 2026	U-3189			
	8906851	Aug 18, 2026	U-3190			
	8906851*PED	Feb 18, 2027				
	9238076	Apr 15, 2024	DP U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			
	9238076	Apr 15, 2024	DP U-2599			
	9238076	Apr 15, 2024	DP U-3188			
	9238076	Apr 15, 2024	DP U-3189			
	9238076	Apr 15, 2024	DP U-3190			
	9238076*PED	Oct 15, 2024				
	9884092	Aug 18, 2026	U-2588			
	9884092	Aug 18, 2026	U-2589			
	9884092	Aug 18, 2026	U-2590			
	9884092	Aug 18, 2026	U-2593			
	9884092	Aug 18, 2026	U-2594			
	9884092	Aug 18, 2026	U-2595			
	9884092	Aug 18, 2026	U-2596			
	9884092	Aug 18, 2026	U-3188			
	9884092	Aug 18, 2026	U-3189			
	9884092	Aug 18, 2026	U-3190			
	9884092*PED	Feb 18, 2027				
<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	6515117	Oct 04, 2025	DS DP U-2588		NPP	Jul 22, 2024
	6515117*PED	Apr 04, 2026			PED	Jan 22, 2025
	7456254	Jun 30, 2025	DP U-2588			
	7456254	Jun 30, 2025	DP U-2589			
	7456254	Jun 30, 2025	DP U-2590			
	7456254	Jun 30, 2025	DP U-3188			
	7456254	Jun 30, 2025	DP U-3189			
	7456254	Jun 30, 2025	DP U-3190			
	7456254*PED	Dec 30, 2025				
	7563871	Apr 15, 2024	DP			
	7563871*PED	Oct 15, 2024				
	7612176	Apr 13, 2025	DP U-2588			
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			

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<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	7612176*PED	Oct 13, 2025				
	8216180	Jan 12, 2028	DP			
	8216180*PED	Jul 12, 2028				
	8329648	Aug 18, 2026		U-2588		
	8329648	Aug 18, 2026		U-2589		
	8329648	Aug 18, 2026		U-2590		
	8329648	Aug 18, 2026		U-2593		
	8329648	Aug 18, 2026		U-2594		
	8329648	Aug 18, 2026		U-2595		
	8329648	Aug 18, 2026		U-2596		
	8329648	Aug 18, 2026		U-3188		
	8329648	Aug 18, 2026		U-3189		
	8329648	Aug 18, 2026		U-3190		
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028		U-2588		
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP	U-2588		
	8431685	Apr 13, 2025	DP	U-2589		
	8431685	Apr 13, 2025	DP	U-2590		
	8431685	Apr 13, 2025	DP	U-3188		
	8431685	Apr 13, 2025	DP	U-3189		
	8431685	Apr 13, 2025	DP	U-3190		
	8431685*PED	Oct 13, 2025				
	8439864	Mar 25, 2028	DP			
	8439864*PED	Sep 25, 2028				
	8461105	Apr 13, 2025	DP	U-2588		
	8461105	Apr 13, 2025	DP	U-2589		
	8461105	Apr 13, 2025	DP	U-2590		
	8461105	Apr 13, 2025	DP	U-3188		
	8461105	Apr 13, 2025	DP	U-3189		
	8461105	Apr 13, 2025	DP	U-3190		
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	DP	U-2588		
	8501698*PED	Dec 20, 2027				
	8690837	May 19, 2029	DP			
	8690837*PED	Nov 19, 2029				
	8721615	Jan 18, 2030	DP			
	8721615*PED	Jul 18, 2030				
	8758292	Nov 12, 2027	DP			
	8758292*PED	May 12, 2028				
	8827963	Feb 04, 2029	DP			
	8827963*PED	Aug 04, 2029				
	8906851	Aug 18, 2026		U-2588		
	8906851	Aug 18, 2026		U-2589		
	8906851	Aug 18, 2026		U-2590		
	8906851	Aug 18, 2026		U-2593		
	8906851	Aug 18, 2026		U-3188		
	8906851	Aug 18, 2026		U-3189		
	8906851	Aug 18, 2026		U-3190		

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<u>EXENATIDE SYNTHETIC - BYDUREON PEN</u>						
N 022200 002	8906851*PED	Feb 18, 2027				
	8998876	Jan 07, 2030	DP			
	8998876*PED	Jul 07, 2030				
	9238076	Apr 15, 2024	DP U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			
	9238076	Apr 15, 2024	DP U-2599			
	9238076	Apr 15, 2024	DP U-3188			
	9238076	Apr 15, 2024	DP U-3189			
	9238076	Apr 15, 2024	DP U-3190			
	9238076*PED	Oct 15, 2024				
	9320853	Mar 25, 2028	DP			
	9320853*PED	Sep 25, 2028				
	9884092	Aug 18, 2026	U-2588			
	9884092	Aug 18, 2026	U-2589			
	9884092	Aug 18, 2026	U-2590			
	9884092	Aug 18, 2026	U-2593			
	9884092	Aug 18, 2026	U-2594			
	9884092	Aug 18, 2026	U-2595			
	9884092	Aug 18, 2026	U-2596			
	9884092	Aug 18, 2026	U-3188			
	9884092	Aug 18, 2026	U-3189			
	9884092	Aug 18, 2026	U-3190			
	9884092*PED	Feb 18, 2027				
<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210 001	6515117	Oct 04, 2025	DS DP U-2588		NPP	Jul 22, 2024
	6515117*PED	Apr 04, 2026			PED	Jan 22, 2025
	7456254	Jun 30, 2025	DP U-2588			
	7456254	Jun 30, 2025	DP U-2589			
	7456254	Jun 30, 2025	DP U-2590			
	7456254	Jun 30, 2025	DP U-3188			
	7456254	Jun 30, 2025	DP U-3189			
	7456254	Jun 30, 2025	DP U-3190			
	7456254*PED	Dec 30, 2025				
	7563871	Apr 15, 2024	DP			
	7563871*PED	Oct 15, 2024				
	7612176	Apr 13, 2025	DP U-2589			
	7612176	Apr 13, 2025	DP U-2590			
	7612176	Apr 13, 2025	DP U-3188			
	7612176	Apr 13, 2025	DP U-3189			
	7612176	Apr 13, 2025	DP U-3190			
	7612176*PED	Oct 13, 2025				
	8329648	Aug 18, 2026	U-2588			
	8329648	Aug 18, 2026	U-2589			
	8329648	Aug 18, 2026	U-2590			
	8329648	Aug 18, 2026	U-2593			
	8329648	Aug 18, 2026	U-2594			
	8329648	Aug 18, 2026	U-2595			
	8329648	Aug 18, 2026	U-2596			

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<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210 001	8329648	Aug 18, 2026	U-3188			
	8329648	Aug 18, 2026	U-3189			
	8329648	Aug 18, 2026	U-3190			
	8329648*PED	Feb 18, 2027				
	8361972	Mar 21, 2028	U-2588			
	8361972*PED	Sep 21, 2028				
	8431685	Apr 13, 2025	DP U-2588			
	8431685	Apr 13, 2025	DP U-2589			
	8431685	Apr 13, 2025	DP U-2590			
	8431685	Apr 13, 2025	DP U-2597			
	8431685	Apr 13, 2025	DP U-3188			
	8431685	Apr 13, 2025	DP U-3189			
	8431685	Apr 13, 2025	DP U-3190			
	8431685*PED	Oct 13, 2025				
	8461105	Apr 13, 2025	DP U-2588			
	8461105	Apr 13, 2025	DP U-2589			
	8461105	Apr 13, 2025	DP U-2590			
	8461105	Apr 13, 2025	DP U-2597			
	8461105	Apr 13, 2025	DP U-3188			
	8461105	Apr 13, 2025	DP U-3189			
	8461105	Apr 13, 2025	DP U-3190			
	8461105*PED	Oct 13, 2025				
	8501698	Jun 20, 2027	U-2588			
	8501698*PED	Dec 20, 2027				
	8895033	Oct 04, 2030	DP U-2589			
	8895033	Oct 04, 2030	DP U-2590			
	8895033	Oct 04, 2030	DP U-2597			
	8895033	Oct 04, 2030	DP U-2601			
	8895033	Oct 04, 2030	DP U-2602			
	8895033	Oct 04, 2030	DP U-3188			
	8895033	Oct 04, 2030	DP U-3189			
	8895033	Oct 04, 2030	DP U-3190			
	8895033*PED	Apr 04, 2031				
	8906851	Aug 18, 2026	U-2588			
	8906851	Aug 18, 2026	U-2589			
	8906851	Aug 18, 2026	U-2590			
	8906851	Aug 18, 2026	U-2593			
	8906851	Aug 18, 2026	U-2597			
	8906851	Aug 18, 2026	U-3188			
	8906851	Aug 18, 2026	U-3189			
	8906851	Aug 18, 2026	U-3190			
	8906851*PED	Feb 18, 2027				
	9238076	Apr 15, 2024	DP U-2588			
	9238076	Apr 15, 2024	DP U-2589			
	9238076	Apr 15, 2024	DP U-2590			
	9238076	Apr 15, 2024	DP U-2597			
	9238076	Apr 15, 2024	DP U-2599			
	9238076	Apr 15, 2024	DP U-3188			
	9238076	Apr 15, 2024	DP U-3189			

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<u>EXENATIDE SYNTHETIC - BYDUREON BCISE</u>						
N 209210	001	9238076	Apr 15, 2024	DP U-3190		
		9238076*PED	Oct 15, 2024			
		9884092	Aug 18, 2026	U-2588		
		9884092	Aug 18, 2026	U-2589		
		9884092	Aug 18, 2026	U-2590		
		9884092	Aug 18, 2026	U-2593		
		9884092	Aug 18, 2026	U-2594		
		9884092	Aug 18, 2026	U-2595		
		9884092	Aug 18, 2026	U-2596		
		9884092	Aug 18, 2026	U-2597		
		9884092	Aug 18, 2026	U-3188		
		9884092	Aug 18, 2026	U-3189		
		9884092	Aug 18, 2026	U-3190		
		9884092*PED	Feb 18, 2027			
<u>EZETIMIBE - ZETIA</u>						
N 021445	001	7612058	Oct 30, 2025	U-1027		
		7612058	Oct 30, 2025	U-1173		
		7612058*PED	Apr 30, 2026			
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	001	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	002	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	003	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>EZETIMIBE; ROSUVASTATIN CALCIUM - ROSZET</u>						
N 213072	004	10376470	May 01, 2033	DP U-3095		
		9763885	May 01, 2033	DP U-3095		
<u>FAMOTIDINE; IBUPROFEN - DUEXIS</u>						
N 022519	001	8067451	Jul 18, 2026	DP U-1196		
		8309127	Jul 18, 2026	DP		
		8318202	Jul 18, 2026	DP		
		8449910	Jul 18, 2026	DP		
		8501228	Jul 18, 2026	U-1196		
<u>FEBUXOSTAT - ULORIC</u>						
N 021856	001	7361676	Mar 08, 2024	DP		
		8372872	Sep 08, 2031	U-1346		
		9107912	Sep 08, 2031	U-1346		
<u>FEBUXOSTAT - ULORIC</u>						
N 021856	002	7361676	Mar 08, 2024	DP		
		8372872	Sep 08, 2031	U-1346		
		9107912	Sep 08, 2031	U-1346		

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<u>FEDRATINIB HYDROCHLORIDE - INREBIC</u>						
N 212327 001	10391094	Jun 04, 2032	DP U-2607		NCE	Aug 16, 2024
	11400092	Sep 24, 2039	U-3409		ODE-259	Aug 16, 2026
	7528143	Nov 16, 2031	DS DP			
	7825246	Dec 16, 2026	DS			
	8138199	Jun 30, 2028	U-2607			
<u>FENFLURAMINE HYDROCHLORIDE - FINTEPLA</u>						
N 212102 001	10452815	Jun 29, 2038	U-2859	Y	I-887	Mar 25, 2025
	10452815*PED	Dec 29, 2038			ODE-312	Jun 25, 2027
	10478441	May 03, 2033	U-2860		ODE-393	Mar 25, 2029
	10478441*PED	Nov 03, 2033			PED	Sep 25, 2025
	10478442	May 03, 2033	U-2860		PED	Dec 25, 2027
	10478442*PED	Nov 03, 2033			PED	Sep 25, 2029
	10603290	Aug 02, 2037	U-2861			
	10603290	Aug 02, 2037	U-3347			
	10603290*PED	Feb 02, 2038				
	10947183	Dec 20, 2036	DS DP			
	10947183*PED	Jun 20, 2037				
	11040018	Aug 02, 2037	U-2861			
	11040018	Aug 02, 2037	U-3347			
	11040018*PED	Feb 02, 2038				
	11406606	Aug 02, 2037	U-3406			
	11406606	Aug 02, 2037	U-3407			
	11406606*PED	Feb 02, 2038				
	11759440	Aug 02, 2037	U-3694			
	11759440*PED	Feb 02, 2038				
	11786487	Aug 02, 2037	U-3733			
	11786487*PED	Feb 02, 2038				
	9549909	May 03, 2033	U-2858			
	9549909*PED	Nov 03, 2033				
	9603814	May 03, 2033	U-2858			
	9603814*PED	Nov 03, 2033				
	9603815	May 03, 2033	U-2858			
	9603815*PED	Nov 03, 2033				
	9610260	May 03, 2033	U-2858			
	9610260*PED	Nov 03, 2033				
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>						
N 021695 004	8026281	Apr 22, 2025	U-1447			
	8026281	Apr 22, 2025	U-1448			
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>						
N 021695 005	8026281	Apr 22, 2025	U-1447			
	8026281	Apr 22, 2025	U-1448			
	9314447	May 31, 2033	DP U-1447			
	9314447	May 31, 2033	DP U-1448			
<u>FENOFIBRATE - FENOGLIDE</u>						
N 022118 001	7658944	Dec 09, 2024	DP			
	8124125	Oct 01, 2024	DP U-1234			
	8481078	Oct 01, 2024	DP U-1416			
	9173847	Oct 01, 2024	DP			

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<u>FENOFIBRATE - FENOGLIDE</u>						
N 022118	001	7658944	Dec 09, 2024	DP		
		8124125	Oct 01, 2024	DP U-1234		
		8481078	Oct 01, 2024	DP U-1416		
		9173847	Oct 01, 2024	DP		
<u>FENOFIBRATE - FENOGLIDE</u>						
N 022118	002	7658944	Dec 09, 2024	DP		
		8124125	Oct 01, 2024	DP U-1234		
		8481078	Oct 01, 2024	DP U-1416		
		9173847	Oct 01, 2024	DP		
<u>FENOFIBRIC ACID - FIBRICOR</u>						
N 022418	001	7569612	Aug 20, 2027	U-1000		
		7741373	Aug 20, 2027	U-1059		
		7741374	Aug 20, 2027	U-1060		
		7741374	Aug 20, 2027	U-1061		
		7915247	Aug 20, 2027	U-1000		
		7915247	Aug 20, 2027	U-1059		
		7915247	Aug 20, 2027	U-1061		
<u>FENOFIBRIC ACID - FIBRICOR</u>						
N 022418	002	7569612	Aug 20, 2027	U-1000		
		7741373	Aug 20, 2027	U-1059		
		7741374	Aug 20, 2027	U-1060		
		7741374	Aug 20, 2027	U-1061		
		7915247	Aug 20, 2027	U-1000		
		7915247	Aug 20, 2027	U-1059		
		7915247	Aug 20, 2027	U-1061		
<u>FENTANYL - SUBSYS</u>						
N 202788	001	10016403	Jan 25, 2027	DP		
		10610523	Jan 25, 2027	DP		
		8486972	Apr 27, 2030	DP		
		8486973	Apr 27, 2030	U-55		
		8835459	Jan 25, 2027	DP		
		8835460	Jan 25, 2027	DP U-55		
		9241935	Jan 25, 2027	DP		
		9289387	Jan 25, 2027	DP U-55		
		9642797	Jan 25, 2027	DP U-55		
		9642844	Jan 25, 2027	DP		
<u>FENTANYL - SUBSYS</u>						
N 202788	002	10016403	Jan 25, 2027	DP		
		10610523	Jan 25, 2027	DP		
		8486972	Apr 27, 2030	DP		
		8486973	Apr 27, 2030	U-55		
		8835460	Jan 25, 2027	DP U-55		
		9241935	Jan 25, 2027	DP		
		9289387	Jan 25, 2027	DP U-55		
		9642797	Jan 25, 2027	DP U-55		
		9642844	Jan 25, 2027	DP		

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<u>FENTANYL - SUBSYS</u>						
N 202788 003	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 004	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 005	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 006	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			
	9642797	Jan 25, 2027	DP U-55			
	9642844	Jan 25, 2027	DP			
<u>FENTANYL - SUBSYS</u>						
N 202788 007	10016403	Jan 25, 2027	DP			
	10610523	Jan 25, 2027	DP			
	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			

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<u>FENTANYL - SUBSYS</u>						
N 202788	007	8835460	Jan 25, 2027	DP U-55		
		9241935	Jan 25, 2027	DP		
		9289387	Jan 25, 2027	DP U-55		
		9642797	Jan 25, 2027	DP U-55		
		9642844	Jan 25, 2027	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	001	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	002	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	003	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	004	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - FENTORA</u>						
N 021947	005	7862832	Jun 15, 2028	DP		
		7862833	Jun 15, 2028	DP		
		8092832	Dec 30, 2024	DP		
		8119158	Dec 30, 2024	DP		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	001	9597288	Jul 23, 2027	DP U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	002	9597288	Jul 23, 2027	DP U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	003	9597288	Jul 23, 2027	DP U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	004	9597288	Jul 23, 2027	DP U-767		
<u>FENTANYL CITRATE - ONSOLIS</u>						
N 022266	005	9597288	Jul 23, 2027	DP U-767		
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569	001	8216604	Oct 03, 2024	U-767		
		8889176	Jan 16, 2024	U-767		
		9078814	Jan 08, 2024	DP		
		9731869	Jan 26, 2032	DP		
		9814705	Jan 08, 2024	DP		

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<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569 002	8216604	Oct 03, 2024	U-767			
	8889176	Jan 16, 2024	U-767			
	9078814	Jan 08, 2024	DP			
	9731869	Jan 26, 2032	DP			
	9814705	Jan 08, 2024	DP			
<u>FENTANYL CITRATE - LAZANDA</u>						
N 022569 003	9731869	Jan 26, 2032	DP			
	9814705	Jan 08, 2024	DP			
<u>FENTANYL HYDROCHLORIDE - IONSYS</u>						
N 021338 001	6975902	Apr 01, 2024	DP			
	8301238	Sep 30, 2031	DP			
	8428708	May 21, 2032	U-736			
	8428709	Jun 11, 2032	DP U-736			
	8781571	Mar 31, 2032	DP U-736			
	9095706	Feb 03, 2033	DP			
	9364656	Sep 30, 2031	U-736			
	9731121	Oct 17, 2031	DP			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 001	11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435		NPP	Nov 19, 2024
	11433091	Jan 08, 2027	U-3436			
	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			
	11433091	Jan 08, 2027	U-3634			
	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2024	DS DP			
	7754702	Feb 15, 2028	U-1432			
	7754702	Feb 15, 2028	U-2555			
	7754702	Feb 15, 2028	U-2556			
	7754702	Feb 15, 2028	U-2557			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-1620			
	8895612	Jan 08, 2027	U-3050			
	8895612	Jan 08, 2027	U-3051			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 002	11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435		NPP	Nov 19, 2024
	11433091	Jan 08, 2027	U-3436			

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<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 002	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			
	11433091	Jan 08, 2027	U-3634			
	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2024	DS DP			
	7754702	Feb 15, 2028	U-2555			
	7754702	Feb 15, 2028	U-2556			
	7754702	Feb 15, 2028	U-2557			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-1620			
	8895612	Jan 08, 2027	U-3050			
	8895612	Jan 08, 2027	U-3051			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			
<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 003	11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435		NPP	Nov 19, 2024
	11433091	Jan 08, 2027	U-3436		NS	Apr 28, 2024
	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			
	11433091	Jan 08, 2027	U-3634			
	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2024	DS DP			
	7754702	Feb 15, 2028	U-2555			
	7754702	Feb 15, 2028	U-2556			
	7754702	Feb 15, 2028	U-2557			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-1620			
	8895612	Jan 08, 2027	U-3050			
	8895612	Jan 08, 2027	U-3051			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			

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<u>FERRIC CARBOXYMALTOSE - INJECTAFER</u>						
N 203565 004	11364260	Jan 08, 2027	U-3637		I-915	May 31, 2026
	11433091	Jan 08, 2027	U-3435			
	11433091	Jan 08, 2027	U-3436			
	11433091	Jan 08, 2027	U-3437			
	11433091	Jan 08, 2027	U-3438			
	11433091	Jan 08, 2027	U-3634			
	11478502	Jan 08, 2027	U-3472			
	11478502	Jan 08, 2027	U-3473			
	11478502	Jan 08, 2027	U-3474			
	7612109	Feb 05, 2024	DS DP			
	7754702	Feb 15, 2028	U-3312			
	7754702	Feb 15, 2028	U-3313			
	7754702	Feb 15, 2028	U-3314			
	7754702	Feb 15, 2028	U-3636			
	8895612	Jan 08, 2027	U-3115			
	8895612	Jan 08, 2027	U-3116			
	8895612	Jan 08, 2027	U-3315			
	8895612	Jan 08, 2027	U-3316			
	8895612	Jan 08, 2027	U-3635			
<u>FERRIC CITRATE - AURYXIA</u>						
N 205874 001	10300039	Jul 21, 2030	U-2549			
	7767851	Feb 18, 2024	DS DP			
	8093423	Apr 21, 2026	U-1577			
	8299298	Feb 18, 2024	DP			
	8338642	Feb 18, 2024	DS DP U-1577			
	8609896	Feb 18, 2024	DP			
	8754257	Feb 18, 2024	DP			
	8754258	Feb 18, 2024	DP			
	8846976	Feb 18, 2024	U-1577			
	8901349	Feb 18, 2024	U-1577			
	9050316	Feb 18, 2024	U-1577			
	9328133	Feb 18, 2024	DS DP U-1577			
	9387191	Jul 21, 2030	DP			
	9757416	Feb 18, 2024	DS DP U-1577			
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171 001	10414831	Mar 25, 2029	DS DP			
	11633489	Jun 22, 2036	U-3594			
	8815301	Aug 14, 2029	DS DP U-2734			
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171 002	10414831	Mar 25, 2029	DS DP			
	11633489	Jun 22, 2036	U-3594			
	8815301	Aug 14, 2029	DS DP U-2734			
<u>FERRIC DERISOMALTOSE - MONOFERRIC</u>						
N 208171 003	10414831	Mar 25, 2029	DS DP			
	11633489	Jun 22, 2036	U-3594			
	8815301	Aug 14, 2029	DS DP U-2734			

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<u>FERRIC MALTOLE - ACCRUFER</u>						
N 212320	001 10179120	Jan 06, 2035		U-2603	NCE	Jul 25, 2024
	9248148	Mar 29, 2031		U-2603		
	9802973	Oct 23, 2035	DS DP	U-2603		
<u>FERRIC OXYHYDROXIDE - VELPHORO</u>						
N 205109	001 10624855	Nov 26, 2034		DP		
	10624855*PED	May 26, 2035				
	10682376	Nov 13, 2028		DP		
	10682376*PED	May 13, 2029				
	10695367	Nov 13, 2028		DP		
	10695367*PED	May 13, 2029				
	10925896	Nov 13, 2028		DP		
	10925896*PED	May 13, 2029				
	10925897	Nov 13, 2028		DP		
	10925897*PED	May 13, 2029				
	10933090	Nov 13, 2028		DP		
	10933090*PED	May 13, 2029				
	11013761	Nov 13, 2028		DP		
	11013761*PED	May 13, 2029				
	11013762	Nov 13, 2028		DP		
	11013762*PED	May 13, 2029				
	11234938	Nov 26, 2034		DP		
	11234938*PED	May 26, 2035				
	11446252	Nov 26, 2034		DP		
	11446252*PED	May 26, 2035				
	9561251	Jan 23, 2030		DP	U-1468	
	9561251*PED	Jul 23, 2030				
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u>						
N 206317	001 7816404	Apr 17, 2029		DP	U-1656	
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u>						
N 208551	001 7816404	Apr 17, 2029		U-1656		
	7857977	Sep 08, 2027		U-1656		
<u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC AVNU</u>						
N 212860	001 7816404	Apr 17, 2029	DS	U-2801		
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030	001 7807715	Jun 07, 2027		DP U-913	I-861	Jun 17, 2024
	8088398	Jun 07, 2027		DP U-913	PED	Dec 17, 2024
	8501723	Jun 07, 2027		DP		
	8501723*PED	Dec 07, 2027				
<u>FESOTERODINE FUMARATE - TOVIAZ</u>						
N 022030	002 7807715	Jun 07, 2027		DP U-913	I-861	Jun 17, 2024
	8088398	Jun 07, 2027		DP U-913	PED	Dec 17, 2024
	8501723	Jun 07, 2027		DP		
	8501723*PED	Dec 07, 2027				
<u>FEXINIDAZOLE - FEXINIDAZOLE</u>						
N 214429	001				NCE	Jul 16, 2026
					ODE-359	Jul 16, 2028

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<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u>						
N 201373	001 8933097	Aug 02, 2030	DP			
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u>						
N 201373	002 8933097	Aug 02, 2030	DP			
<u>FEZOLINETANT - VEOZAH</u>						
N 216578	001 10836768	Mar 28, 2034	U-3621		NCE	May 12, 2028
	8871761	Apr 04, 2031	DS			
	9422299	Mar 28, 2034	DS DP	U-3622		
	9987274	Mar 28, 2034		U-3622		
<u>FIDAXOMICIN - DIFICID</u>						
N 201699	001 7378508	Jul 31, 2027	DS DP		ODE-367	Jan 24, 2027
	7863249	Jul 31, 2027	DS DP			
	7906489	Mar 04, 2027		U-2741		
	7906489	Mar 04, 2027		U-319		
	7906489*PED	Sep 04, 2027				
	8586551*PED	Jan 15, 2024				
	8859510	Jul 31, 2027		U-2741		
	8859510	Jul 31, 2027		U-319		
	8859510*PED	Jan 31, 2028				
<u>FIDAXOMICIN - DIFICID</u>						
N 213138	001 7378508	Jul 31, 2027	DS DP		ODE-367	Jan 24, 2027
	7378508*PED	Jan 31, 2028				
	7863249	Jul 31, 2027		DP		
	7863249*PED	Jan 31, 2028				
	7906489	Mar 04, 2027		U-2741		
	7906489*PED	Sep 04, 2027				
	8586551*PED	Jan 23, 2024				
	8859510	Jul 31, 2027		U-2741		
	8859510*PED	Jan 31, 2028				
	9808530	May 28, 2034		DP		
	9808530*PED	Nov 28, 2034				
<u>FINAFLOXACIN - XTORO</u>						
N 206307	001 8536167	Aug 08, 2031		U-1679		
	9119859	Jul 02, 2030		U-1679		
	9504691	Nov 21, 2033		DP U-1679		
	9993483	Jul 02, 2030		DP		
<u>FINERENONE - KERENDIA</u>						
N 215341	001 8436180	Apr 12, 2029	DS DP		M-279	Sep 01, 2025
					NCE	Jul 09, 2026
<u>FINERENONE - KERENDIA</u>						
N 215341	002 8436180	Apr 12, 2029	DS DP		M-279	Sep 01, 2025
					NCE	Jul 09, 2026
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527	001 10543179	Dec 25, 2027		U-2719		
	8324283	Mar 29, 2026		DP		
	8324283*PED	Sep 29, 2026				
	9187405	Jun 25, 2027		U-2613		

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<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527	001	9187405*PED	Dec 25, 2027			
<u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u>						
N 022527	002	9592208	Mar 30, 2032	DP U-2315		
		9592208*PED	Sep 30, 2032			
<u>FINGOLIMOD LAURYL SULFATE - TASCENSO ODT</u>						
N 214962	001	10555902	Jan 19, 2036	U-3268		
		10925829	Jan 19, 2036	DP		
		9925138	Jan 19, 2036	DP		
<u>FINGOLIMOD LAURYL SULFATE - TASCENSO ODT</u>						
N 214962	002	10555902	Jan 19, 2036	U-3493		
		10925829	Jan 19, 2036	DP		
		9925138	Jan 19, 2036	DP		
<u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u>						
N 210589	001	10350186	Nov 05, 2024	U-2585	ODE-202	Jul 27, 2025
		9566260	Jul 11, 2025	DP U-2366		
		9629821	Jul 11, 2025	DP U-2367		
<u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u>						
N 210589	002	10350186	Nov 05, 2024	U-2585	ODE-202	Jul 27, 2025
		9566260	Jul 11, 2025	DP U-2366		
		9629821	Jul 11, 2025	DP U-2367		
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	001				NPP	Mar 22, 2025
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	002				NPP	Mar 22, 2025
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	003				NPP	Mar 22, 2025
<u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u>						
N 207648	004				NPP	Mar 22, 2025
<u>FLIBANSERIN - ADDYI</u>						
N 022526	001	7151103	May 09, 2028	U-1734		
<u>FLORBETABEN F-18 - NEURACEQ</u>						
N 204677	001	7807135	Mar 18, 2029	DS DP U-1497		
<u>FLORBETAPIR F-18 - AMYVID</u>						
N 202008	001	7687052	Apr 30, 2027	DS DP		
		8506929	Apr 30, 2027	DS DP U-1423		
<u>FLORBETAPIR F-18 - AMYVID</u>						
N 202008	002	7687052	Apr 30, 2027	DS DP		
		8506929	Apr 30, 2027	DS DP U-1423		
<u>FLORBETAPIR F-18 - AMYVID</u>						
N 202008	003	7687052	Apr 30, 2027	DS DP		
		8506929	Apr 30, 2027	DS DP U-1423		
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123	001	8932557	May 26, 2032	DS	NCE	May 28, 2025

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<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123 002	8932557	May 26, 2032	DS		NCE	May 28, 2025
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123 003	8932557	May 26, 2032	DS			
<u>FLORTAUCIPIR F-18 - TAUVID</u>						
N 212123 004	8932557	May 26, 2032	DS			
<u>FLOTUFOLASTAT F-18 GALLIUM - POSLUMA</u>						
N 216023 001	11413360	Nov 22, 2038	DS	U-3614	NCE	May 25, 2028
	11413360	Nov 22, 2038	DS	U-3615		
<u>FLUCICLOVINE F-18 - AXUMIN</u>						
N 208054 001	10010632	Nov 28, 2026	DP			
	10124079	Dec 30, 2035		U-2450		
	10716868	Dec 30, 2035		U-2450		
	10933147	Dec 30, 2035		U-2450		
	10953112	Nov 28, 2026		U-1879		
	10967077	Dec 30, 2035		U-2450		
	9387266	Nov 28, 2026		U-1879		
<u>FLUOCINOLONE ACETONIDE - ILUVIEN</u>						
N 201923 001	8871241	Aug 12, 2027	DP			
<u>FLUOCINOLONE ACETONIDE - YUTIQ</u>						
N 210331 001	7998108	Jan 12, 2028	DP	U-3410		
	8871241	Aug 12, 2027	DP			
<u>FLUORESCEIN SODIUM - FLUORESCEIN SODIUM</u>						
A 215709 001					CGT	Apr 21, 2024
<u>FLUORODOPA F-18 - FLUORODOPA F18</u>						
N 200655 001					NCE	Oct 10, 2024
					W	Oct 10, 2024
<u>FLUOROESTRADIOL F-18 - CERIANNA</u>						
N 212155 001					NCE	May 20, 2025
<u>FLUTEMETAMOL F-18 - VIZAMYL</u>						
N 203137 001	7270800	Sep 03, 2025	DS DP	U-336		
	8236282	May 21, 2024	DS DP			
	8916131	Sep 16, 2028	DP			
<u>FLUTEMETAMOL F-18 - VIZAMYL</u>						
N 203137 002	7270800	Sep 03, 2025	DS DP	U-336		
	8236282	May 21, 2024	DS DP			
	8916131	Sep 16, 2028	DP			
<u>FLUTICASONE FUROATE - FLONASE SENSIMIST ALLERGY RELIEF</u>						
N 022051 002	8062264	Apr 05, 2026	DP			
	8147461	Oct 15, 2028	DP			
	8347879	Jul 15, 2028	DP			
	8752543	Apr 05, 2026	DP			
	9320862	Nov 06, 2024	DP			
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625 001	8201556	Feb 05, 2029	DP			
	8746242	Oct 11, 2030	DP			

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<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625 001	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625 002	8113199	Oct 23, 2027	DP			
	8113199*PED	Apr 23, 2028				
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8201556	Feb 05, 2029	DP			
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
<u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u>						
N 205625 003	8113199	Oct 23, 2027	DP		M-290	Mar 01, 2026
	8113199*PED	Apr 23, 2028				
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8201556	Feb 05, 2029	DP			
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482 001	11090294	Nov 29, 2030		U-3202		
	7439393	May 21, 2025	DS DP	U-2127		
	7439393	May 21, 2025	DS DP	U-2957		
	7439393*PED	Nov 21, 2025				
	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	8113199	Oct 23, 2027	DP			
	8113199*PED	Apr 23, 2028				
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8183257	Jul 27, 2025		U-2128		
	8183257	Jul 27, 2025		U-2129		
	8309572	Apr 27, 2025		U-2129		
	8511304	Jun 14, 2027	DP	U-2954		
	8511304*PED	Dec 14, 2027				
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482	001	9333310*PED	Apr 02, 2028			
		9750726	Nov 29, 2030	DP		
<u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u>						
N 209482	002	7439393	May 21, 2025	DS DP	U-2957	
		7439393*PED	Nov 21, 2025			
		7488827	Dec 18, 2027	DS DP		
		7498440	Apr 27, 2025	DS DP		
		8113199	Oct 23, 2027	DP		
		8113199*PED	Apr 23, 2028			
		8161968	Feb 05, 2028	DP		
		8161968*PED	Aug 05, 2028			
		8183257	Jul 27, 2025		U-2129	
		8309572	Apr 27, 2025		U-2129	
		8511304	Jun 14, 2027	DP	U-2954	
		8511304*PED	Dec 14, 2027			
		8534281	Mar 08, 2030	DP		
		8534281*PED	Sep 08, 2030			
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			
		9750726	Nov 29, 2030	DP		
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275	001	11116721	Feb 26, 2029	DP	U-1401	NPP May 13, 2026
		11116721	Feb 26, 2029	DP	U-1691	PED Nov 13, 2026
		11116721	Feb 26, 2029	DP	U-3623	
		11116721*PED	Aug 26, 2029			
		7439393	May 21, 2025	DS DP	U-1401	
		7439393	May 21, 2025	DS DP	U-1691	
		7439393	May 21, 2025	DS DP	U-2099	
		7439393	May 21, 2025	DS DP	U-2100	
		7439393	May 21, 2025	DS DP	U-3623	
		7439393*PED	Nov 21, 2025			
		8113199	Oct 23, 2027	DP		
		8113199*PED	Apr 23, 2028			
		8161968	Feb 05, 2028	DP		
		8161968*PED	Aug 05, 2028			
		8511304	Jun 14, 2027	DP	U-1424	
		8511304	Jun 14, 2027	DP	U-1691	
		8511304	Jun 14, 2027	DP	U-3623	
		8511304*PED	Dec 14, 2027			
		8534281	Mar 08, 2030	DP		
		8534281*PED	Sep 08, 2030			
		8746242	Oct 11, 2030	DP		
		8746242*PED	Apr 11, 2031			
		9333310	Oct 02, 2027	DP		
		9333310*PED	Apr 02, 2028			

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<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275 002	11116721	Feb 26, 2029	DP U-1691			
	11116721	Feb 26, 2029	DP U-3623			
	11116721*PED	Aug 26, 2029				
	7439393	May 21, 2025	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-2099			
	7439393	May 21, 2025	DS DP U-2100			
	7439393	May 21, 2025	DS DP U-3623			
	7439393*PED	Nov 21, 2025				
	8113199	Oct 23, 2027	DP			
	8113199*PED	Apr 23, 2028				
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8511304	Jun 14, 2027	DP U-1691			
	8511304	Jun 14, 2027	DP U-3623			
	8511304*PED	Dec 14, 2027				
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
<u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u>						
N 204275 003	11116721	Feb 26, 2029	DP U-3623		NS	May 13, 2026
	11116721*PED	Aug 26, 2029			PED	Nov 13, 2026
	7439393	May 21, 2025	DS DP U-3623			
	7439393*PED	Nov 21, 2025				
	8113199	Oct 23, 2027	DP			
	8113199*PED	Apr 23, 2028				
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8511304	Jun 14, 2027	DP U-3623			
	8511304*PED	Dec 14, 2027				
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433 001	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026				
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
N 021433 002	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 001	10022510	May 18, 2031	DP		NPP	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			

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<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 001	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 002	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 003	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				

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<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 003	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 004	10022510	May 18, 2031	DP		NPP	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				

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<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 004	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 005	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				

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<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 005	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 006	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				

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<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 006	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
<u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u>						
N 208798 007	10022510	May 18, 2031	DP		NS	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				

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<u>FLUTICASONE PROPIONATE - ARMONAIR DIGIHALER</u>						
N 208798 008	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11357935*PED	Mar 24, 2039				
	11439777	May 24, 2040	DP			
	11439777*PED	Nov 24, 2040				
	11464923	Jun 19, 2040	DP			
	11464923*PED	Dec 19, 2040				
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Nov 08, 2030	DP			
	8714149*PED	May 08, 2031				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - XHANCE</u>						
N 209022	001	10076614	Oct 20, 2034	DP		
		10076615	Jul 30, 2029	U-2133		
		10124132	Mar 06, 2027	DP U-2133		
		10179216	Jul 08, 2033	DP U-2133		
		10252010	Feb 07, 2031	DP		
		10300229	Jul 07, 2035	DP U-2133		
		10478574	Nov 04, 2033	U-2133		
		11033696	May 20, 2033	DP		
		11554229	Feb 23, 2036	U-2133		
		11602603	Oct 27, 2028	DP U-2133		
		7975690	Dec 29, 2025	U-2133		
		8522778	Apr 20, 2024	DP		
		8550073	Oct 22, 2029	DP		
		8978647	Dec 06, 2030	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N 021254	001	7500444	Feb 26, 2026	DP		
		7500444*PED	Aug 26, 2026			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N 021254	003	7500444	Feb 26, 2026	DP		
		7500444*PED	Aug 26, 2026			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	001	10022510	May 18, 2031	DP	M-61	Jul 09, 2024
		10022510*PED	Nov 18, 2031		PED	Jan 09, 2025
		10124131	May 18, 2031	DP		
		10124131*PED	Nov 18, 2031			
		10195375	Feb 14, 2031	DP		
		10195375*PED	Aug 14, 2031			
		10561808	Jan 01, 2032	DP		
		10561808*PED	Jul 01, 2032			
		10765820	May 19, 2025	DP		
		10765820*PED	Nov 19, 2025			
		8651103	Mar 26, 2028	DP		
		8651103*PED	Sep 26, 2028			
		8714149	Feb 25, 2032	DP		
		8714149*PED	Aug 25, 2032			
		8978966	Jan 13, 2032	DP		
		8978966*PED	Jul 13, 2032			
		9066957	Oct 06, 2034	DP U-645		
		9066957*PED	Apr 06, 2035			
		9216260	Jun 28, 2031	DP		
		9216260*PED	Dec 28, 2031			
		9415008	Oct 06, 2034	DP U-645		
		9415008*PED	Apr 06, 2035			
		9463288	May 19, 2025	DP		
		9463288*PED	Nov 19, 2025			
		9616024	Sep 01, 2024	DP		
		9616024*PED	Mar 01, 2025			
		9731087	May 18, 2031	DP		

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	001	9731087*PED				Nov 18, 2031
		9987229		DP		Sep 01, 2024
		9987229*PED				Mar 01, 2025
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	002	10022510		DP		May 18, 2031
		10022510*PED				Nov 18, 2031
		10124131		DP		May 18, 2031
		10124131*PED				Nov 18, 2031
		10195375		DP		Feb 14, 2031
		10195375*PED				Aug 14, 2031
		10561808		DP		Jan 01, 2032
		10561808*PED				Jul 01, 2032
		10765820		DP		May 19, 2025
		10765820*PED				Nov 19, 2025
		8651103		DP		Mar 26, 2028
		8651103*PED				Sep 26, 2028
		8714149		DP		Feb 25, 2032
		8714149*PED				Aug 25, 2032
		8978966		DP		Jan 13, 2032
		8978966*PED				Jul 13, 2032
		9066957		DP U-645		Oct 06, 2034
		9066957*PED				Apr 06, 2035
		9216260		DP		Jun 28, 2031
		9216260*PED				Dec 28, 2031
		9463288		DP		May 19, 2025
		9463288*PED				Nov 19, 2025
		9616024		DP		Sep 01, 2024
		9616024*PED				Mar 01, 2025
		9731087		DP		May 18, 2031
		9731087*PED				Nov 18, 2031
		9987229		DP		Sep 01, 2024
		9987229*PED				Mar 01, 2025
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799	003	10022510		DP		May 18, 2031
		10022510*PED				Nov 18, 2031
		10124131		DP		May 18, 2031
		10124131*PED				Nov 18, 2031
		10195375		DP		Feb 14, 2031
		10195375*PED				Aug 14, 2031
		10561808		DP		Jan 01, 2032
		10561808*PED				Jul 01, 2032
		10765820		DP		May 19, 2025
		10765820*PED				Nov 19, 2025
		8651103		DP		Mar 26, 2028
		8651103*PED				Sep 26, 2028
		8714149		DP		Feb 25, 2032
		8714149*PED				Aug 25, 2032
		8978966		DP		Jan 13, 2032
		8978966*PED				Jul 13, 2032

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u>						
N 208799 003	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9987229	Sep 01, 2024	DP			
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 004	10022510	May 18, 2031	DP		M-61	Jul 09, 2024
	10022510*PED	Nov 18, 2031			PED	Jan 09, 2025
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				
	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 004	9216260*PED	Dec 28, 2031				
	9415008	Oct 06, 2034	DP U-645			
	9415008*PED	Apr 06, 2035				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
	9987229	Sep 01, 2024	DP			
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 005	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			
	8978966*PED	Jul 13, 2032				

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 005	9066957	Oct 06, 2034	DP U-645			
	9066957*PED	Apr 06, 2035				
	9216260	Jun 28, 2031	DP			
	9216260*PED	Dec 28, 2031				
	9463288	May 19, 2025	DP			
	9463288*PED	Nov 19, 2025				
	9616024	Sep 01, 2024	DP			
	9616024*PED	Mar 01, 2025				
	9731087	May 18, 2031	DP			
	9731087*PED	Nov 18, 2031				
	9782550	Aug 28, 2035	DP			
	9782550*PED	Feb 28, 2036				
	9782551	Aug 28, 2035	DP			
	9782551*PED	Feb 28, 2036				
	9987229	Sep 01, 2024	DP			
	9987229*PED	Mar 01, 2025				
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799 006	10022510	May 18, 2031	DP			
	10022510*PED	Nov 18, 2031				
	10124131	May 18, 2031	DP			
	10124131*PED	Nov 18, 2031				
	10195375	Feb 14, 2031	DP			
	10195375*PED	Aug 14, 2031				
	10561808	Jan 01, 2032	DP			
	10561808*PED	Jul 01, 2032				
	10569034	Aug 16, 2036	DP			
	10569034*PED	Feb 16, 2037				
	10765820	May 19, 2025	DP			
	10765820*PED	Nov 19, 2025				
	10918816	Dec 14, 2035	DP			
	10918816*PED	Jun 14, 2036				
	11000653	Dec 18, 2038	DP			
	11000653*PED	Jun 18, 2039				
	11173259	Jul 06, 2040	DP			
	11173259*PED	Jan 06, 2041				
	11266796	Feb 22, 2041	DP			
	11266796*PED	Aug 22, 2041				
	11344685	Sep 26, 2039	DP			
	11344685*PED	Mar 26, 2040				
	11351317	Feb 10, 2038	DP			
	11351317*PED	Aug 10, 2038				
	11357935	Sep 24, 2038	DP			
	11439777	May 24, 2040	DP			
	11464923	Jun 19, 2040	DP			
	8651103	Mar 26, 2028	DP			
	8651103*PED	Sep 26, 2028				
	8714149	Feb 25, 2032	DP			
	8714149*PED	Aug 25, 2032				
	8978966	Jan 13, 2032	DP			

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<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO DIGIHALER</u>						
N 208799	006	8978966*PED	Jul 13, 2032			
		9066957	Oct 06, 2034	DP	U-645	
		9066957*PED	Apr 06, 2035			
		9216260	Jun 28, 2031	DP		
		9216260*PED	Dec 28, 2031			
		9463288	May 19, 2025	DP		
		9463288*PED	Nov 19, 2025			
		9616024	Sep 01, 2024	DP		
		9616024*PED	Mar 01, 2025			
		9731087	May 18, 2031	DP		
		9731087*PED	Nov 18, 2031			
		9782550	Aug 28, 2035	DP		
		9782550*PED	Feb 28, 2036			
		9782551	Aug 28, 2035	DP		
		9782551*PED	Feb 28, 2036			
		9987229	Sep 01, 2024	DP		
		9987229*PED	Mar 01, 2025			
<u>FOMEPIZOLE - ANTIZOL</u>						
N 020696	001	7553863	Jun 30, 2027	DS	DP	
<u>FORMOTEROL FUMARATE; GLYCOPYRROLATE - BEVESPI AEROSPHERE</u>						
N 208294	001	10716753	May 28, 2030	DP	U-2889	
		8324266	May 28, 2030		U-2889	
		8703806	May 28, 2030		U-2889	
		8808713	May 28, 2030	DP	U-2889	
		8815258	Mar 17, 2031		U-2889	
		9415009	May 28, 2030		U-2889	
		9463161	May 28, 2030	DP	U-2889	
<u>FOSAPREPITANT DIMEGLUMINE - EMEND</u>						
N 022023	001				D-186	May 02, 2025
<u>FOSAPREPITANT DIMEGLUMINE - EMEND</u>						
N 022023	002				D-186	May 02, 2025
<u>FOSAPREPITANT DIMEGLUMINE - FOCINVEZ</u>						
N 216686	001	11065265	Jan 11, 2039	DP		
<u>FOSDENOPTERIN HYDROBROMIDE - NULIBRY</u>						
N 214018	001	7504095	Jan 31, 2025	DP	U-3092	
					M-286	Oct 27, 2025
					NCE	Feb 26, 2026
					ODE-342	Feb 26, 2028
<u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	001	10208073	May 23, 2032		U-2301	
		10624911	Jun 02, 2037	DP		
		10717721	May 23, 2032	DS		
		10828297	Dec 17, 2030		U-2301	
		11312698	May 23, 2032	DS	DP	
		11529362	Jun 02, 2037	DP		
		8426450	May 23, 2032	DS	DP	
		8895586	May 23, 2032		U-2301	
		9186357	Nov 18, 2030		U-2301	

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<u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	001 9403772	May 23, 2032	DS	U-2301		
	9908907	May 23, 2032	DS DP			
<u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 210493	002 10208073	May 23, 2032		U-2301		
	10624911	Jun 02, 2037	DP			
	10717721	May 23, 2032	DS			
	10828297	Dec 17, 2030		U-2301		
	11312698	May 23, 2032	DS DP			
	11529362	Jun 02, 2037	DP			
	8426450	May 23, 2032	DS DP			
	8895586	May 23, 2032		U-2301		
	9186357	Nov 18, 2030		U-2301		
	9403772	May 23, 2032	DS	U-2301		
	9908907	May 23, 2032	DS DP			
<u>FOSPHENYTOIN SODIUM - SESQUIENT</u>						
N 210864	001 7635773	Mar 13, 2029		DP		
	8410077	Mar 13, 2029		DP		
	9200088	Mar 13, 2029		DP		
	9493582	Feb 27, 2033		DP		
	9750822	Mar 13, 2029		DP		
<u>FOSPHENYTOIN SODIUM - SESQUIENT</u>						
N 210864	002 7635773	Mar 13, 2029		DP		
	8410077	Mar 13, 2029		DP		
	9200088	Mar 13, 2029		DP		
	9493582	Feb 27, 2033		DP		
	9750822	Mar 13, 2029		DP		
<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299	001 7449458	Sep 04, 2031	DS		ODE-174	Apr 17, 2025
	7538108	Mar 28, 2026	DS	U-2294		
	7989448	Jun 12, 2026	DS	U-2294		
	8163902	Jun 17, 2026	DS	U-2294		
	8211889	Jan 19, 2026	DS			
	8263122	Nov 24, 2030		DP		
	8445485	Jun 17, 2026		DP		
	8652492	Nov 06, 2028		DP		
	8771648	Jul 27, 2032		DP		
	8912170	Jun 17, 2026		U-2294		
	8951504	Jul 27, 2032		U-2294		
	9266912	Jan 19, 2026		U-2294		
	9283238	Jun 17, 2026		U-2294		
	RE48898	Jan 19, 2026		DP		
<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299	002 7449458	Sep 04, 2031	DS		ODE-174	Apr 17, 2025
	7538108	Mar 28, 2026	DS	U-2294		
	7989448	Jun 12, 2026	DS	U-2294		
	8163902	Jun 17, 2026	DS	U-2294		
	8211889	Jan 19, 2026	DS			
	8263122	Nov 24, 2030		DP		

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<u>FOSTAMATINIB DISODIUM - TAVALISSE</u>						
N 209299	002 8445485	Jun 17, 2026	DP			
	8652492	Nov 06, 2028	DP			
	8771648	Jul 27, 2032	DP			
	8912170	Jun 17, 2026	U-2294			
	8951504	Jul 27, 2032	U-2294			
	9266912	Jan 19, 2026	U-2294			
	9283238	Jun 17, 2026	U-2294			
	RE48898	Jan 19, 2026	DP			
<u>FOSTEMSAVIR TROMETHAMINE - RUKOBIA</u>						
N 212950	001 7745625	Nov 19, 2027	DS		NCE	Jul 02, 2025
	8168615	Feb 25, 2025	DP			
	8461333	Feb 25, 2025	DS			
<u>FRUQUINTINIB - FRUZAOLA</u>						
N 217564	001 10519142	Sep 07, 2035	DS DP U-3753			
	11046674	Sep 07, 2035	U-3753			
	7829574	May 09, 2028	DS DP			
	8212033	May 09, 2028	U-3753			
<u>FRUQUINTINIB - FRUZAOLA</u>						
N 217564	002 10519142	Sep 07, 2035	DS DP U-3753			
	11046674	Sep 07, 2035	U-3753			
	7829574	May 09, 2028	DS DP			
	8212033	May 09, 2028	U-3753			
<u>FULVESTRANT - FULVESTRANT</u>						
N 210326	001 10188663	Feb 14, 2034	DP U-2540			
	9271990	May 17, 2034	DP U-2540			
	9833459	Feb 14, 2034	DP U-2540			
<u>FUROSEMIDE - FUROSCIX</u>						
N 209988	001 10272064	Apr 03, 2034	DP			
	11433044	Apr 03, 2034	U-3462			
	9884039	Apr 03, 2034	U-3462			
<u>FUTIBATINIB - LYTGOBI</u>						
N 214801	001 10434103	Mar 31, 2036	DS DP		NCE	Sep 30, 2027
	11833151	Nov 05, 2039	DP		ODE-410	Sep 30, 2029
	9108973	Feb 23, 2033	DS DP U-3456			
<u>GABAPENTIN - GRALISE</u>						
N 022544	001 7438927	Feb 26, 2024	U-1114			
<u>GABAPENTIN - GRALISE</u>						
N 022544	002 7438927	Feb 26, 2024	U-1114			
<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N 022399	001 6818787	Apr 06, 2025	DS DP			
	8026279	Nov 10, 2026	DS DP			
	8114909	Apr 11, 2026	U-1231			
	8686034	Jan 24, 2025	U-1231			
	8686034	Jan 24, 2025	U-1247			
	8795725	Jun 10, 2029	DP U-1231			
	8795725	Jun 10, 2029	DP U-1247			

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<u>GABAPENTIN ENACARBIL - HORIZANT</u>						
N 022399 002	6818787	Apr 06, 2025	DS DP			
	8026279	Nov 10, 2026	DS DP			
	8114909	Apr 11, 2026		U-1231		
	8686034	Jan 24, 2025		U-1231		
	8686034	Jan 24, 2025		U-1247		
	8795725	Jun 10, 2029	DP	U-1231		
	8795725	Jun 10, 2029	DP	U-1247		
<u>GADOBUTROL - GADOBUTROL</u>						
A 216081 001					CGT	Apr 29, 2024
<u>GADOBUTROL - GADOBUTROL</u>						
A 216081 002					CGT	Apr 29, 2024
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 001	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 002	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 003	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 004	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 005	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 006	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GADOPICLENOL - ELUCIREM</u>						
N 216986 007	10973934	Aug 06, 2039	DS		NCE	Sep 21, 2027
	11590246	Jan 17, 2040	DP			
	8114863	Sep 19, 2028	DS			
<u>GALLIUM DOTATATE GA-68 - NETSPOT</u>						
N 208547 001	9375498	Aug 10, 2032	DP			
<u>GALLIUM GA-68 EDOTREOTIDE - GALLIUM GA 68 EDOTREOTIDE</u>						
N 210828 001					NCE	Aug 21, 2024
					ODE-383	Aug 21, 2026
					W	Aug 21, 2024

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<u>GALLIUM GA-68 EDOTREOTIDE - GALLIUM GA 68 EDOTREOTIDE</u>						
N 210828	001				W	Aug 21, 2026
<u>GALLIUM GA-68 GOZETOTIDE - GALLIUM GA 68 GOZETOTIDE</u>						
N 212642	001				NCE W	Dec 01, 2025 Dec 01, 2025
<u>GALLIUM GA-68 GOZETOTIDE - GALLIUM GA 68 GOZETOTIDE</u>						
N 212643	001				NCE W	Dec 01, 2025 Dec 01, 2025
<u>GALLIUM GA-68 GOZETOTIDE - ILLUCCIX</u>						
N 214032	001	11027031	Jul 28, 2035	U-3317		
<u>GALLIUM GA-68 GOZETOTIDE - LOCAMETZ</u>						
N 215841	001	11369590	Aug 15, 2028	DS DP U-3400	NP	Mar 23, 2025
<u>GANAXOLONE - ZTALMY</u>						
N 215904	001	10603308	Aug 10, 2037	U-3374	NCE	Jun 01, 2027
		7858609	Nov 28, 2026	DP	ODE-395	Jun 01, 2029
		8022054	Nov 28, 2026	DP		
		8318714	Nov 28, 2026	DP		
		8367651	Nov 28, 2026	DP		
		8618087	Nov 28, 2026	U-3374		
		9029355	Nov 28, 2026	DP		
		9056116	Nov 28, 2026	U-3374		
<u>GANCICLOVIR - GANZYK-RTU</u>						
N 209347	001	9486530	Sep 02, 2034	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	001	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	002	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	003	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	004	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	005	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	006	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	007	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	008	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	009	9241948	Jul 01, 2033	DP		
<u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u>						
N 208313	010	9241948	Jul 01, 2033	DP		

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<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164 001	7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028
<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164 002	7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028
<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164 003	7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028
<u>GEPIRONE HYDROCHLORIDE - EXXUA</u>						
N 021164 004	7538116	Sep 02, 2025	U-3699		NCE	Sep 22, 2028
<u>GILTERITINIB FUMARATE - XOSPATA</u>						
N 211349 001	10786500	Jul 01, 2036	DP		ODE-222	Nov 28, 2025
	8969336	Jan 27, 2031	DS DP			
	9487491	Jul 28, 2030	U-2456			
<u>GIVOSIRAN SODIUM - GIVLAARI</u>						
N 212194 001	10119143	Oct 03, 2034	DS DP U-2672		NCE	Nov 20, 2024
	10125364	Mar 15, 2033	DS DP U-2672		ODE-273	Nov 20, 2026
	10131907	Aug 24, 2028	DS DP U-2672			
	10273477	Mar 08, 2024	DS			
	11028392	Oct 03, 2034	DS DP U-2672			
	11530408	May 18, 2024	DS DP			
	8106022	Dec 12, 2029	DS DP U-2672			
	8828956	Dec 04, 2028	DS DP U-2672			
	9133461	Nov 30, 2033	DS DP U-2672			
	9150605	Aug 28, 2025	DS DP			
	9631193	Mar 15, 2033	U-2672			
	9708615	Mar 08, 2024	DS			
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656 001	10414748	Apr 13, 2036	DS DP		ODE-224	Nov 21, 2025
	11168066	Apr 13, 2036	U-3254			
	8148401	Jan 30, 2031	DS DP			
	8431597	Jun 29, 2028	DP			
<u>GLASDEGIB MALEATE - DAURISMO</u>						
N 210656 002	10414748	Apr 13, 2036	DS DP		ODE-224	Nov 21, 2025
	11168066	Apr 13, 2036	U-3254			
	8148401	Jan 30, 2031	DS DP			
	8431597	Jun 29, 2028	DP			
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 209394 001	10028937	Jun 10, 2030	U-2141		ODE-232	Apr 30, 2026
	10028937	Jun 10, 2030	U-3237		ODE-233	Apr 30, 2026
	10028937*PED	Dec 10, 2030			ODE-372	Jun 10, 2028
	10039754	Jun 10, 2030	U-2141		PED	Oct 30, 2026
	10039754	Jun 10, 2030	U-3237		PED	Oct 30, 2026
	10039754*PED	Dec 10, 2030			PED	Dec 10, 2028
	10286029	Mar 14, 2034	U-3237			
	10286029*PED	Sep 14, 2034				
	11246866	Jun 24, 2036	DP			
	11246866*PED	Dec 24, 2036				
	11484534	Mar 14, 2034	U-3237			

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<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 209394	001	11484534*PED	Sep 14, 2034			
		8648037	Jan 19, 2032	DS DP	U-2141	
		8648037	Jan 19, 2032	DS DP	U-3237	
		8648037*PED	Jul 19, 2032			
		8937150	May 18, 2032	DS DP		
		8937150*PED	Nov 18, 2032			
		9321807	Jun 05, 2035	DS		
		9321807*PED	Dec 05, 2035			
		9586978	Nov 06, 2030		U-2141	
		9586978	Nov 06, 2030		U-3237	
		9586978*PED	May 06, 2031			
		RE48923	May 08, 2035	DS		
		RE48923*PED	Nov 08, 2035			
<u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u>						
N 215110	001	10028937	Jun 10, 2030	U-3238	ODE-372	Jun 10, 2028
		10028937*PED	Dec 10, 2030		PED	Dec 10, 2028
		10039754	Jun 10, 2030	U-3238		
		10039754*PED	Dec 10, 2030			
		10286029	Mar 14, 2034	U-3238		
		10286029*PED	Sep 14, 2034			
		8648037	Jan 19, 2032	DS DP	U-3238	
		8648037*PED	Jul 19, 2032			
		8937150	May 18, 2032	DS DP		
		8937150*PED	Nov 18, 2032			
		9321807	Jun 05, 2035	DS DP		
		9321807*PED	Dec 05, 2035			
		9586978	Nov 06, 2030		U-3238	
		9586978*PED	May 06, 2031			
		RE48923	May 08, 2035	DS		
		RE48923*PED	Nov 08, 2035			
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>						
N 021925	001	7700128	Jan 30, 2027	DP		
		8071130	Jun 08, 2028	DP		
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>						
N 021925	002	7700128	Jan 30, 2027	DP		
		8071130	Jun 08, 2028	DP		
<u>GLIPIZIDE - GLIPIZIDE</u>						
A 214874	001				CGT	Apr 15, 2024
<u>GLUCAGON - BAOSIMI</u>						
N 210134	001	10213487	Feb 16, 2036	DP	U-2604	
		10765602	Sep 23, 2039	DP		
		10894133	Jan 03, 2038	DP		
<u>GLUCAGON - GVOKE PFS</u>						
N 212097	001	11590205	Apr 22, 2036	DP	U-2742	
		9649364	Apr 22, 2036	DP	U-2742	

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<u>GLUCAGON - GVOKE PFS</u>						
N 212097	002	11590205	Apr 22, 2036	DP U-2742		
	9649364	Apr 22, 2036	DP U-2742			
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	003	11590205	Apr 22, 2036	DP U-2742		
	9649364	Apr 22, 2036	DP U-2742			
<u>GLUCAGON - GVOKE HYPOPEN</u>						
N 212097	004	11590205	Apr 22, 2036	DP U-2742		
	9649364	Apr 22, 2036	DP U-2742			
<u>GLUCAGON - GVOKE KIT</u>						
N 212097	005	11590205	Apr 22, 2036	DP U-2742		
<u>GLYCEROL PHENYLBUTYRATE - RAVICTI</u>						
N 203284	001	10045958	Sep 22, 2030	U-1816	ODE-157	Apr 28, 2024
		10045959	Sep 22, 2030	U-1816		
		10183002	Sep 22, 2030	U-1816		
		10183003	Sep 22, 2030	U-1816		
		10183004	Sep 22, 2030	U-1816		
		10183005	Sep 22, 2030	U-1816		
		10183006	Sep 22, 2030	U-1816		
		10668040	Sep 22, 2030	U-1816		
		8404215	Mar 09, 2032	U-1383		
		8642012	Sep 22, 2030	U-1383		
		9095559	Mar 09, 2032	U-1383		
		9254278	Mar 09, 2032	U-1816		
		9326966	Mar 09, 2032	U-1816		
		9561197	Sep 22, 2030	U-1383		
		9962359	Sep 22, 2030	U-1816		
		9999608	Sep 22, 2030	U-1816		
<u>GLYCOPYRROLATE - SEEBRI</u>						
N 207923	001	8182838	Oct 20, 2028	DP		
		8479730	Oct 11, 2028	DP		
<u>GLYCOPYRROLATE - LONHALA MAGNAIR KIT</u>						
N 208437	001	10376661	Sep 14, 2035	DP		
		10688518	Nov 12, 2036	DP		
		10744277	Dec 07, 2036	DP U-2941		
		10940110	Feb 26, 2029	DP U-1773		
		11278683	Aug 16, 2026	DP		
		7458372	Nov 18, 2024	DP		
		7931212	Nov 25, 2025	DP		
		9168556	Sep 01, 2032	DP		
		9265900	Dec 07, 2028	DP		
		9604018	May 16, 2033	DP		
		9789270	Oct 30, 2030	DP		
<u>GLYCOPYRROLATE; INDACATEROL MALEATE - UTIBRON</u>						
N 207930	001	6878721	Feb 25, 2025	DS DP U-1773		
		8182838	Oct 20, 2028	DP		
		8479730	Oct 11, 2028	DP		

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<u>GLYCOPYRROLATE; NEOSTIGMINE METHYLSULFATE - PREVDUO</u>						
N 216903	001 10456354	Oct 25, 2038	DP			
	11110054	Oct 25, 2038	DP			
<u>GLYCOPYRRONIUM TOSYLATE - OBREXZA</u>						
N 210361	001 10004717	Feb 28, 2033	DP U-2398			
	10052267	Oct 17, 2028	DP U-2398			
	10543192	Feb 28, 2033	DP			
	10548875	Feb 28, 2033	DS DP U-2398			
	8618160	Dec 10, 2029	DP U-2398			
	8859610	Feb 28, 2033	DP U-2398			
	9259414	Feb 28, 2033	U-2398			
	9744105	Jul 18, 2030	DP U-2398			
<u>GOLODIRSEN - VYONDYS 53</u>						
N 211970	001 10227590	Jun 28, 2025	DS DP		NCE	Dec 12, 2024
	10266827	Jun 28, 2025	U-2675		ODE-280	Dec 12, 2026
	10421966	Jun 28, 2025	DS DP			
	10968450	Jun 28, 2025	DS DP			
	10995337	Jun 28, 2025	DP U-2675			
	9024007	Jun 28, 2025	DS DP			
	9994851	Jun 28, 2025	DS DP			
	RE47691	Jun 28, 2025	DP			
<u>GRANISETRON - SANCUSO</u>						
N 022198	001 7608282	Jan 22, 2025	DP U-1011			
<u>GRANISETRON - SUSTOL</u>						
N 022445	001 10357570	Sep 28, 2024	U-2253			
	8252304	Sep 28, 2024	DP			
	8252305	Sep 28, 2024	U-1891			
	8715710	Sep 28, 2024	DP			
	9913910	Sep 28, 2024	U-2253			
<u>GUAIFENESIN; HYDROCODONE BITARTRATE - OBREDON</u>						
N 205474	001 10105324	Nov 13, 2035	DS DP U-2023			
	9549907	Nov 13, 2035	DS DP U-2023			
	9808431	Nov 13, 2035	DS DP U-2023			
<u>HALOBETASOL PROPIONATE - ULTRAVATE</u>						
N 208183	001 8962028	Jun 19, 2033	DP U-1775			
<u>HALOBETASOL PROPIONATE - BRYHALI</u>						
N 209355	001 10478502	Nov 02, 2031	DP U-2625			
	11839656	Nov 02, 2031	U-2625			
	8809307	Nov 02, 2031	DP			
<u>HALOBETASOL PROPIONATE - LEXETTE</u>						
N 210566	001 10857159	Nov 30, 2036	DP		NPP	Aug 18, 2024
	10857159*PED	May 30, 2037				
	11020407	Nov 30, 2036	DP U-3143			
<u>HALOBETASOL PROPIONATE; TAZAROTENE - DUOBRII</u>						
N 209354	001 10251895	Jun 06, 2036	DP			
	10426787	Jun 06, 2036	U-2625			
	10478502	Nov 02, 2031	DP U-2625			

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<u>HALOBETASOL PROPIONATE; TAZAROTENE - DUOBRII</u>						
N 209354	001	11648256	Jun 06, 2036	DP	U-2625	
		11679115	Jun 06, 2036	DP	U-2625	
		11839656	Nov 02, 2031		U-2625	
		8809307	Nov 02, 2031	DP		
<u>HEPARIN SODIUM; TAUROLIDINE - DEFENCATH</u>						
N 214520	001	11738120	Apr 15, 2042	DS DP		
		7696182	May 16, 2025	DS DP	U-3774	
		8541393	Nov 02, 2024	DS DP	U-3774	
		9339036	Nov 02, 2024	DS DP		
<u>HEPARIN SODIUM; TAUROLIDINE - DEFENCATH</u>						
N 214520	002	11738120	Apr 15, 2042	DS DP		
		7696182	May 16, 2025	DS DP	U-3774	
		8541393	Nov 02, 2024	DS DP	U-3774	
		9339036	Nov 02, 2024	DS DP		
<u>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u>						
N 022555	001	10556010	Dec 19, 2036		U-2250	
		11235168	Jan 04, 2038		U-2250	
		11311620	Dec 19, 2036		U-2250	
<u>HISTRELIN ACETATE - SUPPRELIN LA</u>						
N 022058	001	8062652	Jun 16, 2026		U-1197	
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	001	10028946	Jul 25, 2033		U-1810	
		10092559	Sep 12, 2034		U-55	
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033		U-1810	
		10722511	Jul 25, 2033		U-1810	
		9132096	Sep 12, 2034	DP		
		9265760	Jul 25, 2033		U-1810	
		9326982	Jul 25, 2033		U-1810	
		9333201	Jul 25, 2033		U-1810	
		9339499	Jul 25, 2033		U-1810	
		9421200	Jul 25, 2033		U-1810	
		9433619	Jul 25, 2033		U-1810	
		9452163	Sep 12, 2034		U-55	
		9486451	Sep 12, 2034		U-55	
		9610286	Jul 25, 2033		U-1810	
		9713611	Sep 12, 2034	DP	U-55	
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	002	10028946	Jul 25, 2033		U-1810	
		10092559	Sep 12, 2034		U-55	
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033		U-1810	
		10722511	Jul 25, 2033		U-1810	
		9132096	Sep 12, 2034	DP		
		9265760	Jul 25, 2033		U-1810	
		9326982	Jul 25, 2033		U-1810	
		9333201	Jul 25, 2033		U-1810	

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	002	9339499	Jul 25, 2033	U-1810		
		9421200	Jul 25, 2033	U-1810		
		9433619	Jul 25, 2033	U-1810		
		9452163	Sep 12, 2034	U-55		
		9486451	Sep 12, 2034	U-55		
		9610286	Jul 25, 2033	U-1810		
		9713611	Sep 12, 2034	DP U-55		
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	003	10028946	Jul 25, 2033	U-1810		
		10092559	Sep 12, 2034	U-55		
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033	U-1810		
		10722511	Jul 25, 2033	U-1810		
		9132096	Sep 12, 2034	DP		
		9265760	Jul 25, 2033	U-1810		
		9326982	Jul 25, 2033	U-1810		
		9333201	Jul 25, 2033	U-1810		
		9339499	Jul 25, 2033	U-1810		
		9421200	Jul 25, 2033	U-1810		
		9433619	Jul 25, 2033	U-1810		
		9452163	Sep 12, 2034	U-55		
		9486451	Sep 12, 2034	U-55		
		9610286	Jul 25, 2033	U-1810		
		9713611	Sep 12, 2034	DP U-55		
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	004	10028946	Jul 25, 2033	U-1810		
		10092559	Sep 12, 2034	U-55		
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033	U-1810		
		10722511	Jul 25, 2033	U-1810		
		9132096	Sep 12, 2034	DP		
		9265760	Jul 25, 2033	U-1810		
		9326982	Jul 25, 2033	U-1810		
		9333201	Jul 25, 2033	U-1810		
		9339499	Jul 25, 2033	U-1810		
		9421200	Jul 25, 2033	U-1810		
		9433619	Jul 25, 2033	U-1810		
		9452163	Sep 12, 2034	U-55		
		9486451	Sep 12, 2034	U-55		
		9610286	Jul 25, 2033	U-1810		
		9713611	Sep 12, 2034	DP U-55		
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880	005	10028946	Jul 25, 2033	U-1810		
		10092559	Sep 12, 2034	U-55		
		10322120	Jul 25, 2033	DP		
		10456393	Jul 25, 2033	U-1810		
		10722511	Jul 25, 2033	U-1810		
		9132096	Sep 12, 2034	DP		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880 005	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u>						
N 202880 006	10028946	Jul 25, 2033	U-1810			
	10092559	Sep 12, 2034	U-55			
	10322120	Jul 25, 2033	DP			
	10456393	Jul 25, 2033	U-1810			
	10722511	Jul 25, 2033	U-1810			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627 001	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027	U-1556			
	11844865	Feb 13, 2025	DP U-1556			
	8808740	Dec 21, 2031	DP U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027	U-1556			
	9492391	Aug 24, 2027	U-1556			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
	9872837	Dec 21, 2031	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627 002	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11844865	Feb 13, 2025	DP	U-1556		
	8808740	Dec 21, 2031	DP	U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027		U-1556		
	9492391	Aug 24, 2027		U-1556		
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
	9872837	Dec 21, 2031	DP			
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627 003	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11844865	Feb 13, 2025	DP	U-1556		
	8808740	Dec 21, 2031	DP	U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027		U-1556		
	9492391	Aug 24, 2027		U-1556		
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
	9872837	Dec 21, 2031	DP			
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627 004	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	11844865	Feb 13, 2025	DP	U-1556		
	8808740	Dec 21, 2031	DP	U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	004	9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027		U-1556	
		9492391	Aug 24, 2027		U-1556	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	005	11304908	Aug 24, 2027	DP		
		11304909	Aug 24, 2027		U-1556	
		11844865	Feb 13, 2025	DP	U-1556	
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	DP		
		9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027		U-1556	
		9492391	Aug 24, 2027		U-1556	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	006	11304908	Aug 24, 2027	DP		
		11304909	Aug 24, 2027		U-1556	
		11844865	Feb 13, 2025	DP	U-1556	
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	DP		
		9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027		U-1556	
		9492391	Aug 24, 2027		U-1556	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		

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<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	006	9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - HYSINGLA ER</u>						
N 206627	007	11304908	Aug 24, 2027	DP		
		11304909	Aug 24, 2027		U-1556	
		11844865	Feb 13, 2025	DP	U-1556	
		8808740	Dec 21, 2031	DP	U-1556	
		9084816	Aug 24, 2027	DP		
		9095614	Aug 24, 2027		U-1556	
		9095615	Aug 24, 2027	DP		
		9486412	Aug 24, 2027	DP		
		9486413	Aug 24, 2027	DP		
		9492389	Aug 24, 2027	DP		
		9492390	Aug 24, 2027		U-1556	
		9492391	Aug 24, 2027		U-1556	
		9545380	Aug 24, 2027		U-1556	
		9572779	Dec 21, 2031	DP		
		9750703	Dec 21, 2031	DP		
		9763933	Aug 24, 2027	DP		
		9770416	Aug 24, 2027	DP		
		9775809	Aug 24, 2027	DP		
		9861584	Dec 21, 2031	DP		
		9872837	Dec 21, 2031	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	001	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	002	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	003	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	004	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		
<u>HYDROCODONE BITARTRATE - VANTRELA ER</u>						
N 207975	005	8445018	Jul 31, 2029	DP		
		9216176	Sep 13, 2027	DP		
		9572803	Sep 13, 2027	DP		

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<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	001 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	002 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	003 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE - ALKINDI SPRINKLE</u>						
N 213876	004 9649280	May 12, 2034	DP U-3075			
	9675559	Jan 10, 2033	U-3075			
	9717740	Nov 19, 2032	U-3075			
<u>HYDROCORTISONE BUTYRATE - LOCOID</u>						
N 022076	001 7378405	Dec 19, 2026	DP			
	7981877	Jan 23, 2025	DP			
<u>HYDROGEN PEROXIDE - ESKATA</u>						
N 209305	001 10098910	Apr 21, 2035	DP U-2205			
	10493103	Apr 21, 2035	DP			
	10729720	Apr 21, 2035	DP			
	9675639	Jul 04, 2035	DP U-2205			
	9980983	Apr 21, 2035	U-2205			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N 019034	001 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N 019034	002 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	003 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	004 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	005 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	006 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034	007 9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			

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<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N 019034 007	9248229	Mar 12, 2034	DP			
	9731082	Apr 23, 2032	DP			
<u>HYDROXYUREA - SIKLOS</u>						
N 208843 001					ODE-177	Dec 21, 2024
<u>HYDROXYUREA - SIKLOS</u>						
N 208843 002					ODE-177	Dec 21, 2024
<u>IBREXAFUNGERP CITRATE - BREXAFEMME</u>						
N 214900 001	10174074	Jan 19, 2035	DS DP		I-903	Nov 30, 2025
	10370406	Jan 19, 2035		U-3159	NCE	Jun 01, 2026
	10927142	Jan 19, 2035	DS		GAIN	Jun 01, 2031
	11534433	Jun 10, 2039		U-3159		
	11534433	Jun 10, 2039		U-3508		
	8188085	Aug 28, 2030	DS DP	U-3159		
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 001	10004746	Jun 03, 2031		U-1684	NPP	Aug 24, 2025
	10004746	Jun 03, 2031		U-1946	ODE-152	Aug 02, 2024
	10004746	Jun 03, 2031		U-2241	ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031		U-2242	PED	Feb 02, 2025
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2026
	10016435	Jun 03, 2031		U-1650	PED	Feb 24, 2030
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10294231	Jun 03, 2033		DP		
	10294231*PED	Dec 03, 2033				
	10294232	Jun 03, 2033		DP		
	10294232*PED	Dec 03, 2033				
	10463668	Oct 24, 2034		U-2654		
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031		U-1650		
	10478439	Jun 03, 2031		U-1684		
	10478439	Jun 03, 2031		U-1946		
	10478439	Jun 03, 2031		U-2241		
	10478439	Jun 03, 2031		U-2242		
	10478439	Jun 03, 2031		U-2665		
	10478439	Jun 03, 2031		U-3422		
	10478439*PED	Dec 03, 2031				
	10653696	Jun 03, 2031		U-1456		
	10653696*PED	Dec 03, 2031				
	10695350	Oct 24, 2034		U-2846		
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031		U-1491		
	10751342	Jun 03, 2031		U-1946		
	10751342	Jun 03, 2031		U-2943		
	10751342	Jun 03, 2031		U-2944		
	10751342*PED	Dec 03, 2031				

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<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 001	10752634	Jun 03, 2033	DP			
	10752634*PED	Dec 03, 2033				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031		U-1684		
	11672803	Jun 03, 2031		U-1946		
	11672803	Jun 03, 2031		U-2241		
	11672803	Jun 03, 2031		U-2242		
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026		U-1456		
	8476284	Dec 28, 2026		U-1650		
	8476284	Dec 28, 2026		U-1946		
	8476284	Dec 28, 2026		U-1947		
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026		U-1456		
	8497277	Dec 28, 2026		U-1491		
	8497277	Dec 28, 2026		U-1650		
	8497277	Dec 28, 2026		U-1946		
	8497277	Dec 28, 2026		U-1947		
	8497277	Dec 28, 2026		U-2241		
	8497277	Dec 28, 2026		U-2242		
	8497277	Dec 28, 2026		U-3422		
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027		U-1491		
	8563563	Apr 26, 2027		U-1650		
	8563563	Apr 26, 2027		U-1946		
	8563563	Apr 26, 2027		U-2219		
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026		U-1491		
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031		U-1456		
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026		U-1456		
	8952015	Dec 28, 2026		U-1491		
	8952015	Dec 28, 2026		U-1650		
	8952015	Dec 28, 2026		U-1946		
	8952015	Dec 28, 2026		U-1947		
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				

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<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 001	8999999	Jun 03, 2031	U-1683			
	8999999	Jun 03, 2031	U-1684			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1745			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS DP			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS DP			
	9296753*PED	Apr 30, 2034				
	9540382	Aug 18, 2033	U-1456			
	9540382	Aug 18, 2033	U-1650			
	9540382	Aug 18, 2033	U-1684			
	9540382	Aug 18, 2033	U-1946			
	9540382	Aug 18, 2033	U-1947			
	9540382*PED	Feb 18, 2034				
	9713617	Jun 03, 2033	DP			
	9713617*PED	Dec 03, 2033				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2150			
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883*PED	Dec 03, 2031				
	9814721	Jun 03, 2031	U-1947			
	9814721*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	10004746	Jun 03, 2031	U-1684		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		ODE*	Aug 02, 2024
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10016435	Jun 03, 2031	U-1650			
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10294231	Jun 03, 2033	DP			
	10294231*PED	Dec 03, 2033				
	10294232	Jun 03, 2033	DP			
	10294232*PED	Dec 03, 2033				
	10463668	Oct 24, 2034	U-2654			
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031	U-1650			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	10478439	Jun 03, 2031				U-1684
	10478439	Jun 03, 2031				U-1946
	10478439	Jun 03, 2031				U-2241
	10478439	Jun 03, 2031				U-2242
	10478439	Jun 03, 2031				U-2665
	10478439	Jun 03, 2031				U-3422
	10478439*PED	Dec 03, 2031				
	10653696	Jun 03, 2031				U-1456
	10653696*PED	Dec 03, 2031				
	10695350	Oct 24, 2034				U-2846
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031				U-1491
	10751342	Jun 03, 2031				U-1946
	10751342	Jun 03, 2031				U-2943
	10751342	Jun 03, 2031				U-2944
	10751342*PED	Dec 03, 2031				
	10961251	Jun 03, 2033		DP		
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031				U-1684
	11672803	Jun 03, 2031				U-1946
	11672803	Jun 03, 2031				U-2241
	11672803	Jun 03, 2031				U-2242
	7514444	Dec 28, 2026		DS DP		
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027		DS DP		
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026				U-1456
	8476284	Dec 28, 2026				U-1650
	8476284	Dec 28, 2026				U-1946
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026				U-1456
	8497277	Dec 28, 2026				U-1491
	8497277	Dec 28, 2026				U-1650
	8497277	Dec 28, 2026				U-1946
	8497277	Dec 28, 2026				U-2241
	8497277	Dec 28, 2026				U-2242
	8497277	Dec 28, 2026				U-3422
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027				U-1491
	8563563	Apr 26, 2027				U-1650
	8563563	Apr 26, 2027				U-1946
	8563563	Apr 26, 2027				U-2219
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026		DS DP		
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026				U-1491
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026		DS DP		
	8735403*PED	Jun 28, 2027				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 205552 002	8754090	Jun 03, 2031	U-1456			
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1456			
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2228			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9540382	Aug 18, 2033	U-1456			
	9540382	Aug 18, 2033	U-1491			
	9540382	Aug 18, 2033	U-1650			
	9540382	Aug 18, 2033	U-1946			
	9540382*PED	Feb 18, 2034				
	9713617	Jun 03, 2033	DP			
	9713617*PED	Dec 03, 2033				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883*PED	Dec 03, 2031				
	9814721	Jun 03, 2031	U-1947			
	9814721*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	10004746	Jun 03, 2031	U-1684		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		ODE*	Aug 02, 2024
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				
	10016435	Jun 03, 2031	U-1650			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034	U-2654			
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10478439	Jun 03, 2031	U-3422			
	10478439*PED	Dec 03, 2031				
	10653696	Jun 03, 2031	U-1456			
	10653696*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2846			
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10752634	Jun 03, 2033	DP			
	10752634*PED	Dec 03, 2033				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-1684			
	11672803	Jun 03, 2031	U-1946			
	11672803	Jun 03, 2031	U-2241			
	11672803	Jun 03, 2031	U-2242			
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026	U-1456			
	8476284	Dec 28, 2026	U-1650			
	8476284	Dec 28, 2026	U-1946			
	8476284	Dec 28, 2026	U-2241			
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026	U-1456			
	8497277	Dec 28, 2026	U-1491			
	8497277	Dec 28, 2026	U-1650			
	8497277	Dec 28, 2026	U-1946			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	8497277	Dec 28, 2026	U-2241			
	8497277	Dec 28, 2026	U-2242			
	8497277	Dec 28, 2026	U-3422			
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-1491			
	8563563	Apr 26, 2027	U-1650			
	8563563	Apr 26, 2027	U-1946			
	8563563	Apr 26, 2027	U-2241			
	8563563	Apr 26, 2027	U-2242			
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-1491			
	8703780	Dec 28, 2026	U-2242			
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031	U-1456			
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1456			
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015	Dec 28, 2026	U-2241			
	8952015	Dec 28, 2026	U-2242			
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 001	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9801883*PED	Dec 03, 2031				
	9814721	Jun 03, 2031	U-1947			
	9814721*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	10004746	Jun 03, 2031	U-1684		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		ODE*	Aug 02, 2024
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				
	10016435	Jun 03, 2031	U-1650			
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034	U-2654			
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10478439	Jun 03, 2031	U-3422			
	10478439*PED	Dec 03, 2031				
	10653696	Jun 03, 2031	U-1456			
	10653696*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2846			
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-1684			
	11672803	Jun 03, 2031	U-1946			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	11672803	Jun 03, 2031				U-2241
	11672803	Jun 03, 2031				U-2242
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026				U-1456
	8476284	Dec 28, 2026				U-1650
	8476284	Dec 28, 2026				U-1946
	8476284	Dec 28, 2026				U-2241
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026				U-1456
	8497277	Dec 28, 2026				U-1491
	8497277	Dec 28, 2026				U-1650
	8497277	Dec 28, 2026				U-1946
	8497277	Dec 28, 2026				U-2241
	8497277	Dec 28, 2026				U-2242
	8497277	Dec 28, 2026				U-3422
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027				U-1491
	8563563	Apr 26, 2027				U-1650
	8563563	Apr 26, 2027				U-1946
	8563563	Apr 26, 2027				U-2241
	8563563	Apr 26, 2027				U-2242
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026				U-1491
	8703780	Dec 28, 2026				U-2242
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031				U-1456
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026				U-1456
	8952015	Dec 28, 2026				U-1491
	8952015	Dec 28, 2026				U-1650
	8952015	Dec 28, 2026				U-1946
	8952015	Dec 28, 2026				U-2241
	8952015	Dec 28, 2026				U-2242
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031				U-1491
	8999999	Jun 03, 2031				U-1946
	8999999	Jun 03, 2031				U-2241
	8999999	Jun 03, 2031				U-2242

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 002	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9801883*PED	Dec 03, 2031				
	9814721	Jun 03, 2031	U-1947			
	9814721*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 003	10004746	Jun 03, 2031	U-1684		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		ODE*	Aug 02, 2024
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				
	10016435	Jun 03, 2031	U-1650			
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			
	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034	U-2654			
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031	U-1650			
	10478439	Jun 03, 2031	U-1684			
	10478439	Jun 03, 2031	U-1946			
	10478439	Jun 03, 2031	U-2241			
	10478439	Jun 03, 2031	U-2242			
	10478439	Jun 03, 2031	U-2665			
	10478439	Jun 03, 2031	U-3422			
	10478439*PED	Dec 03, 2031				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 003	10653696	Jun 03, 2031	U-1456			
	10653696*PED	Dec 03, 2031				
	10695350	Oct 24, 2034	U-2846			
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031	U-1491			
	10751342	Jun 03, 2031	U-1946			
	10751342	Jun 03, 2031	U-2943			
	10751342	Jun 03, 2031	U-2944			
	10751342*PED	Dec 03, 2031				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	11672803	Jun 03, 2031	U-1684			
	11672803	Jun 03, 2031	U-1946			
	11672803	Jun 03, 2031	U-2241			
	11672803	Jun 03, 2031	U-2242			
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026	U-1456			
	8476284	Dec 28, 2026	U-1650			
	8476284	Dec 28, 2026	U-1946			
	8476284	Dec 28, 2026	U-2241			
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026	U-1456			
	8497277	Dec 28, 2026	U-1491			
	8497277	Dec 28, 2026	U-1650			
	8497277	Dec 28, 2026	U-1946			
	8497277	Dec 28, 2026	U-2241			
	8497277	Dec 28, 2026	U-2242			
	8497277	Dec 28, 2026	U-3422			
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027	U-1491			
	8563563	Apr 26, 2027	U-1650			
	8563563	Apr 26, 2027	U-1946			
	8563563	Apr 26, 2027	U-2241			
	8563563	Apr 26, 2027	U-2242			
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026	U-1491			
	8703780	Dec 28, 2026	U-2242			
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031	U-1456			
	8754090*PED	Dec 03, 2031				

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 003	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026	U-1456			
	8952015	Dec 28, 2026	U-1491			
	8952015	Dec 28, 2026	U-1650			
	8952015	Dec 28, 2026	U-1946			
	8952015	Dec 28, 2026	U-2241			
	8952015	Dec 28, 2026	U-2242			
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031	U-1491			
	8999999	Jun 03, 2031	U-1946			
	8999999	Jun 03, 2031	U-2241			
	8999999	Jun 03, 2031	U-2242			
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031	U-1650			
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034	U-2969			
	9795604	Oct 24, 2034	U-2970			
	9795604	Oct 24, 2034	U-3422			
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031	U-1491			
	9801881	Jun 03, 2031	U-2242			
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031	U-2159			
	9801883	Jun 03, 2031	U-2243			
	9801883*PED	Dec 03, 2031				
	9814721	Jun 03, 2031	U-1947			
	9814721*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	10004746	Jun 03, 2031	U-1684		NPP	Aug 24, 2025
	10004746	Jun 03, 2031	U-1946		ODE*	Aug 02, 2024
	10004746	Jun 03, 2031	U-2241		ODE-405	Aug 24, 2029
	10004746	Jun 03, 2031	U-2242		PED	Feb 24, 2026
	10004746*PED	Dec 03, 2031			PED	Feb 24, 2030
	10010507	Mar 03, 2036	DP			
	10010507*PED	Sep 03, 2036				
	10016435	Jun 03, 2031	U-1650			
	10016435*PED	Dec 03, 2031				
	10106548	Jun 03, 2033	DS DP			

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	10106548*PED	Dec 03, 2033				
	10125140	Jun 03, 2033	DS DP			
	10125140*PED	Dec 03, 2033				
	10213386	Mar 03, 2036	DP			
	10213386*PED	Sep 03, 2036				
	10463668	Oct 24, 2034		U-2654		
	10463668*PED	Apr 24, 2035				
	10478439	Jun 03, 2031		U-1456		
	10478439	Jun 03, 2031		U-1650		
	10478439	Jun 03, 2031		U-1684		
	10478439	Jun 03, 2031		U-1946		
	10478439	Jun 03, 2031		U-1947		
	10478439	Jun 03, 2031		U-2241		
	10478439	Jun 03, 2031		U-2242		
	10478439	Jun 03, 2031		U-2665		
	10478439*PED	Dec 03, 2031				
	10653696	Jun 03, 2031		U-1456		
	10653696*PED	Dec 03, 2031				
	10695350	Oct 24, 2034		U-2846		
	10695350*PED	Apr 24, 2035				
	10751342	Jun 03, 2031		U-1491		
	10751342	Jun 03, 2031		U-1946		
	10751342	Jun 03, 2031		U-2943		
	10751342	Jun 03, 2031		U-2944		
	10751342*PED	Dec 03, 2031				
	10828259	Mar 03, 2036	DP			
	10828259*PED	Sep 03, 2036				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8476284	Dec 28, 2026		U-1456		
	8476284	Dec 28, 2026		U-1650		
	8476284	Dec 28, 2026		U-1946		
	8476284	Dec 28, 2026		U-1947		
	8476284	Dec 28, 2026		U-2241		
	8476284*PED	Jun 28, 2027				
	8497277	Dec 28, 2026		U-1456		
	8497277	Dec 28, 2026		U-1491		
	8497277	Dec 28, 2026		U-1650		
	8497277	Dec 28, 2026		U-1946		
	8497277	Dec 28, 2026		U-1947		
	8497277	Dec 28, 2026		U-2241		
	8497277	Dec 28, 2026		U-2242		
	8497277*PED	Jun 28, 2027				
	8563563	Apr 26, 2027		U-1491		
	8563563	Apr 26, 2027		U-1650		

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	8563563	Apr 26, 2027				U-1946
	8563563	Apr 26, 2027				U-2241
	8563563	Apr 26, 2027				U-2242
	8563563*PED	Oct 26, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8703780	Dec 28, 2026				U-1491
	8703780	Dec 28, 2026				U-2242
	8703780*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754090	Jun 03, 2031				U-1456
	8754090*PED	Dec 03, 2031				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8952015	Dec 28, 2026				U-1456
	8952015	Dec 28, 2026				U-1491
	8952015	Dec 28, 2026				U-1650
	8952015	Dec 28, 2026				U-1946
	8952015	Dec 28, 2026				U-1947
	8952015	Dec 28, 2026				U-2241
	8952015	Dec 28, 2026				U-2242
	8952015*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	8999999	Jun 03, 2031				U-1491
	8999999	Jun 03, 2031				U-1946
	8999999	Jun 03, 2031				U-2241
	8999999	Jun 03, 2031				U-2242
	8999999*PED	Dec 03, 2031				
	9125889	Jun 03, 2031				U-1650
	9125889*PED	Dec 03, 2031				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9655857	Mar 03, 2036	DP			
	9655857*PED	Sep 03, 2036				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034				U-2969
	9795604	Oct 24, 2034				U-2970
	9795604*PED	Apr 24, 2035				
	9801881	Jun 03, 2031				U-1491
	9801881	Jun 03, 2031				U-2242
	9801881*PED	Dec 03, 2031				
	9801883	Jun 03, 2031				U-2159
	9801883	Jun 03, 2031				U-2243
	9801883*PED	Dec 03, 2031				

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<u>IBRUTINIB - IMBRUVICA</u>						
N 210563 004	9814721	Jun 03, 2031		U-1947		
	9814721*PED	Dec 03, 2031				
<u>IBRUTINIB - IMBRUVICA</u>						
N 217003 001	10106548	Jun 03, 2033	DS DP		NP	Aug 24, 2025
	10106548*PED	Dec 03, 2033			ODE-405	Aug 24, 2029
	10125140	Jun 03, 2033	DS DP		PED	Feb 24, 2026
	10125140*PED	Dec 03, 2033			PED	Feb 24, 2030
	10478439	Jun 03, 2031		U-3422		
	10478439*PED	Dec 03, 2031				
	10961251	Jun 03, 2033	DP			
	10961251*PED	Dec 03, 2033				
	7514444	Dec 28, 2026	DS DP			
	7514444*PED	Jun 28, 2027				
	8008309	Nov 13, 2027	DS DP			
	8008309*PED	May 13, 2028				
	8497277	Dec 28, 2026		U-3422		
	8497277*PED	Jun 28, 2027				
	8697711	Dec 28, 2026	DS DP			
	8697711*PED	Jun 28, 2027				
	8735403	Dec 28, 2026	DS DP			
	8735403*PED	Jun 28, 2027				
	8754091	Dec 28, 2026	DP			
	8754091*PED	Jun 28, 2027				
	8957079	Dec 28, 2026	DS DP			
	8957079*PED	Jun 28, 2027				
	9181257	Dec 28, 2026	DS			
	9181257*PED	Jun 28, 2027				
	9296753	Oct 30, 2033	DS			
	9296753*PED	Apr 30, 2034				
	9725455	Jun 03, 2033	DS			
	9725455*PED	Dec 03, 2033				
	9795604	Oct 24, 2034		U-3422		
	9795604*PED	Apr 24, 2035				
<u>IBUPROFEN - CHILDREN'S ADVIL-FLAVORED</u>						
N 020589 002	10238640	May 25, 2024	DP			
<u>IBUPROFEN - CALDOLOR</u>						
N 022348 001					M-128	Nov 19, 2024
<u>IBUPROFEN - CALDOLOR</u>						
N 022348 002	11806400	Mar 16, 2032		U-3746	M-128	Nov 19, 2024
	8735452	Sep 30, 2029		U-981		
	8871810	Sep 30, 2029		U-981		
	9012508	Sep 14, 2030		U-981		
	9114068	Sep 30, 2029		U-1735		
	9138404	Sep 30, 2029		U-1756		
	9295639	Sep 30, 2029		U-1756		
	9649284	Sep 30, 2029		U-2018		

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<u>IBUPROFEN - CALDOLOR</u>						
N 022348 003	11806400	Mar 16, 2032	U-3746		M-128	Nov 19, 2024
	8735452	Sep 30, 2029	U-981			
	8871810	Sep 30, 2029	U-981			
	9012508	Sep 14, 2030	U-981			
	9072661	Mar 16, 2032	U-2264			
	9072710	Mar 16, 2032	U-2266			
<u>IBUPROFEN LYSINE - NEOPROFEN</u>						
N 021903 001	8415337	Mar 02, 2032	DS DP			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 001	10010517	Apr 29, 2030	U-2690			
	10265287	Apr 29, 2030	U-2700			
	10278935	Jun 28, 2033	U-2701			
	10278936	Jun 28, 2033	U-2702			
	10278937	Jun 28, 2033	U-2703			
	10383840	Jun 28, 2033	U-2704			
	10555924	Jun 28, 2033	U-2743			
	10555925	Jun 28, 2033	U-2744			
	10568861	Jun 28, 2033	U-2756			
	10576054	Jun 28, 2033	U-2762			
	10668042	Jun 28, 2033	U-2841			
	10786478	Jun 28, 2033	U-2959			
	10786478	Jun 28, 2033	U-2960			
	10792267	Apr 29, 2030	U-2961			
	10792270	Jun 28, 2033	U-2962			
	10842766	Apr 29, 2030	U-2997			
	10842768	Jun 15, 2030	U-2688			
	10881632	Apr 29, 2030	U-3052			
	10894028	Jun 28, 2033	U-3053			
	11000499	Jun 28, 2033	U-3126			
	11103477	Apr 29, 2030	U-3209			
	11116742	Jun 28, 2033	U-3221			
	11154526	Apr 29, 2030	U-3240			
	11213504	Apr 29, 2030	U-3292			
	11298333	Jun 28, 2033	U-3358			
	11369582	Jun 28, 2033	U-2841			
	11717504	Apr 29, 2030	U-3669			
	8293727	Feb 09, 2030	U-1287			
	8293728	Feb 09, 2030	U-1287			
	8298554	Apr 29, 2030	DP			
	8314086	Feb 09, 2030	U-1287			
	8318715	Feb 09, 2030	U-1287			
	8357677	Feb 09, 2030	U-1287			
	8367652	Feb 09, 2030	U-1287			
	8377920	Feb 09, 2030	U-1287			
	8399446	Feb 09, 2030	U-1287			
	8410086	Jun 15, 2030	U-2688			
	8415335	Feb 09, 2030	U-1287			
	8426399	Feb 09, 2030	U-1287			
	8431560	Feb 09, 2030	U-1287			

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<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 001	8440650	Feb 09, 2030	U-1287			
	8445003	Apr 29, 2030	U-1287			
	8445013	Apr 29, 2030	U-1287			
	8454994	Apr 29, 2030	U-2689			
	8455472	Jun 15, 2030	U-2690			
	8501225	Apr 29, 2030	U-1287			
	8518929	Feb 09, 2030	U-1287			
	8524698	Feb 09, 2030	U-1287			
	8546372	Feb 09, 2030	U-1287			
	8551521	Apr 29, 2030	U-1287			
	8563608	Apr 29, 2030	U-1287			
	8617593	Apr 29, 2030	U-1478			
	8617593	Apr 29, 2030	U-2691			
	8617594	Apr 29, 2030	U-1287			
	8618166	Apr 29, 2030	U-2689			
	8623406	Apr 29, 2030	U-1478			
	8623406	Apr 29, 2030	U-2692			
	8642077	Apr 29, 2030	U-2693			
	8669245	Jun 15, 2030	U-2694			
	8680144	Feb 09, 2030	U-2695			
	8691871	Apr 29, 2030	U-2689			
	8703185	Apr 29, 2030	U-2691			
	8709475	Apr 29, 2030	U-2689			
	8710041	Jun 15, 2030	U-2690			
	9198892	Sep 25, 2027	U-2706			
	9603826	Jun 28, 2033	U-2696			
	9610272	Jun 28, 2033	U-2697			
	9623001	Jun 28, 2033	U-2698			
	9693984	Jun 28, 2033	U-2697			
	9693985	Jun 28, 2033	U-2696			
	9693986	Jun 28, 2033	U-2698			
	9700537	May 31, 2027	U-2707			
	9918954	Jun 28, 2033	U-2699			
<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 002	10010517	Apr 29, 2030	U-2690			
	10265287	Apr 29, 2030	U-2700			
	10278935	Jun 28, 2033	U-2701			
	10278936	Jun 28, 2033	U-2702			
	10278937	Jun 28, 2033	U-2703			
	10383840	Jun 28, 2033	U-2704			
	10555924	Jun 28, 2033	U-2743			
	10555925	Jun 28, 2033	U-2744			
	10568861	Jun 28, 2033	U-2756			
	10576054	Jun 28, 2033	U-2762			
	10668042	Jun 28, 2033	U-2841			
	10786478	Jun 28, 2033	U-2959			
	10786478	Jun 28, 2033	U-2960			
	10792267	Apr 29, 2030	U-2961			
	10792270	Jun 28, 2033	U-2962			

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<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057 002	10842766	Apr 29, 2030				U-2997
	10842768	Jun 15, 2030				U-2688
	10881632	Apr 29, 2030				U-3052
	10894028	Jun 28, 2033				U-3053
	11000499	Jun 28, 2033				U-3126
	11103477	Apr 29, 2030				U-3209
	11116742	Jun 28, 2033				U-3221
	11154526	Apr 29, 2030				U-3240
	11213504	Apr 29, 2030				U-3292
	11298333	Jun 28, 2033				U-3358
	11369582	Jun 28, 2033				U-2841
	11717504	Apr 29, 2030				U-3669
	8293727	Feb 09, 2030				U-1287
	8293728	Feb 09, 2030				U-1287
	8298554	Apr 29, 2030	DP			
	8314086	Feb 09, 2030				U-1287
	8318715	Feb 09, 2030				U-1287
	8357677	Feb 09, 2030				U-1287
	8367652	Feb 09, 2030				U-1287
	8377920	Feb 09, 2030				U-1287
	8399446	Feb 09, 2030				U-1287
	8410086	Jun 15, 2030				U-2688
	8415335	Feb 09, 2030				U-1287
	8426399	Feb 09, 2030				U-1287
	8440650	Feb 09, 2030				U-1287
	8445003	Apr 29, 2030				U-1287
	8445013	Apr 29, 2030				U-1287
	8454994	Apr 29, 2030				U-2689
	8501225	Apr 29, 2030				U-1287
	8518929	Feb 09, 2030				U-1287
	8524698	Feb 09, 2030				U-1287
	8546372	Feb 09, 2030				U-1287
	8551521	Apr 29, 2030				U-1287
	8563608	Apr 29, 2030				U-1287
	8617593	Apr 29, 2030				U-1287
	8617593	Apr 29, 2030				U-2691
	8617594	Apr 29, 2030				U-1287
	8623406	Apr 29, 2030				U-1287
	8623406	Apr 29, 2030				U-2692
	8642077	Apr 29, 2030				U-2693
	8669245	Jun 15, 2030				U-2694
	8680144	Feb 09, 2030				U-2695
	8691871	Apr 29, 2030				U-2689
	8703185	Apr 29, 2030				U-2691
	8709475	Apr 29, 2030				U-2689
	8710041	Jun 15, 2030				U-2690
	9198892	Sep 25, 2027				U-2706
	9603826	Jun 28, 2033				U-2696
	9610272	Jun 28, 2033				U-2697

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<u>ICOSAPENT ETHYL - VASCEPA</u>						
N 202057	002 9623001	Jun 28, 2033				U-2698
	9693984	Jun 28, 2033				U-2697
	9693985	Jun 28, 2033				U-2696
	9693986	Jun 28, 2033				U-2698
	9700537	May 31, 2027				U-2707
	9918954	Jun 28, 2033				U-2699
<u>IDELALISIB - ZYDELIG</u>						
N 205858	001 10730879	Mar 05, 2033		DS DP		
	8865730	Mar 05, 2033		DS DP		U-1615
	8980901	May 12, 2025				U-1678
	9149477	May 12, 2025				U-1757
	9469643	Sep 02, 2033		DS		
	9492449	Mar 11, 2030				U-1914
	RE44599	Jul 21, 2025				U-1558
	RE44599	Jul 21, 2025				U-1615
	RE44638	Aug 05, 2025		DS DP		
<u>IDELALISIB - ZYDELIG</u>						
N 205858	002 10730879	Mar 05, 2033		DS DP		
	8865730	Mar 05, 2033		DS DP		U-1615
	8980901	May 12, 2025				U-1678
	9149477	May 12, 2025				U-1757
	9469643	Sep 02, 2033		DS		
	9492449	Mar 11, 2030				U-1914
	RE44599	Jul 21, 2025				U-1558
	RE44599	Jul 21, 2025				U-1615
	RE44638	Aug 05, 2025		DS DP		
<u>ILOPERIDONE - FANAPT</u>						
N 022192	001 8586610	Nov 02, 2027				U-1625
	8652776	Aug 31, 2030				U-1685
	8999638	Oct 28, 2030				U-1674
	9072742	Jan 16, 2031				U-1674
	9074254	Dec 28, 2031				U-1674
	9074255	Dec 17, 2030				U-1674
	9074256	Feb 10, 2031				U-1674
	9138432	Sep 30, 2025				U-1737
	9157121	Apr 05, 2030				U-1674
<u>ILOPERIDONE - FANAPT</u>						
N 022192	002 8586610	Nov 02, 2027				U-1625
	8652776	Aug 31, 2030				U-1685
	8999638	Oct 28, 2030				U-1674
	9072742	Jan 16, 2031				U-1674
	9074254	Dec 28, 2031				U-1674
	9074255	Dec 17, 2030				U-1674
	9074256	Feb 10, 2031				U-1674
	9138432	Sep 30, 2025				U-1737
	9157121	Apr 05, 2030				U-1674

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<u>ILOPERIDONE - FANAPT</u>						
N 022192	003	8586610				
		Nov 02, 2027	U-1625			
		8652776				
		Aug 31, 2030	U-1685			
		8999638				
		Oct 28, 2030	U-1674			
		9072742				
		Jan 16, 2031	U-1674			
		9074254				
		Dec 28, 2031	U-1674			
		9074255				
		Dec 17, 2030	U-1674			
		9074256				
		Feb 10, 2031	U-1674			
		9138432				
		Sep 30, 2025	U-1737			
		9157121				
		Apr 05, 2030	U-1674			
<u>ILOPERIDONE - FANAPT</u>						
N 022192	004	8586610				
		Nov 02, 2027	U-1625			
		8652776				
		Aug 31, 2030	U-1685			
		8999638				
		Oct 28, 2030	U-1674			
		9072742				
		Jan 16, 2031	U-1674			
		9074254				
		Dec 28, 2031	U-1674			
		9074255				
		Dec 17, 2030	U-1674			
		9074256				
		Feb 10, 2031	U-1674			
		9138432				
		Sep 30, 2025	U-1737			
		9157121				
		Apr 05, 2030	U-1674			
<u>ILOPERIDONE - FANAPT</u>						
N 022192	005	8586610				
		Nov 02, 2027	U-1625			
		8652776				
		Aug 31, 2030	U-1685			
		8999638				
		Oct 28, 2030	U-1674			
		9072742				
		Jan 16, 2031	U-1674			
		9074254				
		Dec 28, 2031	U-1674			
		9074255				
		Dec 17, 2030	U-1674			
		9074256				
		Feb 10, 2031	U-1674			
		9138432				
		Sep 30, 2025	U-1737			
		9157121				
		Apr 05, 2030	U-1674			
<u>ILOPERIDONE - FANAPT</u>						
N 022192	006	8586610				
		Nov 02, 2027	U-1625			
		8652776				
		Aug 31, 2030	U-1685			
		8999638				
		Oct 28, 2030	U-1674			
		9072742				
		Jan 16, 2031	U-1674			
		9074254				
		Dec 28, 2031	U-1674			
		9074255				
		Dec 17, 2030	U-1674			
		9074256				
		Feb 10, 2031	U-1674			
		9138432				
		Sep 30, 2025	U-1737			
		9157121				
		Apr 05, 2030	U-1674			
<u>ILOPERIDONE - FANAPT</u>						
N 022192	007	8586610				
		Nov 02, 2027	U-1625			
		8652776				
		Aug 31, 2030	U-1685			
		8999638				
		Oct 28, 2030	U-1674			
		9072742				
		Jan 16, 2031	U-1674			
		9074254				
		Dec 28, 2031	U-1674			
		9074255				
		Dec 17, 2030	U-1674			
		9074256				
		Feb 10, 2031	U-1674			
		9138432				
		Sep 30, 2025	U-1737			

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<u>ILOPERIDONE - FANAPT</u>						
N 022192	007 9157121	Apr 05, 2030	U-1674			
<u>IMIQUIMOD - ALDARA</u>						
N 020723	001 7696159	Apr 01, 2024	DS U-1047			
	7696159	Apr 01, 2024	DS U-1048			
<u>IMIQUIMOD - ZYCLARA</u>						
N 022483	001 10238644	Dec 11, 2029	U-68			
	10238645	Aug 18, 2029	U-1455			
	10238645	Aug 18, 2029	U-172			
	10918635	Apr 30, 2030	U-1455			
	10918635	Apr 30, 2030	U-172			
	11202752	Apr 30, 2030	U-1455			
	11202752	Apr 30, 2030	U-172			
	8236816	Dec 11, 2029	U-68			
	8299109	Dec 11, 2029	U-68			
	8598196	Aug 18, 2029	U-1455			
	8598196	Aug 18, 2029	U-172			
<u>IMIQUIMOD - ZYCLARA</u>						
N 022483	002 11318130	Dec 11, 2029	U-68			
	8222270	Dec 11, 2029	U-68			
<u>INCLISIRAN SODIUM - LEOVIO</u>						
N 214012	001 10125369	Aug 18, 2034	DS DP U-3652		NCE	Dec 22, 2026
	10131907	Aug 24, 2028	DS DP U-3652			
	10273477	Mar 08, 2024	DS			
	10669544	Mar 08, 2024	DS			
	10806791	Dec 04, 2028	DS			
	10851377	Aug 25, 2036	U-3652			
	11530408	May 18, 2024	DS			
	8106022	Dec 12, 2029	DS DP U-3652			
	8222222	Dec 29, 2027	U-3652			
	8809292	May 10, 2027	DS DP U-3652			
	8828956	Dec 04, 2028	DS DP U-3652			
	9370582	Dec 04, 2028	DS DP U-3652			
	9708615	Mar 08, 2024	DS			
<u>INDACATEROL MALEATE - ARCAPTA NEOHALER</u>						
N 022383	001 6878721	Feb 25, 2025	DS DP U-1168			
	8479730	Oct 11, 2028	DP			
<u>INDIGOTINDISULFONATE SODIUM - BLUDIGO</u>						
N 216264	001 10927258	Dec 23, 2037	DS		NCE	Jul 08, 2027
	11499050	Dec 23, 2037	DS			
	11845867	Nov 25, 2036	DS			
<u>INDOCYANINE GREEN - SPY AGENT GREEN KIT</u>						
N 211580	001 10631746	Aug 04, 2035	DP U-2815		I-911	Jun 05, 2026
	11712320	Jul 14, 2039	DP			
	8185176	Jun 04, 2028	U-2462			
	8406860	Apr 09, 2029	U-2463			
	8647605	Feb 11, 2029	U-2464			
	8647605	Feb 11, 2029	U-2468			

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<u>INDOCYANINE GREEN - SPY AGENT GREEN KIT</u>						
N 211580	001 9421280	Nov 24, 2025	U-2466			
	9421280	Nov 24, 2025	U-2467			
<u>INDOMETHACIN - INDOMETHACIN</u>						
A 216184	001				CGT	Jan 30, 2024
<u>INDOMETHACIN - TIVORBEX</u>						
N 204768	001 8734847	Apr 23, 2030	DP			
	8992982	Apr 23, 2030	DP			
	9089471	Apr 23, 2030	U-55			
<u>INDOMETHACIN - TIVORBEX</u>						
N 204768	002 8734847	Apr 23, 2030	DP			
	8992982	Apr 23, 2030	DP			
	9089471	Apr 23, 2030	U-55			
<u>INFIGRATINIB PHOSPHATE - TRUSELTIO</u>						
N 214622	001 10278969	Dec 11, 2034	DP		NCE	May 28, 2026
	11160804	Dec 11, 2034	DP		ODE-353	May 28, 2028
	8552002	Aug 25, 2029	DS DP			
	9067896	Aug 06, 2028	DS			
<u>INFIGRATINIB PHOSPHATE - TRUSELTIO</u>						
N 214622	002 10278969	Dec 11, 2034	DP		NCE	May 28, 2026
	11160804	Dec 11, 2034	DP		ODE-353	May 28, 2028
	8552002	Aug 25, 2029	DS DP			
	9067896	Aug 06, 2028	DS			
<u>INGENOL MEBUTATE - PICATO</u>						
N 202833	001 8278292	Jul 06, 2027	DP			
	8372827	Dec 18, 2026	DP			
	8372828	Dec 18, 2026	DP			
	8377919	Dec 18, 2026	DP			
	8536163	Dec 18, 2026	U-1440			
	8716271	Dec 18, 2026	U-1440			
	8735375	Dec 18, 2026	U-1440			
	9789078	May 15, 2033	U-2138			
	9820959	Dec 18, 2026	DP U-1440			
	9833428	Dec 18, 2026	DP			
	9833429	Dec 18, 2026	DP			
	9861603	Dec 18, 2026	U-1440			
<u>INGENOL MEBUTATE - PICATO</u>						
N 202833	002 8278292	Jul 06, 2027	DP			
	8372827	Dec 18, 2026	DP			
	8372828	Dec 18, 2026	DP			
	8377919	Dec 18, 2026	DP			
	8536163	Dec 18, 2026	U-1440			
	8716271	Dec 18, 2026	U-1440			
	8735375	Dec 18, 2026	U-1440			
	9820959	Dec 18, 2026	DP U-1440			
	9833428	Dec 18, 2026	DP			
	9833429	Dec 18, 2026	DP			
	9861603	Dec 18, 2026	U-1440			

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<u>INGENOL MEBUTATE - PICATO</u>						
N 202833	002	8278292	Jul 06, 2027	DP		
		8372827	Dec 18, 2026	DP		
		8372828	Dec 18, 2026	DP		
		8377919	Dec 18, 2026	DP		
		8536163	Dec 18, 2026	U-1440		
		8716271	Dec 18, 2026	U-1440		
		8735375	Dec 18, 2026	U-1440		
		9820959	Dec 18, 2026	DP U-1440		
		9833428	Dec 18, 2026	DP		
		9833429	Dec 18, 2026	DP		
		9861603	Dec 18, 2026	U-1440		
<u>INOTERSEN SODIUM - TEGSEDI</u>						
N 211172	001	8101743	Apr 01, 2025	DS DP	ODE-212	Oct 05, 2025
		8697860	Apr 29, 2031	DP		
		9061044	Apr 29, 2031	DS		
		9399774	Apr 29, 2031	U-2430		
<u>IOBENGUANE I-131 - AZEDRA</u>						
N 209607	001				ODE-204	Jul 30, 2025
<u>IPRATROPIUM BROMIDE - ATROVENT HFA</u>						
N 021527	001	8474447	Jan 17, 2030	DP		
<u>IPRATROPIUM BROMIDE - ATROVENT HFA</u>						
N 218276	001	11603363	May 25, 2041	DS DP		
		9682968	Jul 14, 2034	DS DP		
<u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u>						
N 207793	001	10456360	Oct 15, 2036	DP		
		10722508	May 02, 2025	DS DP		
		10980795	Jun 12, 2033	U-1848		
		10993914	Oct 15, 2036	DP		
		11369597	Jun 12, 2033	U-1848		
		8147867	Aug 29, 2028	DS DP		
		8329213	Jan 06, 2027	DS DP		
		8703181	May 02, 2025	U-1434		
		8992970	May 02, 2025	DS DP		
		9339497	Jun 12, 2033	U-1848		
		9364473	Jun 12, 2033	U-1856		
		9452162	Jun 12, 2033	U-1899		
		9492442	Jun 12, 2033	U-1848		
		9492442	Jun 12, 2033	U-1899		
		9492442	Jun 12, 2033	U-1917		
		9717724	Jun 12, 2033	U-1848		
		9717724	Jun 12, 2033	U-2091		
		9724303	May 02, 2025	DS DP		
		9730891	May 02, 2025	U-1848		
		9782349	May 02, 2025	DS DP		
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207500	001	10206879	Sep 14, 2027	DP	NCE	Mar 06, 2020
		10206879*PED	Mar 14, 2028		ODE-305	Mar 06, 2022

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<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207500 001	10603280	Sep 14, 2027	DP		ODE-90	Mar 06, 2022
	10603280*PED	Mar 14, 2028			PED	Sep 06, 2020
	6812238	Oct 31, 2025	DS		PED	Sep 06, 2022
	6812238*PED	May 01, 2026			PED	Sep 06, 2022
					GAIN	Sep 06, 2025
					GAIN	Sep 06, 2027
					GAIN	Sep 06, 2027
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207500 002	10206879	Sep 14, 2027	DP		NCE	Mar 06, 2020
	10206879*PED	Mar 14, 2028			ODE*	Mar 06, 2022
	10603280	Sep 14, 2027	DP		PED	Sep 06, 2020
	10603280*PED	Mar 14, 2028			PED	Sep 06, 2022
	6812238	Oct 31, 2025	DS		GAIN	Sep 06, 2025
	6812238*PED	May 01, 2026			GAIN	Sep 06, 2027
<u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u>						
N 207501 001	6812238	Oct 31, 2025	DS		NCE	Mar 06, 2020
	6812238*PED	May 01, 2026			ODE-305	Mar 06, 2022
					ODE-90	Mar 06, 2022
					PED	Sep 06, 2020
					PED	Sep 06, 2022
					PED	Sep 06, 2022
					GAIN	Sep 06, 2025
					GAIN	Sep 06, 2027
					GAIN	Sep 06, 2027
<u>ISOPROPYL ALCOHOL - ZURAGARD</u>						
N 210872 001	10688291	Dec 20, 2034	DP U-1397		M-268	Jun 24, 2024
	8226971	May 06, 2025	DP			
	8389583	Aug 09, 2029	U-1397			
	8703828	May 23, 2028	DP			
	9011897	Feb 08, 2025	DP			
	9629368	May 23, 2028	U-1397			
	9844654	Apr 24, 2036	DP U-1397			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 001	9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 002	9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 003	9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 004	9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913 005	9700535	Aug 04, 2035	DP			

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<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	005 9750711	May 29, 2035	DP			
<u>ISOTRETINOIN - ABSORICA LD</u>						
N 211913	006 9700535	Aug 04, 2035	DP			
	9750711	May 29, 2035	DP			
<u>ISTRADEFYLLINE - NOURIANZ</u>						
N 022075	001 7541363	Nov 13, 2024	DS DP		NCE	Aug 27, 2024
	7727993	Jan 28, 2028	DP U-2623			
	8318201	Sep 05, 2027	DP			
<u>ISTRADEFYLLINE - NOURIANZ</u>						
N 022075	002 7541363	Nov 13, 2024	DS DP		NCE	Aug 27, 2024
	7727993	Jan 28, 2028	DP U-2623			
	8318201	Sep 05, 2027	DP			
<u>ITRACONAZOLE - ONMEL</u>						
N 022484	001 8486456	Oct 03, 2028	DP U-1054			
<u>ITRACONAZOLE - TOLSURA</u>						
N 208901	001 10463740	Jun 21, 2033	DP U-2453			
	10806792	Jun 21, 2033	DP			
	8921374	Jun 21, 2033	DP			
	9272046	Jun 21, 2033	DP			
	9713642	Jun 21, 2033	U-2453			
<u>IVABRADINE - CORLANOR</u>						
N 209964	001 7361649	Feb 22, 2026	DS DP U-1694		ODE-234	Apr 22, 2026
	7361650	Feb 22, 2026	DS DP U-1694		PED	Oct 22, 2026
	7867996	Dec 12, 2026	DS DP U-1694			
	7879842	Feb 22, 2026	DS DP U-1694			
<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143	001 7361649	Feb 22, 2026	DS DP U-1694			
	7361649*PED	Aug 22, 2026				
	7361650	Feb 22, 2026	DS DP U-1694			
	7361650*PED	Aug 22, 2026				
	7867996	Dec 12, 2026	DS DP U-1694			
	7867996*PED	Jun 12, 2027				
	7879842	Feb 22, 2026	DS DP U-1694			
	7879842*PED	Aug 22, 2026				
<u>IVABRADINE HYDROCHLORIDE - CORLANOR</u>						
N 206143	002 7361649	Feb 22, 2026	DS DP U-1694			
	7361649*PED	Aug 22, 2026				
	7361650	Feb 22, 2026	DS DP U-1694			
	7361650*PED	Aug 22, 2026				
	7867996	Dec 12, 2026	DS DP U-1694			
	7867996*PED	Jun 12, 2027				
	7879842	Feb 22, 2026	DS DP U-1694			
	7879842*PED	Aug 22, 2026				
<u>IVACAFTOR - KALYDECO</u>						
N 203188	001 10646481	Aug 13, 2029	DP		ODE-189	Jul 31, 2024
	11564916	Aug 13, 2029	U-3530		ODE-190	May 17, 2024

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<u>IVACAFTOR - KALYDECO</u>						
N 203188 001	7495103	May 20, 2027	DS DP		ODE-199	Aug 15, 2025
	8324242	Aug 05, 2027		U-1311	ODE-338	Dec 21, 2027
	8324242	Aug 05, 2027		U-1906		
	8354427	Jul 06, 2026		U-1311		
	8354427	Jul 06, 2026		U-1905		
	8410274	Dec 28, 2026		DP		
	8629162	Jun 24, 2025		U-2234		
	8754224	Dec 28, 2026	DS DP			
	9670163	Dec 28, 2026	DP	U-1311		
<u>IVACAFTOR - KALYDECO</u>						
N 207925 001	10272046	Feb 27, 2033	DP	U-2531	ODE-189	Jul 31, 2024
	10646481	Aug 13, 2029		DP	ODE-190	May 17, 2024
	11147770	Feb 27, 2033	DP	U-3339	ODE-199	Aug 15, 2025
	11564916	Aug 13, 2029		U-3528	ODE-236	Apr 29, 2026
	11752106	Feb 27, 2033	DP	U-3697	ODE-338	Dec 21, 2027
	7495103	May 20, 2027	DS DP		ODE-435	May 03, 2030
	8324242	Aug 05, 2027		U-1311		
	8324242	Aug 05, 2027		U-1906		
	8324242	Aug 05, 2027		U-2527		
	8354427	Jul 06, 2026		U-1311		
	8354427	Jul 06, 2026		U-1905		
	8354427	Jul 06, 2026		U-2528		
	8410274	Dec 28, 2026		DP		
	8629162	Jun 24, 2025		U-2234		
	8629162	Jun 24, 2025		U-2529		
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033		DP		
	9670163	Dec 28, 2026	DP	U-1311		
	9670163	Dec 28, 2026	DP	U-2530		
<u>IVACAFTOR - KALYDECO</u>						
N 207925 002	10272046	Feb 27, 2033	DP	U-2531	ODE-189	Jul 31, 2024
	10646481	Aug 13, 2029		DP	ODE-190	May 17, 2024
	11147770	Feb 27, 2033	DP	U-3339	ODE-199	Aug 15, 2025
	11564916	Aug 13, 2029		U-3528	ODE-236	Apr 29, 2026
	11752106	Feb 27, 2033	DP	U-3697	ODE-338	Dec 21, 2027
	7495103	May 20, 2027	DS DP		ODE-435	May 03, 2030
	8324242	Aug 05, 2027		U-1311		
	8324242	Aug 05, 2027		U-1906		
	8324242	Aug 05, 2027		U-2527		
	8354427	Jul 06, 2026		U-1311		
	8354427	Jul 06, 2026		U-1905		
	8354427	Jul 06, 2026		U-2528		
	8410274	Dec 28, 2026		DP		
	8629162	Jun 24, 2025		U-2234		
	8629162	Jun 24, 2025		U-2529		
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033		DP		
	9670163	Dec 28, 2026	DP	U-1311		
	9670163	Dec 28, 2026	DP	U-2530		

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<u>IVACAFTOR - KALYDECO</u>						
N 207925 002	10272046	Feb 27, 2033	DP U-2531		ODE-189	Jul 31, 2024
	10646481	Aug 13, 2029	DP		ODE-190	May 17, 2024
	11147770	Feb 27, 2033	DP U-3339		ODE-199	Aug 15, 2025
	11564916	Aug 13, 2029	U-3528		ODE-236	Apr 29, 2026
	11752106	Feb 27, 2033	DP U-3697		ODE-338	Dec 21, 2027
	7495103	May 20, 2027	DS DP		ODE-435	May 03, 2030
	8324242	Aug 05, 2027	U-1311			
	8324242	Aug 05, 2027	U-1906			
	8324242	Aug 05, 2027	U-2527			
	8354427	Jul 06, 2026	U-1311			
	8354427	Jul 06, 2026	U-1905			
	8354427	Jul 06, 2026	U-2528			
	8410274	Dec 28, 2026	DP			
	8629162	Jun 24, 2025	U-2234			
	8629162	Jun 24, 2025	U-2529			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-1311			
	9670163	Dec 28, 2026	DP U-2530			
<u>IVACAFTOR - KALYDECO</u>						
N 207925 003	10272046	Feb 27, 2033	DP U-2967		ODE-189	Jul 31, 2024
	10646481	Aug 13, 2029	DP		ODE-190	May 17, 2024
	11147770	Feb 27, 2033	DP U-3339		ODE-199	Aug 15, 2025
	11564916	Aug 13, 2029	U-3528		ODE-236	Apr 29, 2026
	11752106	Feb 27, 2033	DP U-3697		ODE-338	Dec 21, 2027
	7495103	May 20, 2027	DS DP		ODE-435	May 03, 2030
	8324242	Aug 05, 2027	U-1311			
	8324242	Aug 05, 2027	U-1906			
	8324242	Aug 05, 2027	U-2963			
	8354427	Jul 06, 2026	U-1311			
	8354427	Jul 06, 2026	U-1905			
	8354427	Jul 06, 2026	U-2964			
	8410274	Dec 28, 2026	DP			
	8629162	Jun 24, 2025	U-2234			
	8629162	Jun 24, 2025	U-2965			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-1311			
	9670163	Dec 28, 2026	DP U-2966			
<u>IVACAFTOR - KALYDECO</u>						
N 207925 004	10272046	Feb 27, 2033	DP U-3605		NPP	May 03, 2026
	10646481	Aug 13, 2029	DP		ODE-435	May 03, 2030
	11147770	Feb 27, 2033	DP U-3604			
	11564916	Aug 13, 2029	U-3603			
	11752106	Feb 27, 2033	DP U-3697			
	7495103	May 20, 2027	DS DP			
	8324242	Aug 05, 2027	U-3609			
	8354427	Jul 06, 2026	U-3608			
	8410274	Dec 28, 2026	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IVACAFTOR - KALYDECO</u>						
N 207925	004	8629162	Jun 24, 2025	U-3607		
		8754224	Dec 28, 2026	DS DP		
		8883206	Feb 27, 2033	DP		
		9670163	Dec 28, 2026	DP U-3606		
<u>IVACAFTOR - KALYDECO</u>						
N 207925	005	10272046	Feb 27, 2033	DP U-3605	NPP	May 03, 2026
		10646481	Aug 13, 2029	DP	ODE-435	May 03, 2030
		11147770	Feb 27, 2033	DP U-3604		
		11564916	Aug 13, 2029	U-3603		
		11752106	Feb 27, 2033	DP U-3697		
		7495103	May 20, 2027	DS DP		
		8324242	Aug 05, 2027	U-3609		
		8354427	Jul 06, 2026	U-3608		
		8410274	Dec 28, 2026	DP		
		8629162	Jun 24, 2025	U-3607		
		8754224	Dec 28, 2026	DS DP		
		8883206	Feb 27, 2033	DP		
		9670163	Dec 28, 2026	DP U-3606		
<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	001	10022352	Apr 09, 2027	DP U-2343	ODE-173	Feb 12, 2025
		10022352	Apr 09, 2027	DP U-2573	ODE-247	Jun 21, 2026
		10058546	Jul 15, 2033	U-2399	ODE-335	Dec 21, 2027
		10058546	Jul 15, 2033	U-2572		
		10058546	Jul 15, 2033	U-3022		
		10058546	Jul 15, 2033	U-3023		
		10081621	Mar 25, 2031	DP U-2420		
		10081621	Mar 25, 2031	DP U-2571		
		10081621	Mar 25, 2031	DP U-3024		
		10081621	Mar 25, 2031	DP U-3025		
		10206877	Apr 14, 2035	DP U-2498		
		10206877	Apr 14, 2035	DP U-2570		
		10206877	Apr 14, 2035	DP U-3026		
		10206877	Apr 14, 2035	DP U-3027		
		10239867	Apr 09, 2027	DS DP U-2512		
		10239867	Apr 09, 2027	DS DP U-2569		
		10646481	Aug 13, 2029	DP		
		11564916	Aug 13, 2029	U-3527		
		11578062	Mar 25, 2031	DP U-3545		
		11639347	Apr 09, 2027	DS DP U-2569		
		7495103	May 20, 2027	DS DP		
		7645789	May 01, 2027	DS DP		
		7776905	Jun 03, 2027	DS DP		
		8324242	Aug 05, 2027	U-2246		
		8354427	Jul 06, 2026	U-3021		
		8410274	Dec 28, 2026	DP		
		8415387	Nov 12, 2027	U-2246		
		8598181	May 01, 2027	U-2246		
		8623905	May 01, 2027	DS DP		
		8629162	Jun 24, 2025	U-2247		

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<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	001	8754224	Dec 28, 2026	DS DP		
		9012496	Jul 15, 2033		U-2248	
		9670163	Dec 28, 2026	DP	U-2246	
		9931334	Dec 28, 2026	DP	U-2275	
		9931334	Dec 28, 2026	DP	U-2575	
		9974781	Apr 09, 2027	DP	U-2318	
		9974781	Apr 09, 2027	DP	U-2574	
<u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u>						
N 210491	002	10022352	Apr 09, 2027	DP	U-2343	ODE-173 Feb 12, 2025
		10022352	Apr 09, 2027	DP	U-2573	ODE-247 Jun 21, 2026
		10058546	Jul 15, 2033		U-2399	ODE-335 Dec 21, 2027
		10058546	Jul 15, 2033		U-2572	
		10058546	Jul 15, 2033		U-3022	
		10058546	Jul 15, 2033		U-3023	
		10081621	Mar 25, 2031	DP	U-2420	
		10081621	Mar 25, 2031	DP	U-2571	
		10081621	Mar 25, 2031	DP	U-3024	
		10081621	Mar 25, 2031	DP	U-3025	
		10206877	Apr 14, 2035	DP	U-2498	
		10206877	Apr 14, 2035	DP	U-2570	
		10206877	Apr 14, 2035	DP	U-3026	
		10206877	Apr 14, 2035	DP	U-3027	
		10239867	Apr 09, 2027	DS DP	U-2512	
		10239867	Apr 09, 2027	DS DP	U-2569	
		10646481	Aug 13, 2029	DP		
		11564916	Aug 13, 2029		U-3527	
		11578062	Mar 25, 2031	DP	U-3545	
		11639347	Apr 09, 2027	DS DP	U-2569	
		7495103	May 20, 2027	DS DP		
		7645789	May 01, 2027	DS DP		
		7776905	Jun 03, 2027	DS DP		
		8324242	Aug 05, 2027		U-2246	
		8354427	Jul 06, 2026		U-3021	
		8410274	Dec 28, 2026	DP		
		8415387	Nov 12, 2027		U-2246	
		8598181	May 01, 2027		U-2246	
		8623905	May 01, 2027	DS DP		
		8629162	Jun 24, 2025		U-2247	
		8754224	Dec 28, 2026	DS DP		
		9012496	Jul 15, 2033		U-2248	
		9670163	Dec 28, 2026	DP	U-2246	
		9931334	Dec 28, 2026	DP	U-2275	
		9931334	Dec 28, 2026	DP	U-2575	
		9974781	Apr 09, 2027	DP	U-2318	
		9974781	Apr 09, 2027	DP	U-2574	
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 206038	001	10076513	Dec 04, 2028	DP	U-2411	
		10597384	Dec 04, 2028	DS DP	U-2777	
		10646481	Aug 13, 2029	DP		

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<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 206038	001	11052075	Dec 04, 2028	DP	U-3181	
		11564916	Aug 13, 2029		U-3529	
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026		U-1973	
		8324242	Aug 05, 2027		U-1311	
		8324242	Aug 05, 2027		U-1911	
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		
		8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP	U-1718	
		8716338	Sep 20, 2030	DP	U-1910	
		8741933	Nov 08, 2026		U-1717	
		8741933	Nov 08, 2026		U-1909	
		8754224	Dec 28, 2026	DS DP		
		8846718	Jul 02, 2029		U-1717	
		8846718	Jul 02, 2029		U-1908	
		8993600	Dec 11, 2030	DP		
		9150552	Dec 04, 2028		U-1908	
		9192606	Sep 29, 2029	DP	U-1912	
		9216969	Nov 08, 2026	DS DP		
		9670163	Dec 28, 2026	DP	U-1911	
		9931334	Dec 28, 2026	DP	U-2276	
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 206038	002	10597384	Dec 04, 2028	DS DP	U-2777	
		10646481	Aug 13, 2029	DP		
		11052075	Dec 04, 2028	DP	U-3181	
		11564916	Aug 13, 2029		U-3529	
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026		U-1973	
		8324242	Aug 05, 2027		U-1911	
		8410274	Dec 28, 2026	DP		
		8507534	Sep 20, 2030	DS DP		
		8653103	Dec 04, 2028	DP		
		8716338	Sep 20, 2030	DP	U-1910	
		8741933	Nov 08, 2026		U-1909	
		8754224	Dec 28, 2026	DS DP		
		8846718	Jul 02, 2029		U-1908	
		8993600	Dec 11, 2030	DP		
		9150552	Dec 04, 2028		U-1908	
		9192606	Sep 29, 2029	DP	U-1912	
		9216969	Nov 08, 2026	DP		
		9670163	Dec 28, 2026	DP	U-1911	
		9931334	Dec 28, 2026	DP	U-2276	
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	001	10597384	Dec 04, 2028	DS DP	U-2778	NPP Sep 02, 2025
		10646481	Aug 13, 2029	DP		ODE-195 Aug 07, 2025
		11564916	Aug 13, 2029		U-3526	ODE-408 Sep 02, 2029
		7495103	May 20, 2027	DS DP		
		7973038	Nov 08, 2026		U-2374	

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<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358 001	8324242	Aug 05, 2027	U-2374			
	8410274	Dec 28, 2026	DP			
	8507534	Sep 20, 2030	DS DP			
	8653103	Dec 04, 2028	DP			
	8716338	Sep 20, 2030	DP U-2396			
	8741933	Nov 08, 2026	U-2374			
	8754224	Dec 28, 2026	DS DP			
	8846718	Jul 02, 2029	U-2375			
	8993600	Dec 11, 2030	DP			
	9150552	Dec 04, 2028	U-2375			
	9192606	Sep 29, 2029	DP U-2397			
	9216969	Nov 08, 2026	DP			
	9670163	Dec 28, 2026	DP U-2376			
	9931334	Dec 28, 2026	DP U-2376			
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358 002	10597384	Dec 04, 2028	DS DP U-2778		NPP	Sep 02, 2025
	10646481	Aug 13, 2029	DP		ODE-195	Aug 07, 2025
	11564916	Aug 13, 2029	U-3526		ODE-408	Sep 02, 2029
	7495103	May 20, 2027	DS DP			
	7973038	Nov 08, 2026	U-2374			
	8324242	Aug 05, 2027	U-2374			
	8410274	Dec 28, 2026	DP			
	8507534	Sep 20, 2030	DS DP			
	8653103	Dec 04, 2028	DP			
	8716338	Sep 20, 2030	DP U-2396			
	8741933	Nov 08, 2026	U-2374			
	8754224	Dec 28, 2026	DS DP			
	8846718	Jul 02, 2029	U-2375			
	8993600	Dec 11, 2030	DP			
	9150552	Dec 04, 2028	U-2375			
	9192606	Sep 29, 2029	DP U-2397			
	9216969	Nov 08, 2026	DP			
	9670163	Dec 28, 2026	DP U-2376			
	9931334	Dec 28, 2026	DP U-2376			
<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358 003	10597384	Dec 04, 2028	DS DP U-3430		NS	Sep 02, 2025
	10646481	Aug 13, 2029	DP		ODE-408	Sep 02, 2029
	11564916	Aug 13, 2029	U-3526			
	7495103	May 20, 2027	DS DP			
	7973038	Nov 08, 2026	U-3424			
	8324242	Aug 05, 2027	U-3424			
	8410274	Dec 28, 2026	DP			
	8507534	Sep 20, 2030	DS DP			
	8653103	Dec 04, 2028	DP			
	8716338	Sep 20, 2030	DP U-3426			
	8741933	Nov 08, 2026	U-3424			
	8754224	Dec 28, 2026	DS DP			
	8846718	Jul 02, 2029	U-3427			
	8993600	Dec 11, 2030	DP			

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<u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u>						
N 211358	003 9150552	Dec 04, 2028	U-3427			
	9192606	Sep 29, 2029	DP U-3428			
	9216969	Nov 08, 2026	DP			
	9670163	Dec 28, 2026	DP U-3429			
	9931334	Dec 28, 2026	DP U-3429			
<u>IVERMECTIN - SOOLANTRA</u>						
N 206255	001 10206939	Mar 13, 2034	U-1631			
	11033565	Apr 22, 2024	DP			
	7550440	Apr 22, 2024	DP U-1631			
	8080530	Apr 22, 2024	DP U-1631			
	8093219	Apr 22, 2024	DP U-1631			
	8415311	Apr 22, 2024	DP U-1631			
	8470788	Apr 22, 2024	DP U-1631			
	8815816	Apr 22, 2024	DP U-1631			
	9089587	Mar 13, 2034	U-1631			
	9233117	Mar 13, 2034	U-1631			
	9233118	Mar 13, 2034	U-1631			
	9782425	Mar 13, 2034	U-1631			
<u>IVOSIDENIB - TIBSOVO</u>						
N 211192	001 10449184	Mar 13, 2035	DP		I-875	Aug 25, 2024
	10610125	Jun 21, 2030	U-2784		I-893	May 25, 2025
	10610125	Jun 21, 2030	U-2785		I-924	Oct 24, 2026
	10610125	Jun 21, 2030	U-3385		ODE-203	Jul 20, 2025
	10653710	Oct 18, 2036	U-3387		ODE-242	May 02, 2026
	10717764	Jan 18, 2033	U-3215		ODE-368	Aug 25, 2028
	10799490	Mar 13, 2035	DP U-2981		ODE-447	Oct 24, 2030
	10799490	Mar 13, 2035	DP U-2982			
	10799490	Mar 13, 2035	DP U-3384			
	10980788	Jun 07, 2039	U-3112			
	10980788	Jun 07, 2039	U-3113			
	10980788	Jun 07, 2039	U-3214			
	10980788	Jun 07, 2039	U-3383			
	10980788	Jun 07, 2039	U-3743			
	11667673	Jan 18, 2033	U-3742			
	9474779	Aug 19, 2033	DS DP U-2350			
	9474779	Aug 19, 2033	DS DP U-2533			
	9474779	Aug 19, 2033	DS DP U-2534			
	9474779	Aug 19, 2033	DS DP U-3213			
	9474779	Aug 19, 2033	DS DP U-3386			
	9474779	Aug 19, 2033	DS DP U-3742			
	9850277	Jan 18, 2033	DS DP U-2350			
	9850277	Jan 18, 2033	DS DP U-2533			
	9850277	Jan 18, 2033	DS DP U-2534			
	9850277	Jan 18, 2033	DS DP U-3213			
	9850277	Jan 18, 2033	DS DP U-3386			
	9850277	Jan 18, 2033	DS DP U-3742			
	9968595	Mar 13, 2035	DP U-2351			
	9968595	Mar 13, 2035	DP U-2533			
	9968595	Mar 13, 2035	DP U-2534			

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<u>IVOSIDENIB - TIBSOVO</u>						
N 211192	001 9968595	Mar 13, 2035	DP U-3384			
<u>IXABEPILONE - IXEMPRA KIT</u>						
N 022065	001 7312237	Aug 21, 2024	U-965			
<u>IXABEPILONE - IXEMPRA KIT</u>						
N 022065	002 7312237	Aug 21, 2024	U-965			
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462	001 7442830	Nov 20, 2029	DS DP U-2434			
	7687662	Aug 06, 2027	DS DP			
	8003819	Aug 06, 2027	DS DP U-2434			
	8530694	Aug 06, 2027	DS DP U-2434			
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS DP			
	8871745	Aug 06, 2027	U-2434			
	9175017	Jun 16, 2029	U-2434			
	9233115	Aug 12, 2024	U-2434			
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462	002 7442830	Nov 20, 2029	DS DP U-2434			
	7687662	Aug 06, 2027	DS DP			
	8003819	Aug 06, 2027	DS DP U-2434			
	8530694	Aug 06, 2027	DS DP U-2434			
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS DP			
	8871745	Aug 06, 2027	U-2434			
	9175017	Jun 16, 2029	U-2434			
	9233115	Aug 12, 2024	U-2434			
<u>IXAZOMIB CITRATE - NINLARO</u>						
N 208462	003 7442830	Nov 20, 2029	DS DP U-2434			
	7687662	Aug 06, 2027	DS DP			
	8003819	Aug 06, 2027	DS DP U-2434			
	8530694	Aug 06, 2027	DS DP U-2434			
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS DP			
	8871745	Aug 06, 2027	U-2434			
	9175017	Jun 16, 2029	U-2434			
	9233115	Aug 12, 2024	U-2434			
<u>KETOROLAC TROMETHAMINE - ACULAR LS</u>						
N 021528	001 8008338	May 24, 2027	DS DP U-1181			
	8207215	May 28, 2024	U-1251			
	8377982	May 28, 2024	U-1363			
	8377982*PED	Nov 28, 2024				
	8541463	May 28, 2024	U-1441			
	8541463*PED	Nov 28, 2024				
	8648107	May 28, 2024	DP			
	8906950	May 28, 2024	U-1626			
	8946281	May 28, 2024	U-1662			
	9216167	May 28, 2024	U-1800			

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<u>KETOROLAC TROMETHAMINE - ACUVAIL</u>						
N 022427	001 7842714	Aug 15, 2029	DS DP			
	8512717	Mar 07, 2028	DP			
	8992952	Aug 05, 2024	DP			
	9192571	Mar 07, 2028	DP			
<u>KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE - OMIIDRIA</u>						
N 205388	001 8173707*PED	Jan 30, 2024				
	8586633*PED	Jan 30, 2024				
	9066856	Oct 23, 2033	DP			
	9066856*PED	Apr 23, 2034				
	9278101*PED	Jan 30, 2024				
	9399040*PED	Jan 30, 2024				
	9486406	Oct 23, 2033	DP			
	9486406*PED	Apr 23, 2034				
	9855246	Oct 23, 2033	DP			
<u>KETOTIFEN FUMARATE - ACUVUE THERAVISION WITH KETOTIFEN</u>						
N 022388	001 9474746	Mar 27, 2028	DP		NP	Feb 25, 2025
	9962376	Jun 27, 2030	DP			
<u>L-GLUTAMINE - ENDARI</u>						
N 208587	001				ODE-150	Jul 07, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	001				D-188	Apr 28, 2026
					NPP	Oct 14, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	002				D-188	Apr 28, 2026
					NPP	Oct 14, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	003				D-188	Apr 28, 2026
					NPP	Oct 14, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022253	004				D-188	Apr 28, 2026
					NPP	Oct 14, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022254	001				D-188	Apr 28, 2026
					NPP	Oct 14, 2024
<u>LACOSAMIDE - VIMPAT</u>						
N 022255	001				D-188	Apr 28, 2026
					NPP	Oct 14, 2024
<u>LACOSAMIDE - MOTPOLY XR</u>						
N 216185	001 11337943	Jun 05, 2040	DP U-3660			
<u>LACOSAMIDE - MOTPOLY XR</u>						
N 216185	002 11337943	Jun 05, 2040	DP U-3660			
<u>LACOSAMIDE - MOTPOLY XR</u>						
N 216185	003 11337943	Jun 05, 2040	DP U-3660			

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<u>LACTITOL - PIZENSY</u>						
N 211281	001 10806743	May 12, 2037	U-1516		NCE	Feb 12, 2025
<u>LAMIVUDINE; RALTEGRAVIR POTASSIUM - DUTREBIS</u>						
N 206510	001 7169780*PED	Apr 03, 2024				
	7754731	Mar 11, 2029	DS DP U-1663			
	7754731*PED	Sep 11, 2029				
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	001 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	002 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	003 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	004 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	005 8637512	Jun 14, 2028	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N 022115	006 8637512	Jun 14, 2028	DP			
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	001				ODE-156	Sep 15, 2024
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	002				ODE-156	Sep 15, 2024
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
N 022074	003				ODE-156	Sep 15, 2024
<u>LANSOPRAZOLE - LANSOPRAZOLE</u>						
N 208025	001 11077055	Apr 21, 2036	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	001 7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	002 7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	003 7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 021468	004 7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 204734	001 7465465	Aug 26, 2024	DP			
	8980327	Dec 01, 2030	DP			
	9023397	Dec 01, 2030	DP			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 204734	002 7465465	Aug 26, 2024	DP			

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<u>LANTHANUM CARBONATE - FOSRENOL</u>						
N 204734	002 8980327	Dec 01, 2030	DP			
	9023397	Dec 01, 2030	DP			
<u>LAPATINIB DITOSYLATE - TYKERB</u>						
N 022059	001 8821927	Sep 18, 2029	DS DP			
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861	001 10005783	Oct 21, 2029		U-2472	ODE-215	Nov 26, 2025
	10047097	Oct 21, 2029		U-2474	ODE-220	Nov 26, 2025
	10172861	Nov 16, 2035	DS DP		ODE-221	Nov 26, 2025
	10285993	Nov 16, 2035		U-2470		
	10774085	Oct 21, 2029		U-2470		
	10799505	Aug 15, 2036	DS DP			
	10813936	Nov 16, 2035		U-2987		
	8513263	Dec 23, 2029	DS DP			
	8865698	Oct 21, 2029		U-2469		
	9127013	Oct 21, 2029	DS DP			
	9447104	Oct 21, 2029		U-2470		
	9676783	Oct 21, 2029		U-2469		
	9782414	Nov 16, 2035		U-2475		
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 210861	002 10005783	Oct 21, 2029		U-2472	ODE-215	Nov 26, 2025
	10047097	Oct 21, 2029		U-2474	ODE-220	Nov 26, 2025
	10172861	Nov 16, 2035	DS DP		ODE-221	Nov 26, 2025
	10285993	Nov 16, 2035		U-2470		
	10774085	Oct 21, 2029		U-2470		
	10799505	Aug 15, 2036	DS DP			
	10813936	Nov 16, 2035		U-2987		
	8513263	Dec 23, 2029	DS DP			
	8865698	Oct 21, 2029		U-2469		
	9127013	Oct 21, 2029	DS DP			
	9447104	Oct 21, 2029		U-2470		
	9676783	Oct 21, 2029		U-2469		
	9782414	Nov 16, 2035		U-2475		
<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 211710	001 10005783	Oct 21, 2029		U-2472	ODE-215	Nov 26, 2025
	10045991	Apr 04, 2037		U-2473	ODE-220	Nov 26, 2025
	10047097	Oct 21, 2029		U-2474	ODE-221	Nov 26, 2025
	10137127	Apr 04, 2037	DP			
	10172861	Nov 16, 2035	DS			
	10668072	Apr 04, 2037	DP			
	10774085	Oct 21, 2029		U-2470		
	10799505	Aug 15, 2036	DS			
	11191766	Apr 04, 2037		U-2471		
	11484535	Apr 04, 2037		U-2470		
	8513263	Dec 23, 2029	DS DP			
	8865698	Oct 21, 2029		U-2469		
	9127013	Oct 21, 2029	DS DP			
	9447104	Oct 21, 2029		U-2470		
	9676783	Oct 21, 2029		U-2469		

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<u>LAROTRECTINIB SULFATE - VITRAKVI</u>						
N 211710	001 9782414	Nov 16, 2035	U-2471			
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280	001 11053214	Dec 05, 2037	DS DP U-1719		NCE	Jan 31, 2025
	7423050	Feb 17, 2028	DS DP U-1719			
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280	002 11053214	Dec 05, 2037	DS DP U-1719		NCE	Jan 31, 2025
	7423050	Feb 17, 2028	DS DP U-1719			
<u>LASMIDITAN SUCCINATE - REYVOW</u>						
N 211280	003 11053214	Dec 05, 2037	DS DP U-1719		NCE	Jan 31, 2025
	7423050	Feb 17, 2028	DS DP U-1719			
<u>LATANOPROST - XELPROS</u>						
N 206185	001 9539262	Oct 15, 2028	U-2400			
	9629852	Sep 12, 2029	DP			
<u>LATANOPROST; NETARSUDIL DIMESYLATE - ROCKLATAN</u>						
N 208259	001 10174017	Jan 27, 2030	DS DP U-1524			
	10532993	Jul 11, 2026	U-1524			
	10588901	Mar 14, 2034	DS DP U-1524			
	10654844	Jan 27, 2030	DS DP U-1524			
	10882840	Jul 11, 2026	U-1524			
	11021456	Jul 11, 2026	U-1524			
	11028081	Jan 27, 2030	U-1524			
	11185538	Mar 14, 2034	DP			
	11197853	Mar 14, 2034	DP			
	11618748	Jan 27, 2030	U-1524			
	8394826	Nov 10, 2030	DS DP U-1524			
	8450344	Jul 11, 2026	DS DP U-1524			
	9096569	Jul 11, 2026	DS DP U-1524			
	9415043	Mar 14, 2034	DS			
	9931336	Mar 14, 2034	DS DP U-1524			
	9993470	Mar 14, 2034	DS DP U-1524			
<u>LATANOPROSTENE BUNOD - VYZULTA</u>						
N 207795	001 7273946	Oct 03, 2025	DS DP U-2144			
	7629345	Jan 05, 2025	DP U-2144			
	7910767	Jan 05, 2025	DS DP U-2144			
	8058467	Jan 05, 2025	DS U-2144			
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834	001 10039779	Jan 30, 2034	DS DP U-2369		ODE*	Aug 28, 2026
	10039779	Jan 30, 2034	DS DP U-2370		ODE-136	Apr 07, 2024
	10039779*PED	Jul 30, 2034			PED	Oct 07, 2024
	10456414	Sep 14, 2032	DP			
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			

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<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834 001	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	DP U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834 002	10039779	Jan 30, 2034	DS DP U-1470		ODE*	Apr 07, 2024
	10039779*PED	Jul 30, 2034			ODE*	Aug 28, 2026
	10456414	Sep 14, 2032	DP		PED	Oct 07, 2024
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				

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<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 205834	002 9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477	001 10456414	Sep 14, 2032	DP		ODE-262	Aug 28, 2026
	7964580	Mar 26, 2029	DS DP U-1470		ODE-263	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-264	Aug 28, 2026
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477	002 10456414	Sep 14, 2032	DP		ODE-262	Aug 28, 2026
	7964580	Mar 26, 2029	DS DP U-1470		ODE-263	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-264	Aug 28, 2026
	8088368	May 12, 2030	DS DP			
	8088368*PED	Nov 12, 2030				
	8273341	May 12, 2030	U-1470			
	8273341*PED	Nov 12, 2030				
	8334270	Mar 21, 2028	DS DP U-1470			

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<u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u>						
N 212477 002	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8822430	May 12, 2030	DS DP U-1470			
	8822430*PED	Nov 12, 2030				
	8841278	May 12, 2030	DP U-1470			
	8841278*PED	Nov 12, 2030				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9393256	Sep 14, 2032	U-1470			
	9393256*PED	Mar 14, 2033				
	9511056	May 12, 2030	U-1470			
	9511056*PED	Nov 12, 2030				
<u>LEFAMULIN ACETATE - XENLETA</u>						
N 211672 001	8071643	Jan 16, 2029	DS DP		NCE	Aug 19, 2024
	8153689	Mar 19, 2028	DS DP		GAIN	Aug 19, 2029
	9120727	May 23, 2031	DS DP			
<u>LEFAMULIN ACETATE - XENLETA</u>						
N 211673 001	8071643	Jan 16, 2029	DS DP		NCE	Aug 19, 2024
	8153689	Mar 19, 2028	DS DP		GAIN	Aug 19, 2029
<u>LEMBorexant - DAYVIGO</u>						
N 212028 001	10188652	Oct 21, 2035	DP U-2791		M-293	Apr 20, 2026
	8268848	Sep 20, 2031	DS DP U-2791		NCE	Apr 07, 2025
<u>LEMBorexant - DAYVIGO</u>						
N 212028 002	10188652	Oct 21, 2035	DP U-2791		M-293	Apr 20, 2026
	8268848	Sep 20, 2031	DS DP U-2791		NCE	Apr 07, 2025
<u>LENACAPAVIR SODIUM - SUNLENCA</u>						
N 215973 001	10071985	Aug 17, 2037	DS DP		NCE	Dec 22, 2027
	10654827	Aug 17, 2037	U-3507			
	11267799	Aug 16, 2038	DS			
	9951043	Feb 28, 2034	DS DP U-3507			
<u>LENACAPAVIR SODIUM - SUNLENCA</u>						
N 215974 001	10071985	Aug 17, 2037	DS DP		NCE	Dec 22, 2027
	10654827	Aug 17, 2037	U-3507			
	11267799	Aug 16, 2038	DS			
	9951043	Feb 28, 2034	DS DP U-3507			

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<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 001	7465800	Apr 27, 2027	DS DP		ODE-131	Feb 22, 2024
	7855217	Nov 24, 2024	DS DP		ODE-241	May 28, 2026
	8741929	Mar 08, 2028		U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 002	7465800	Apr 27, 2027	DS DP		ODE-131	Feb 22, 2024
	7855217	Nov 24, 2024	DS DP		ODE-241	May 28, 2026
	8741929	Mar 08, 2028		U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 003	7465800	Apr 27, 2027	DS DP		ODE-131	Feb 22, 2024
	7855217	Nov 24, 2024	DS DP		ODE-241	May 28, 2026
	8741929	Mar 08, 2028		U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 004	7465800	Apr 27, 2027	DS DP		ODE-131	Feb 22, 2024
	7855217	Nov 24, 2024	DS DP		ODE-241	May 28, 2026
	8741929	Mar 08, 2028		U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 005	7465800	Apr 27, 2027	DS DP		ODE-131	Feb 22, 2024
	7855217	Nov 24, 2024	DS DP		ODE-241	May 28, 2026
	8741929	Mar 08, 2028		U-1983	ODE-245	May 28, 2026
<u>LENALIDOMIDE - REVLIMID</u>						
N 021880 006	7465800	Apr 27, 2027	DS DP		ODE-131	Feb 22, 2024
	7855217	Nov 24, 2024	DS DP		ODE-241	May 28, 2026
	8741929	Mar 08, 2028		U-1983	ODE-245	May 28, 2026
<u>LENIOLISIB PHOSPHATE - JOENJA</u>						
N 217759 001	8653092	Feb 19, 2032	DS DP		NCE	Mar 24, 2028
					ODE-430	Mar 24, 2030
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947 001	10259791	Aug 26, 2035	DS		I-868	Aug 10, 2024
	10407393	Aug 26, 2035	DS		M-269	Jul 21, 2024
	11090386	Feb 23, 2036		U-3519	M-272	Dec 19, 2024
	11186547	Aug 26, 2035	DS		ODE-196	Aug 15, 2025
	7253286	Oct 24, 2025	DS DP			
	7612208	Sep 19, 2026	DS DP			
	9006256	Jul 27, 2027		U-1695		
<u>LENVATINIB MESYLATE - LENVIMA</u>						
N 206947 002	10259791	Aug 26, 2035	DS		I-868	Aug 10, 2024
	10407393	Aug 26, 2035	DS		M-269	Jul 21, 2024
	11090386	Feb 23, 2036		U-3519	M-272	Dec 19, 2024
	11186547	Aug 26, 2035	DS		ODE-196	Aug 15, 2025
	7253286	Oct 24, 2025	DS DP			
	7612208	Sep 19, 2026	DS DP			
	9006256	Jul 27, 2027		U-1695		
<u>LESINURAD - ZURAMPIC</u>						
N 207988 001	10183012	Nov 26, 2028		U-2311		
	8003681	Aug 25, 2025	DS			
	8084483	Aug 17, 2029		U-1801		

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<u>LESINURAD - ZURAMPIC</u>						
N 207988	001	8283369	Nov 26, 2028	U-1802		
		8283369	Nov 26, 2028	U-1804		
		8357713	Dec 22, 2029	DP U-1801		
		8357713	Dec 22, 2029	DP U-1802		
		8357713	Dec 22, 2029	DP U-1803		
		8546436	Feb 29, 2032	DS DP		
		8546437	Apr 29, 2029	U-1803		
		9216179	Aug 01, 2031	U-1806		
		9956205	Dec 28, 2031	U-2311		
<u>LETERMOVIR - PREVYMIS</u>						
N 209939	001	8513255	May 22, 2024	DS DP	D-189	Aug 02, 2026
		RE46791	May 22, 2024	DS DP	I-916	Jun 05, 2026
					ODE-165	Nov 08, 2024
					ODE-423	Jun 05, 2030
<u>LETERMOVIR - PREVYMIS</u>						
N 209939	002	8513255	May 22, 2024	DS DP	D-189	Aug 02, 2026
		RE46791	May 22, 2024	DS DP	I-916	Jun 05, 2026
					ODE-165	Nov 08, 2024
					ODE-423	Jun 05, 2030
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	001	10603384	Feb 28, 2033	DP	D-189	Aug 02, 2026
		8513255	May 22, 2024	DS DP	I-916	Jun 05, 2026
		RE46791	May 22, 2024	DS DP	ODE-165	Nov 08, 2024
					ODE-423	Jun 05, 2030
<u>LETERMOVIR - PREVYMIS</u>						
N 209940	002	10603384	Feb 28, 2033	DP	D-189	Aug 02, 2026
		8513255	May 22, 2024	DS DP	I-916	Jun 05, 2026
		RE46791	May 22, 2024	DS DP	ODE-165	Nov 08, 2024
					ODE-423	Jun 05, 2030
<u>LETROZOLE; RIBOCICLIB SUCCINATE - KISQALI FEMARA CO-PACK (COPACKAGED)</u>						
N 209935	001	10799506	Apr 14, 2036	DP	NPP	Dec 10, 2024
		8324225	Jun 17, 2028	DS DP		
		8415355	Mar 13, 2031	DS DP		
		8685980	May 25, 2030	DS DP		
		8962630	Dec 09, 2029	U-2505		
		8962630	Dec 09, 2029	U-3264		
		9193732	Nov 09, 2031	DS DP		
		9416136	Aug 20, 2029	U-2505		
		9416136	Aug 20, 2029	U-3264		
		9868739	Nov 09, 2031	U-2505		
		9868739	Nov 09, 2031	U-3264		
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED KIT</u>						
N 020263	009	8921326	Feb 05, 2031	DP	NS	Apr 14, 2026
		9617303	Mar 22, 2028	U-3611		
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N 020517	003	8921326	Feb 05, 2031	DP	U-1666	

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<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021343	001 11771841	Dec 22, 2041	DS DP			
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021379	001 11771841	Dec 22, 2041	DS DP			
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021488	001 11771841	Dec 22, 2041	DS DP			
<u>LEUPROLIDE ACETATE - ELIGARD KIT</u>						
N 021731	001 11771841	Dec 22, 2041	DS DP			
<u>LEUPROLIDE ACETATE - FENSOLVI KIT</u>						
N 213150	001 11771841	Dec 22, 2041	DS DP			
<u>LEUPROLIDE MESYLATE - CAMCEVI KIT</u>						
N 211488	001 10646572	Jan 16, 2027	DP		NP	May 25, 2024
	11717555	Jan 01, 2039	DP			
	9572857	Jan 16, 2027	DP			
	9744207	Jan 16, 2027	DP			
<u>LEVALBUTEROL TARTRATE - XOPENEX HFA</u>						
N 021730	001 7256310	Oct 08, 2024	DS DP U-636			
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	001 8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	002 8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	003 8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<u>LEVETIRACETAM - KEPPRA</u>						
N 021035	004 8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<u>LEVETIRACETAM - KEPPRA XR</u>						
N 022285	001 7858122	Sep 17, 2028	DP			
<u>LEVETIRACETAM - KEPPRA XR</u>						
N 022285	002 7858122	Sep 17, 2028	DP			
<u>LEVETIRACETAM - ELEPSIA XR</u>						
N 204417	001 8163306	Sep 03, 2027	DP			
	8425938	Feb 22, 2026	DP			
	8431156	Oct 31, 2027	DP			
	8470367	Oct 31, 2027	DP			
	8535717	Feb 22, 2026	DP			
<u>LEVETIRACETAM - ELEPSIA XR</u>						
N 204417	002 8163306	Sep 03, 2027	DP			
	8425938	Feb 22, 2026	DP			
	8431156	Oct 31, 2027	DP			
	8470367	Oct 31, 2027	DP			
	8535717	Feb 22, 2026	DP			

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<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	001	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	002	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	003	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVETIRACETAM - SPRITAM</u>						
N 207958	004	11160786	Mar 14, 2034	DP		
		9339489	Mar 14, 2034	DP	U-1850	
		9669009	Mar 14, 2034		U-1850	
		9669009	Mar 14, 2034		U-2021	
		9669009	Mar 14, 2034		U-2022	
<u>LEVOCETIRIZINE DIHYDROCHLORIDE - XYZAL ALLERGY 24HR</u>						
N 209090	001	8633194	Oct 16, 2027	DP		
<u>LEVODOPA - INBRIJA</u>						
N 209184	001	7182961	Feb 22, 2024	DP		
		8545878	Nov 16, 2032	DP		
		8685442	Nov 16, 2032	DP		
		8945612	Nov 16, 2032	DP		
		9393210	Nov 16, 2032	DP		
		RE43711	Feb 03, 2029		U-2484	
<u>LEVOKETOCONAZOLE - RECORLEV</u>						
N 214133	001	10098877	Jan 10, 2026	U-3283	ODE-385	Dec 30, 2028
		10517868	Jan 10, 2026	U-3283		
		10835530	Jan 10, 2026	U-3283		
		11020393	Mar 02, 2040	U-3282		
		11278547	Mar 02, 2040	U-3282		
		11478471	Jan 10, 2026	U-3283		
		9918984	Jan 10, 2026	U-3283		
<u>LEVOLEUCOVORIN - KHAPZORY</u>						
N 211226	001	11541012	Mar 25, 2039	DP		
<u>LEVOLEUCOVORIN - KHAPZORY</u>						
N 211226	002	11541012	Mar 25, 2039	DP		

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<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	001	8481598	Mar 02, 2031		U-839	
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026		U-839	
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	002	8481598	Mar 02, 2031		U-839	
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026		U-839	
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	003	8481598	Mar 02, 2031		U-839	
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026		U-839	
<u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u>						
N 204168	004	8481598	Mar 02, 2031		U-839	
		8865937	May 23, 2032	DS DP		
		RE43879	Jan 11, 2026		U-839	
<u>LEVONORGESTREL - MIRENA</u>						
N 2021225	001	10561524	Sep 16, 2029		U-2948	
		10987244	Apr 01, 2031	DP		
		9615965	Sep 16, 2029	DP	U-2003	
		9668912	Apr 01, 2031	DP		
<u>LEVONORGESTREL - SKYLA</u>						
N 203159	001	10561524	Sep 16, 2029		U-2948	
		10987244	Apr 01, 2031	DP		
		11628088	Feb 07, 2027	DP		
		9615965	Sep 16, 2029	DP	U-2003	
		9668912	Apr 01, 2031	DP		
<u>LEVONORGESTREL - LILETTA</u>						
N 206229	001	10028858	Mar 22, 2034	DP	U-2348	I-917 Jun 29, 2026
		11090186	Oct 24, 2033		U-2348	
		11571328	Sep 07, 2040	DP		
<u>LEVONORGESTREL - KYLEENA</u>						
N 208224	001	10561524	Sep 16, 2029		U-2948	
		10987244	Apr 01, 2031	DP		
		11628088	Feb 07, 2027	DP		
		9615965	Sep 16, 2029	DP	U-2003	
		9668912	Apr 01, 2031	DP		
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	003	10231931	Mar 23, 2038		DP	
		10406108	Mar 23, 2038		DP	
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	004	10231931	Mar 23, 2038		DP	
		10406108	Mar 23, 2038		DP	
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	005	10231931	Mar 23, 2038		DP	
		10406108	Mar 23, 2038		DP	

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<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	006 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	007 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	008 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	009 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	010 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	011 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - LEVOLET</u>						
N 021137	012 10231931	Mar 23, 2038	DP			
	10406108	Mar 23, 2038	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	002 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	003 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	004 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	005 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	006 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	007 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	008 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	009 7691411	Mar 14, 2024	DP			
	7723390	Mar 14, 2024	DP			

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<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	010	7691411	Mar 14, 2024	DP		
		7723390	Mar 14, 2024	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	011	7691411	Mar 14, 2024	DP		
		7723390	Mar 14, 2024	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	012	7691411	Mar 14, 2024	DP		
		7723390	Mar 14, 2024	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N 021924	013	7691411	Mar 14, 2024	DP		
		7723390	Mar 14, 2024	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 202231	001	9006289	Oct 03, 2032	DP		
		9168238	Aug 29, 2032	DP		
		9168239	Aug 29, 2032	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 202231	002	9006289	Oct 03, 2032	DP		
		9168238	Aug 29, 2032	DP		
		9168239	Aug 29, 2032	DP		
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 202231	003	9006289	Oct 03, 2032	DP		
		9168238	Aug 29, 2032	DP		
		9168239	Aug 29, 2032	DP		
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	001	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	002	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	003	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	004	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	005	10537538	Feb 28, 2037	DP		

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<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	005	11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	006	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	007	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	008	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	009	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	010	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	011	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	012	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	013	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		
		11241382	Sep 17, 2039		U-3757	
		11241382	Sep 17, 2039		U-3758	
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	014	10537538	Feb 28, 2037	DP		
		11096913	Feb 28, 2037	DP		

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<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	014 11241382	Sep 17, 2039	U-3757			
	11241382	Sep 17, 2039	U-3758			
<u>LEVOTHYROXINE SODIUM - TIROSINT-SOL</u>						
N 206977	015 10537538	Feb 28, 2037	DP			
	11096913	Feb 28, 2037	DP			
	11241382	Sep 17, 2039	U-3757			
	11241382	Sep 17, 2039	U-3758			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	001 10398669	Dec 01, 2036	DP			
	11135190	Dec 01, 2036	DP			
	9782376	Dec 01, 2036	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	002 10398669	Dec 01, 2036	DP			
	11135190	Dec 01, 2036	DP			
	9782376	Dec 01, 2036	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 210632	003 10398669	Dec 01, 2036	DP			
	11135190	Dec 01, 2036	DP			
	9782376	Dec 01, 2036	DP			
<u>LEVOTHYROXINE SODIUM - THYQUIDITY</u>						
N 214047	001 9050307	Aug 06, 2031	DP			
<u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u>						
N 214253	001 11154498	Jul 20, 2036	DP			
<u>LEVOTHYROXINE SODIUM - ERMEZA</u>						
N 215809	001 9345772	Feb 27, 2035	DP			
<u>LIDOCAINE - ZTLIDO</u>						
N 207962	001 10765640	May 10, 2031	DP			
	10765749	May 10, 2031	DP			
	11278623	May 10, 2031	DP			
	11786455	May 10, 2031	DP	U-2267		
	11793766	May 10, 2031	DP	U-2267		
	9283174	May 10, 2031	DP			
	9925264	May 10, 2031	DP	U-2267		
	9931403	May 10, 2031	DP			
<u>LIDOCAINE HYDROCHLORIDE - ZINGO</u>						
N 022114	001 8540665	Oct 22, 2029	U-1438			
	9358338	Apr 27, 2035	U-1870			
	9370622	Sep 28, 2035	U-1870			
<u>LIDOCAINE HYDROCHLORIDE - AKTEN</u>						
N 022221	001 8759401	Jul 24, 2026	DP	U-1523		
<u>LIDOCAINE; TETRACAINE - PLIAGLIS</u>						
N 021717	001 10350180	Jan 14, 2031	DP			
	10603293	Jan 14, 2031	DP			
	10751305	Jan 14, 2031	DP			

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<u>LIFITEGRAST - XIIDRA</u>						
N 208073 001	10124000	Nov 05, 2024	U-1900			
	11058677	Dec 18, 2033	DP			
	7314938	Mar 10, 2025	DS DP			
	7745460	Nov 05, 2024	DS DP U-1880			
	7790743	Nov 05, 2024	U-1880			
	7928122	Nov 05, 2024	DS DP			
	8084047	May 17, 2026	DS DP			
	8168655	May 09, 2029	U-1880			
	8367701	Apr 15, 2029	DP U-1880			
	8592450	May 17, 2026	U-1880			
	8927574	Nov 12, 2030	DP			
	9085553	Jul 25, 2033	DP			
	9216174	Nov 05, 2024	DP			
	9353088	Oct 21, 2030	DP			
	9447077	Apr 15, 2029	U-1900			
	9890141	Oct 21, 2030	DS			
<u>LINACLOTIDE - LINZESS</u>						
N 202811 001	7304036	Aug 30, 2026	DS DP U-1278		I-921	Jun 12, 2026
	7304036	Aug 30, 2026	DS DP U-1516			
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024	U-1278			
	8748573	Oct 30, 2031	U-1515			
	8748573	Oct 30, 2031	U-1516			
	8802628	Oct 30, 2031	DP			
	8933030	Feb 17, 2031	DP			
	9708371	Aug 16, 2033	DP U-1515			
	9708371	Aug 16, 2033	DP U-1516			
<u>LINACLOTIDE - LINZESS</u>						
N 202811 002	7304036	Aug 30, 2026	DS DP U-1278			
	7304036	Aug 30, 2026	DS DP U-1516			
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024	U-1278			
	8748573	Oct 30, 2031	U-1515			
	8748573	Oct 30, 2031	U-1516			
	8802628	Oct 30, 2031	DP			
	8933030	Feb 17, 2031	DP			
	9708371	Aug 16, 2033	DP U-1515			
<u>LINACLOTIDE - LINZESS</u>						
N 202811 003	10675325	Aug 11, 2031	DP		I-921	Jun 12, 2026
	10702576	Aug 11, 2031	U-1516			
	10702576	Aug 11, 2031	U-3644			
	7304036	Aug 30, 2026	DS DP U-1516			

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<u>LINACLOTIDE - LINZESS</u>						
N 202811 003	7304036	Aug 30, 2026	DS DP U-3644			
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024		U-1516		
	8110553	Jan 28, 2024		U-3644		
	8933030	Feb 17, 2031	DP U-1516			
	8933030	Feb 17, 2031	DP U-3644			
	9708371	Aug 16, 2033	DP U-1516			
	9708371	Aug 16, 2033	DP U-3644			
<u>LINAGLIPTIN - TRADJENTA</u>						
N 201280 001	10034877	Aug 05, 2029		U-2347	M-295	Jun 20, 2026
	10034877*PED	Feb 05, 2030			PED	Dec 20, 2026
	11033552	May 04, 2027	DP			
	11033552*PED	Nov 04, 2027				
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030		U-1503	Y	
	8846695*PED	Dec 04, 2030				
	8853156	Mar 05, 2031	DP U-1642			
	8853156*PED	Sep 05, 2031				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9486526	Aug 05, 2029		U-1915		
	9486526*PED	Feb 05, 2030				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 001	10022379	Apr 02, 2029		U-2339	M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	10973827	Apr 02, 2029	DP			
	10973827*PED	Oct 02, 2029				
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030		U-1503		
	8846695*PED	Dec 04, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 002	10022379	Apr 02, 2029		U-2339	M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026

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<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 002	10973827	Apr 02, 2029	DP			
	10973827*PED	Oct 02, 2029				
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030		U-1503		
	8846695*PED	Dec 04, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u>						
N 201281 003	10022379	Apr 02, 2029		U-2339	M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	10973827	Apr 02, 2029	DP			
	10973827*PED	Oct 02, 2029				
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8846695	Jun 04, 2030		U-1503	Y	
	8846695*PED	Dec 04, 2030				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 001	10022379	Apr 02, 2029		U-2339	M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027		U-1503	Y	
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9173859	May 04, 2027	DP U-1503		Y	
	9173859*PED	Nov 04, 2027				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
	9555001*PED	Sep 06, 2033				

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<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 001	10022379	Apr 02, 2029	U-2339		M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027	U-1503	Y		
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9173859	May 04, 2027	DP U-1503	Y		
	9173859*PED	Nov 04, 2027				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
	9555001*PED	Sep 06, 2033				
<u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u>						
N 208026 002	10022379	Apr 02, 2029	U-2339		M-295	Jun 20, 2026
	10022379*PED	Oct 02, 2029			PED	Dec 20, 2026
	7407955	May 02, 2025	DS DP			
	7407955*PED	Nov 02, 2025				
	8673927	May 04, 2027	U-1503	Y		
	8673927*PED	Nov 04, 2027				
	8883805	Nov 26, 2025	DP			
	8883805*PED	May 26, 2026				
	9155705	May 21, 2030	DP			
	9155705*PED	Nov 21, 2030				
	9173859	May 04, 2027	DP U-1503	Y		
	9173859*PED	Nov 04, 2027				
	9415016	Apr 02, 2029	DP			
	9415016*PED	Oct 02, 2029				
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
	9555001*PED	Sep 06, 2033				
<u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u>						
N 022341 001	7762994	May 23, 2024	DP			
	7762994*PED	Nov 23, 2024				
	8114833	Aug 13, 2025	DS DP			
	8114833*PED	Feb 13, 2026				
	9265893	Sep 23, 2032	DP			
	9265893*PED	Mar 23, 2033				
	9968659	Jan 09, 2037	U-2313			
	9968659*PED	Jul 09, 2037				
<u>LIRAGLUTIDE RECOMBINANT - SAXENDA</u>						
N 206321 001	10220155	Jul 17, 2026	DP			
	10220155*PED	Jan 17, 2027				
	10357616	Jan 20, 2026	DP			

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<u>LIRAGLUTIDE RECOMBINANT - SAXENDA</u>						
N 206321 001	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	11311679	Jan 20, 2026	DP			
	11446443	Oct 20, 2025	DP			
	7762994	May 23, 2024	DP			
	7762994*PED	Nov 23, 2024				
	8114833	Aug 13, 2025	DP			
	8114833*PED	Feb 13, 2026				
	8684969	Oct 20, 2025	DP			
	8684969*PED	Apr 20, 2026				
	8920383	Jul 17, 2026	DP			
	8920383*PED	Jan 17, 2027				
	9108002	Jan 26, 2026	DP			
	9108002*PED	Jul 26, 2026				
	9132239	Feb 01, 2032	DP			
	9132239*PED	Aug 01, 2032				
	9457154	Sep 27, 2027	DP			
	9457154*PED	Mar 27, 2028				
	9616180	Jan 20, 2026	DP			
	9616180*PED	Jul 20, 2026				
	9687611	Feb 27, 2027	DP			
	9687611*PED	Aug 27, 2027				
	9775953	Jul 17, 2026	DP			
	9775953*PED	Jan 17, 2027				
	9861757	Jan 20, 2026	DP			
	9861757*PED	Jul 20, 2026				
	9968659	Jan 09, 2037		U-2438		
	9968659*PED	Jul 09, 2037				
	RE46363	Aug 03, 2026	DP			
	RE46363*PED	Feb 03, 2027				
<u>LISINAPRIL - OBRELIS</u>						
N 208401 001	10039800	Nov 06, 2035		U-1723		
	10039800	Nov 06, 2035		U-185		
	10039800	Nov 06, 2035		U-1864		
	10039800	Nov 06, 2035		U-1991		
	10039800	Nov 06, 2035		U-3		
	10039800	Nov 06, 2035		U-71		
	10039800	Nov 06, 2035		U-8		
	10265370	Nov 06, 2035	DP			
	10406199	Nov 06, 2035		U-1723		
	10406199	Nov 06, 2035		U-185		
	10406199	Nov 06, 2035		U-1864		
	10406199	Nov 06, 2035		U-1991		
	10406199	Nov 06, 2035		U-3		
	10406199	Nov 06, 2035		U-71		
	10406199	Nov 06, 2035		U-8		
	10940177	Nov 06, 2035	DP			
	11179434	Nov 06, 2035	DP			
	11771733	Nov 06, 2035	DP			

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<u>LISINOPRIL - OBRELIS</u>						
N 208401	001	9463183	Nov 06, 2035	DP		
		9616096	Nov 06, 2035	U-1723		
		9616096	Nov 06, 2035	U-185		
		9616096	Nov 06, 2035	U-1864		
		9616096	Nov 06, 2035	U-1991		
		9616096	Nov 06, 2035	U-3		
		9616096	Nov 06, 2035	U-71		
		9616096	Nov 06, 2035	U-8		
		9814751	Nov 06, 2035	DP		
<u>LITHIUM CITRATE - LITHIUM CITRATE</u>						
A 218036	001				CGT	Mar 18, 2024
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	001	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	002	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	003	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	004	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		

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<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	005	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LOMITAPIDE MESYLATE - JUXTAPID</u>						
N 203858	006	10016404	Mar 07, 2025	U-1316		
		10555938	Mar 07, 2025	U-1316		
		7932268	Aug 19, 2027	U-1316		
		8618135	Mar 07, 2025	U-1316		
		9265758	Mar 07, 2025	U-1316		
		9364470	Mar 07, 2025	U-1851		
		9433617	Mar 07, 2025	U-1316		
		9861622	Mar 07, 2025	U-1316		
<u>LONAFARNIB - ZOKINVY</u>						
N 213969	001	7838531	Jul 26, 2024	U-3070	NCE	Nov 20, 2025
		8828356	Oct 17, 2024	U-3070	ODE-324	Nov 20, 2027
<u>LONAFARNIB - ZOKINVY</u>						
N 213969	002	7838531	Jul 26, 2024	U-3070	NCE	Nov 20, 2025
		8828356	Oct 17, 2024	U-3070	ODE-324	Nov 20, 2027
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	001	8999393	Jan 08, 2034	DP U-3210		
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	002	8999393	Jan 08, 2034	DP U-3210		
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	003	8999393	Jan 08, 2034	DP U-3210		
<u>LORAZEPAM - LOREEV XR</u>						
N 214826	004	8999393	Jan 08, 2034	DP U-3210		
<u>LORLATINIB - LOREBENA</u>						
N 210868	001	10420749	Jul 27, 2036	DS DP U-2633	I-847	Mar 03, 2024
		10420749	Jul 27, 2036	DS DP U-3096	ODE-217	Nov 02, 2025
		11020376	Jul 27, 2036	DP	ODE-218	Nov 02, 2025
		11299500	Oct 04, 2038	DS	ODE-219	Nov 02, 2025
		8680111	Mar 05, 2033	DS DP	ODE-349	Mar 03, 2028
<u>LORLATINIB - LOREBENA</u>						
N 210868	002	10420749	Jul 27, 2036	DS DP U-2633	I-847	Mar 03, 2024
		10420749	Jul 27, 2036	DS DP U-3096	ODE-217	Nov 02, 2025
		11020376	Jul 27, 2036	DP	ODE-218	Nov 02, 2025
		11299500	Oct 04, 2038	DS	ODE-219	Nov 02, 2025
		8680111	Mar 05, 2033	DS DP	ODE-349	Mar 03, 2028
<u>LOTEPREDNOL ETABONATE - LOTEMAX SM</u>						
N 208219	001	10596107	Dec 23, 2036	DP U-2764		
		11534395	Jan 26, 2036	DP U-2764		

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<u>LOTEPREDNOL ETABONATE - LOTEMAX SM</u>						
N 208219	001 10596107	Dec 23, 2036	DP U-2764			
	11534395	Jan 26, 2036	DP U-2764			
<u>LOTEPREDNOL ETABONATE - INVELTYS</u>						
N 210565	001 10058511	May 03, 2033	DP U-2492			
	10646437	May 03, 2033	DP			
	10688045	May 03, 2033	DP			
	10864219	May 03, 2033	U-3011			
	11219597	May 03, 2033	U-3278			
	11219597	May 03, 2033	U-3279			
	11642317	May 03, 2033	DP			
	9056057	May 03, 2033	DP U-2491			
	9393213	May 03, 2033	DP			
	9532955	May 03, 2033	U-2491			
	9737491	May 03, 2033	U-2492			
	9827191	May 03, 2033	DP U-2493			
<u>LOTEPREDNOL ETABONATE - EYSUVIS</u>						
N 210933	001 10058511	May 03, 2033	DP U-2492			
	10646436	May 03, 2033	DP			
	10688045	May 03, 2033	DP			
	10857096	May 03, 2033	U-2985			
	10940108	May 03, 2033	U-2985			
	10945948	May 03, 2033	U-2985			
	10993908	May 03, 2033	U-3117			
	11219596	May 03, 2033	U-2985			
	11596599	May 03, 2033	U-2985			
	11642317	May 03, 2033	DP			
	9056057	May 03, 2033	DP U-2491			
	9393213	May 03, 2033	DP			
	9532955	May 03, 2033	U-2491			
	9737491	May 03, 2033	U-2492			
	9827191	May 03, 2033	DP U-2985			
<u>LOTILANER - XDEMVY</u>						
N 217603	001 10835517	Dec 14, 2038	U-3674		NCE	Jul 24, 2028
	11197847	Dec 14, 2038	U-3674			
	11690826	Dec 14, 2038	U-3674			
	11690827	Dec 14, 2038	U-3674			
	11752137	Dec 14, 2038	DP			
	8383659	Jan 17, 2030	DS DP			
<u>LOXAPINE - ADASUVE</u>						
N 022549	001 7458374	Aug 18, 2024	DP			
	7537009	Oct 28, 2024	DP			
	8387612	Oct 23, 2026	DP			
	8991387	May 21, 2024	DP			
	9370629	May 20, 2024	DP			
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	001 8026393	Oct 25, 2027	DP			
	8338639	Jan 23, 2027	DP			
	8748481	Sep 01, 2025	U-1520			

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<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	001 8779187	Jan 23, 2027	DP			
<u>LUBIPROSTONE - AMITIZA</u>						
N 021908	002 7795312	Sep 17, 2024	U-1085			
	8026393	Oct 25, 2027	DP			
	8338639	Jan 23, 2027	DP			
	8748481	Sep 01, 2025	U-1519			
	8779187	Jan 23, 2027	DP			
<u>LULICONAZOLE - LUZU</u>						
N 204153	001 8980931	Apr 28, 2034	DP			
	9012484	Sep 06, 2033	DS DP U-540			
	9199977	Sep 06, 2033	DS DP			
	9453006	Sep 06, 2033	DS			
<u>LUMASIRAN SODIUM - OXLUMO</u>						
N 214103	001 10131907	Aug 24, 2028	DS DP U-2995		I-901	Oct 06, 2025
	10435692	Dec 26, 2034	U-2995		NCE	Nov 23, 2025
	10465195	Dec 26, 2034	DS DP U-2995		ODE-339	Nov 23, 2027
	10478500	Oct 09, 2035	DS DP U-2995		ODE-415	Oct 06, 2029
	10487330	Dec 26, 2034	DS DP U-2995			
	10612024	Aug 14, 2035	DS DP U-2995			
	10612027	Aug 14, 2035	DS DP U-2995			
	11060093	Dec 26, 2034	DS DP U-2995			
	11261447	Nov 20, 2038	DS DP U-2995			
	11401517	Aug 14, 2035	DS DP U-2995			
	11446380	Oct 09, 2035	DS DP			
	8106022	Dec 12, 2029	DS DP U-2995			
	8828956	Dec 04, 2028	DS DP U-2995			
	9828606	Dec 26, 2034	DS DP			
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500	001 10117867	May 27, 2029	U-3271		I-882	Dec 17, 2024
	10464938	Mar 12, 2028	DP		NCE	Dec 20, 2024
	10695345	Aug 30, 2039	DP U-543			
	10960009	Dec 03, 2034	U-814			
	11026951	Dec 03, 2034	U-3274			
	11690842	Aug 30, 2039	DP U-3362			
	11690842	Aug 30, 2039	DP U-3363			
	11753419	Dec 10, 2040	DP			
	11806348	Aug 30, 2039	DP U-3362			
	11806348	Aug 30, 2039	DP U-3363			
	8598119	Dec 28, 2029	U-543			
	8648077	Dec 01, 2029	DS DP			
	9199995	Mar 12, 2029	U-2713			
	9586960	Mar 12, 2029	DS DP			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034	U-2714			
	RE48825	Mar 12, 2029	DS DP			
	RE48839	Aug 19, 2033	U-3271			
	RE48839	Aug 19, 2033	U-814			

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<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 002	10117867	May 27, 2029	U-3271		NCE	Dec 20, 2024
	10464938	Mar 12, 2028	DP			
	10695345	Aug 30, 2039	DP U-814			
	11026951	Dec 03, 2034	U-3364			
	11052084	Aug 30, 2039	DP U-3362			
	11052084	Aug 30, 2039	DP U-3363			
	11690842	Aug 30, 2039	DP U-3362			
	11690842	Aug 30, 2039	DP U-3363			
	11753419	Dec 10, 2040	DP			
	11806348	Aug 30, 2039	DP U-3362			
	11806348	Aug 30, 2039	DP U-3363			
	8648077	Dec 01, 2029	DS DP			
	9168258	May 27, 2029	DP			
	9199995	Mar 12, 2029	U-2713			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034	U-2714			
	RE48825	Feb 12, 2029	DS			
	RE48839	Aug 19, 2033	U-3271			
	RE48839	Aug 19, 2033	U-814			
<u>LUMATEPERONE TOSYLATE - CAPLYTA</u>						
N 209500 003	10117867	May 27, 2029	U-3271		NCE	Dec 20, 2024
	10464938	Mar 12, 2028	DP			
	10695345	Aug 30, 2039	DP U-814			
	11026951	Dec 03, 2034	U-3364			
	11052084	Aug 30, 2039	DP U-3362			
	11052084	Aug 30, 2039	DP U-3363			
	11690842	Aug 30, 2039	DP U-3362			
	11690842	Aug 30, 2039	DP U-3363			
	11753419	Dec 10, 2040	DP			
	11806348	Aug 30, 2039	DP U-3362			
	11806348	Aug 30, 2039	DP U-3363			
	8648077	Dec 01, 2029	DS DP			
	9199995	Mar 12, 2029	U-2713			
	9616061	May 27, 2029	DP			
	9956227	Dec 03, 2034	U-2714			
	RE48825	Feb 12, 2029	DS			
	RE48839	Aug 19, 2033	U-3271			
	RE48839	Aug 19, 2033	U-814			
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 001	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9174975	Feb 20, 2024	U-1770			
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			

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<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	001	9815827	Feb 20, 2024		U-543	
		9827242	May 23, 2031		U-2199	
		9827242	May 23, 2031		U-2201	
		9907794	May 26, 2026	DP		
		9907794*PED	Nov 26, 2026			
		RE45573	Jun 23, 2025	DS		
		RE45573*PED	Dec 23, 2025			
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	002	8729085	May 26, 2026	DP		
		8729085*PED	Nov 26, 2026			
		8883794	May 26, 2026	DP		
		8883794*PED	Nov 26, 2026			
		9174975	Feb 20, 2024		U-1770	
		9174975*PED	Aug 20, 2024			
		9259423	May 23, 2031		U-1822	
		9259423*PED	Nov 23, 2031			
		9555027	May 26, 2026	DP	U-543	
		9815827	Feb 20, 2024		U-2166	
		9815827	Feb 20, 2024		U-543	
		9827242	May 23, 2031		U-2199	
		9827242	May 23, 2031		U-2201	
		9907794	May 26, 2026	DP		
		9907794*PED	Nov 26, 2026			
		RE45573	Jun 23, 2025	DS		
		RE45573*PED	Dec 23, 2025			
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	003	8729085	May 26, 2026	DP		
		8729085*PED	Nov 26, 2026			
		8883794	May 26, 2026	DP		
		8883794*PED	Nov 26, 2026			
		9174975	Feb 20, 2024		U-1770	
		9174975*PED	Aug 20, 2024			
		9259423	May 23, 2031		U-1822	
		9259423*PED	Nov 23, 2031			
		9555027	May 26, 2026	DP	U-543	
		9815827	Feb 20, 2024		U-2166	
		9815827	Feb 20, 2024		U-543	
		9827242	May 23, 2031		U-2199	
		9827242	May 23, 2031		U-2201	
		9907794	May 26, 2026	DP		
		9907794*PED	Nov 26, 2026			
		RE45573	Jun 23, 2025	DS		
		RE45573*PED	Dec 23, 2025			
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603	004	8729085	May 26, 2026	DP		
		8729085*PED	Nov 26, 2026			
		8883794	May 26, 2026	DP		
		8883794*PED	Nov 26, 2026			

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<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 004	9174975	Feb 20, 2024	U-1770			
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N 200603 005	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9174975	Feb 20, 2024	U-1770			
	9174975*PED	Aug 20, 2024				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	9907794	May 26, 2026	DP			
	9907794*PED	Nov 26, 2026				
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<u>LURBINECTEDIN - ZEPZELCA</u>						
N 213702 001	7763615	Dec 13, 2029	DS DP U-2836		NCE ODE-304	Jun 15, 2025 Jun 15, 2027
<u>LUSUTROMBOPAG - MULPLETA</u>						
N 210923 001	7601746	Sep 05, 2024	DS DP U-2344			
	8530668	Jan 21, 2030	DS DP			
	8889722	Jul 29, 2028	DS DP			
	9427402	Sep 29, 2031	DP			
<u>LUTETIUM LU 177 DOTATATE - LUTATHERA</u>						
N 208700 001	10596276	Jul 25, 2038	DP		ODE-166	Jan 26, 2025
	10596278	Jul 25, 2038	DP			
<u>LUTETIUM LU-177 VIPIVOTIDE TETRAJETAN - PLUVICTO</u>						
N 215833 001	10398791	Oct 17, 2034	DS DP		NCE	Mar 23, 2027
	10406240	Aug 15, 2028	DS DP U-3345			
	11318121	Aug 15, 2028	DS DP U-3345			

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<u>MACIMORELIN ACETATE - MACRILEN</u>						
N 205598	001 8192719	Oct 12, 2027	U-2220		ODE-170	Dec 20, 2024
<u>MACITENTAN - OPSUMIT</u>						
N 204410	001 10946015	Sep 11, 2026	DP U-1445			
	7094781	Dec 05, 2025	DS DP			
	8268847	Apr 18, 2029	U-1446			
	8367685	Oct 04, 2028	DP U-1445			
	9265762	May 29, 2027	DP U-1820			
<u>MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM SULFATE - SUFLAVE</u>						
N 215344	001				NP	Jun 15, 2026
<u>MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM SULFATE - SUTAB</u>						
N 213135	001 10143656	Aug 04, 2037	DP			
	11033498	Aug 04, 2037	U-3164			
	11382864	Aug 04, 2037	U-3164			
	11638697	Aug 04, 2037	DP			
<u>MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE - SUPREP BOWEL PREP KIT</u>						
N 022372	001				ODE-315	Aug 05, 2027
<u>MALATHION - OVIDE</u>						
N 018613	001 7560445	Feb 01, 2027	DS DP U-986			
	7977324	Aug 14, 2026	DP			
<u>MANNITOL - BRONCHITOL</u>						
N 202049	001				ODE-327	Oct 30, 2027
<u>MARALIXIBAT CHLORIDE - LIVMARLI</u>						
N 214662	001 11229647	Feb 12, 2040	U-3290		NCE	Sep 29, 2026
	11260053	May 26, 2031	U-3290		NPP	Mar 13, 2026
	11376251	Oct 26, 2032	U-3290		ODE-379	Sep 29, 2028
	11497745	Feb 12, 2040	U-3290		ODE-429	Mar 13, 2030
<u>MARAVIROC - SELZENTRY</u>						
N 208984	001				NPP	Oct 30, 2023
					PED	Apr 30, 2024
<u>MARIBAVIR - LIVTENCITY</u>						
N 215596	001 11684632	Jan 04, 2032	U-3650		NCE	Nov 23, 2026
					ODE-388	Nov 23, 2028
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	001 9181200	Jun 19, 2034	DS DP		M-297	Jun 15, 2026
	9585883	Jun 19, 2034	U-3373		NCE	Apr 28, 2027
					ODE-398	Apr 28, 2029
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	002 9181200	Jun 19, 2034	DS DP		M-297	Jun 15, 2026
	9585883	Jun 19, 2034	U-3373		NCE	Apr 28, 2027
					ODE-398	Apr 28, 2029
<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	003 9181200	Jun 19, 2034	DS DP		M-297	Jun 15, 2026
	9585883	Jun 19, 2034	U-3373		NCE	Apr 28, 2027
					ODE-398	Apr 28, 2029

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<u>MAVACAMTEN - CAMZYOS</u>						
N 214998	004 9181200	Jun 19, 2034	DS DP		M-297	Jun 15, 2026
	9585883	Jun 19, 2034		U-3373	NCE	Apr 28, 2027
					ODE-398	Apr 28, 2029
<u>MECHLORETHAMINE HYDROCHLORIDE - VALCHLOR</u>						
N 202317	001 7838564	Mar 07, 2026	DP			
	7872050	Jul 08, 2029		U-1427		
	8450375	Mar 07, 2026	DP			
	8501818	Mar 07, 2026	DP			
	8501819	Mar 07, 2026		U-1427		
	9382191	Mar 07, 2026	DP			
<u>MEGESTROL ACETATE - MEGACE ES</u>						
N 021778	001 7101576	Apr 22, 2024		U-755		
	9040088	Apr 22, 2024		U-755		
	9101540	Apr 22, 2024	DP	U-755		
	9101549	Apr 22, 2024		U-755		
	9107827	Apr 22, 2024		U-755		
<u>MELOXICAM - VIVLODEX</u>						
N 207233	001 9526734	Mar 31, 2033	DP			
	9649318	Mar 31, 2035	DP			
	9808468	Mar 31, 2035		U-2160		
	9808468	Mar 31, 2035		U-2165		
<u>MELOXICAM - VIVLODEX</u>						
N 207233	002 9526734	Mar 31, 2033	DP			
	9649318	Mar 31, 2035	DP			
	9808468	Mar 31, 2035		U-2160		
	9808468	Mar 31, 2035		U-2165		
<u>MELOXICAM - ANJESO</u>						
N 210583	001 10463673	Feb 24, 2024	DP	U-2750		
	10471067	Feb 24, 2024	DP	U-2750		
	10709713	May 26, 2030		U-2750		
	10881663	Mar 08, 2039		U-3038		
	11253478	May 26, 2030	DP	U-3318		
	11458145	Mar 08, 2039		U-3318		
	9974746	May 26, 2030	DP			
<u>MELOXICAM - OMIIZ ODT</u>						
N 211210	001 8545879	Aug 31, 2030	DP			
<u>MELOXICAM - OMIIZ ODT</u>						
N 211210	002 8545879	Aug 31, 2030	DP			
<u>MELPHALAN FLUFENAMIDE HYDROCHLORIDE - PEPAXTO</u>						
N 214383	001 10285946	Apr 25, 2032	DP		NCE	Feb 26, 2026
	10322182	Apr 25, 2032	DP		ODE-348	Feb 26, 2028
	10543274	Apr 25, 2032		U-3093		
	10869928	Apr 25, 2032	DP			
	11344622	Apr 25, 2032		U-3093		
	6992207	Jun 25, 2024	DS DP	U-3093		

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<u>MELPHALAN HYDROCHLORIDE - HEPZATO</u>						
N 201848	001	10098997	Nov 07, 2032	DP	NP	Aug 14, 2026
		10195334	Jan 16, 2033	DP	ODE-438	Aug 14, 2030
		10369264	Nov 07, 2032	DP		
		10569004	Nov 07, 2032	U-3680		
		10569004	Nov 07, 2032	U-3683		
		11083831	Dec 30, 2032	DP		
		11241522	Nov 07, 2032	DP		
		11633528	Nov 07, 2032	U-3675		
		11833286	Dec 30, 2032	DP		
		9314561	Feb 07, 2034	DP		
		9707331	Sep 17, 2034	DP		
<u>MELPHALAN HYDROCHLORIDE - EVOMELA</u>						
N 207155	001	10040872	Jan 30, 2034	DP		
		10864183	May 28, 2030	DP		
		10940128	Jun 14, 2030	DP	U-3086	
		11020363	May 28, 2030	DP		
		8410077	Mar 13, 2029	DP		
		9200088	Mar 13, 2029	DP		
		9493582	Feb 27, 2033	DP		
<u>MELPHALAN HYDROCHLORIDE - MELPHALAN HYDROCHLORIDE</u>						
N 217110	001	10537520	Jun 29, 2036	DP		
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	001	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	002	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	003	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N 022525	004	8039009	Mar 24, 2029	U-539		
		8039009*PED	Sep 24, 2029			
<u>MENTHOL; METHYL SALICYLATE - SALONPAS</u>						
N 022029	001	8809615	Jan 03, 2030	DP		
		9233184	Aug 01, 2027	DP		
<u>MENTHOL; METHYL SALICYLATE - SALONPAS</u>						
N 022029	002	8809615	Jan 03, 2030	DP		
		9233184	Aug 01, 2027	DP		
<u>MEROPENEM; VABORBACTAM - VABOMERE</u>						
N 209776	001	10172874	Aug 08, 2031	DP	NCE	Aug 29, 2022
		10183034	Aug 08, 2031	U-2490	GAIN	Aug 29, 2027
		10561675	Aug 08, 2031	U-2490		
		11007206	Aug 08, 2031	U-3128		
		11376237	Apr 06, 2039	U-3421		
		8680136	Aug 29, 2031	DS DP		

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<u>MEROPENEM; VABORBACTAM - VABOMERE</u>						
N 209776	001 9694025	Aug 08, 2031			U-2120	
<u>MESALAMINE - SFROWASA</u>						
N 019618	002 7645801	Jul 24, 2027	DS DP			
<u>MESALAMINE - CANASA</u>						
N 021252	002 8217083	Jun 06, 2028			DP	
	8436051	Jun 06, 2028			DP	
<u>MESALAMINE - APRISO</u>						
N 022301	001 8865688	May 01, 2030			U-1310	
<u>METAXALONE - SKELAXIN</u>						
N 013217	003 7122566	Feb 06, 2026			U-915	
<u>METFORMIN HYDROCHLORIDE - GLUMETZA</u>						
N 021748	002 7780987	Mar 23, 2025	DS DP			
<u>METFORMIN HYDROCHLORIDE - RIOMET ER</u>						
N 212595	001 9962336	May 01, 2035			DP	
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u>						
N 021842	001 9101660	Jan 22, 2027			DP	
	9320714	Feb 03, 2029			DP	
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u>						
N 021842	002 9101660	Jan 22, 2027			DP	
	9320714	Feb 03, 2029			DP	
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
N 022024	001 7785627	Jul 31, 2026			DP	
	7959946	Jul 31, 2026			DP	
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
N 022024	002 7785627	Jul 31, 2026			DP	
	7959946	Jul 31, 2026			DP	
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	001 8628799	Jul 13, 2025			DP	
	9339472	Jul 13, 2025			DP	
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	002 8628799	Jul 13, 2025			DP	
	9339472	Jul 13, 2025			DP	
<u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u>						
N 200678	003 8628799	Jul 13, 2025			DP	
	9339472	Jul 13, 2025			DP	
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	001 7326708	Nov 24, 2026	DS DP		U-802	M-187
	7326708*PED	May 24, 2027				PED
	8414921	Jul 21, 2028			DP U-1036	
	8414921*PED	Jan 21, 2029				
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	002 7326708	Nov 24, 2026	DS DP		U-802	M-187
	7326708*PED	May 24, 2027				PED
	8414921	Jul 21, 2028			DP U-1036	

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<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>						
N 022044	002	8414921*PED	Jan 21, 2029			
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	001	7326708	Nov 24, 2026	DS DP U-1227	M-187	Dec 04, 2023
		7326708*PED	May 24, 2027		PED	Jun 04, 2024
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	002	7326708	Nov 24, 2026	DS DP U-1227	M-187	Dec 04, 2023
		7326708*PED	May 24, 2027		PED	Jun 04, 2024
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u>						
N 202270	003	7326708	Nov 24, 2026	DS DP U-1227		
		7326708*PED	May 24, 2027			
<u>METHOTREXATE - OTREXUP</u>						
N 204824	001	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	002	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	003	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		

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<u>METHOTREXATE - OTREXUP</u>						
N 204824	003	8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP	U-1442	
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP	U-1442	
		9421333	Mar 19, 2030	DP	U-1442	
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	004	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP	U-1442	
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP	U-1442	
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP	U-1442	
		9421333	Mar 19, 2030	DP	U-1442	
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	005	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP	U-1442	
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP	U-1442	
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP	U-1442	
		9421333	Mar 19, 2030	DP	U-1442	
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	006	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP	U-1442	
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP	U-1442	
		8814834	May 27, 2031	DP		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>METHOTREXATE - OTREXUP</u>						
N 204824	006	8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	007	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - OTREXUP</u>						
N 204824	008	10709844	Mar 10, 2029	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11684723	Mar 10, 2029	DP		
		8021335	Oct 04, 2026	DP		
		8480631	Mar 19, 2030	DP U-1442		
		8562564	Jan 24, 2026	DP		
		8579865	Mar 19, 2030	DP U-1442		
		8814834	May 27, 2031	DP		
		8945063	Mar 19, 2030	DP U-1442		
		9421333	Mar 19, 2030	DP U-1442		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9867949	Mar 10, 2029	DP		
<u>METHOTREXATE - RASUVO</u>						
N 205776	001	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	002	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	003	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	004	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	005	8664231	Jun 01, 2029	U-1442		

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>METHOTREXATE - RASUVO</u>						
N 205776	006	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	007	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	008	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	009	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - RASUVO</u>						
N 205776	010	8664231	Jun 01, 2029	U-1442		
<u>METHOTREXATE - JYLAMVO</u>						
N 212479	001	11129833	Oct 28, 2035	DP		
		11771701	Oct 29, 2034	DP	U-3700	
		11771701	Oct 29, 2034	DP	U-3701	
		11771701	Oct 29, 2034	DP	U-3702	
		11771701	Oct 29, 2034	DP	U-3703	
		11771701	Oct 29, 2034	DP	U-3704	
<u>METHOTREXATE SODIUM - XATMEP</u>						
N 208400	001	10231927	Jan 02, 2033	U-1349	ODE-137	Apr 25, 2024
		10231927	Jan 02, 2033	U-1699	ODE-138	Apr 25, 2024
		10610485	Jan 02, 2033	DP		
		11116724	Jan 02, 2033	U-1349		
		11116724	Jan 02, 2033	U-1699		
		9259427	Jan 02, 2033	DP		
		9855215	Jan 02, 2033	DP		
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	001	10376584	Apr 08, 2024	DP	U-1185	
		8247425	Dec 31, 2030	U-1185		
		8420663	Sep 30, 2029	U-1185		
		8552025	Apr 08, 2024	DP		
		8822490	Sep 30, 2029	DP	U-1185	
		9180125	Sep 30, 2029	DP	U-1185	
		9492445	Sep 30, 2029	DP	U-1185	
		9669096	Apr 08, 2024	DP		
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	002	10376584	Apr 08, 2024	DP	U-1185	
		8247425	Dec 31, 2030	U-1185		
		8420663	Sep 30, 2029	U-1185		
		8552025	Apr 08, 2024	DP		
		8822490	Sep 30, 2029	DP	U-1185	
		9180125	Sep 30, 2029	DP	U-1185	
		9492445	Sep 30, 2029	DP	U-1185	
		9669096	Apr 08, 2024	DP		
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	003	10376584	Apr 08, 2024	DP	U-1185	
		8247425	Dec 31, 2030	U-1185		
		8420663	Sep 30, 2029	U-1185		

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<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 021964	003 8552025	Apr 08, 2024	DP			
	8822490	Sep 30, 2029	DP U-1185			
	9180125	Sep 30, 2029	DP U-1185			
	9492445	Sep 30, 2029	DP U-1185			
	9669096	Apr 08, 2024	DP			
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>						
N 208271	001 10307417	Mar 10, 2031	DP			
	10376505	Mar 10, 2031	DP			
	8420663	Sep 30, 2029	U-1185			
	8524276	Mar 10, 2031	DP			
	8956651	Mar 10, 2031	DP			
	9180125	Sep 30, 2029	DP U-1185			
	9314461	Mar 10, 2031	DP			
	9492445	Sep 30, 2029	DP U-1185			
	9724343	Sep 30, 2029	DP U-1185			
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	001 8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9668981	Oct 07, 2025	U-2024			
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	002 8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9668981	Oct 07, 2025	U-2024			
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	003 8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9668981	Oct 07, 2025	U-2024			
<u>METHYLPHENIDATE - DAYTRANA</u>						
N 021514	004 8632802	Oct 07, 2025	DP			
	9034370	Oct 07, 2025	DP			
	9668981	Oct 07, 2025	U-2024			
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	001 11166947	Jan 25, 2038	U-3299			
	8840924	Jun 05, 2026	DP			
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	002 11166947	Jan 25, 2038	U-3299			
	8840924	Jun 05, 2026	DP			
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<u>METHYLPHENIDATE - COTEMPLA XR-ODT</u>						
N 205489	003 11166947	Jan 25, 2038	U-3299			
	8840924	Jun 05, 2026	DP			
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			

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<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u>						
N 021419	001	7691880	Oct 07, 2024	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u>						
N 021419	002	7691880	Oct 07, 2024	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLIVANT XR</u>						
N 202100	001	8062667	Mar 29, 2029	DP		
		8287903	Feb 15, 2031	DP		
		8465765	Feb 15, 2031	DP	U-1415	
		8563033	Feb 15, 2031	DP	U-1415	
		8778390	Feb 15, 2031	DP	U-1543	
		8956649	Feb 15, 2031	DP	U-1665	
		9040083	Feb 15, 2031	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960	001	10857143	Aug 14, 2033	DP	U-2993	
		11103494	Aug 14, 2033	DP		
		11103495	Aug 14, 2033	DP	U-2993	
		11633389	Aug 14, 2033	DP		
		8202537	Mar 15, 2027	DP		
		8287903	Feb 15, 2031	DP		
		8999386	Aug 14, 2033	DP		
		9295642	Aug 14, 2033	DP	U-1827	
		9545399	Aug 14, 2033	DP	U-1827	
		9844544	Aug 14, 2033	DP	U-2203	
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960	002	10857143	Aug 14, 2033	DP	U-2993	
		11103494	Aug 14, 2033	DP		
		11103495	Aug 14, 2033	DP	U-2993	
		11633389	Aug 14, 2033	DP		
		8202537	Mar 15, 2027	DP		
		8287903	Feb 15, 2031	DP		
		8999386	Aug 14, 2033	DP		
		9295642	Aug 14, 2033	DP	U-1827	
		9545399	Aug 14, 2033	DP	U-1827	
		9844544	Aug 14, 2033	DP	U-2203	
<u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u>						
N 207960	003	10857143	Aug 14, 2033	DP	U-2993	
		11103494	Aug 14, 2033	DP		
		11103495	Aug 14, 2033	DP	U-2993	
		11633389	Aug 14, 2033	DP		
		8202537	Mar 15, 2027	DP		
		8287903	Feb 15, 2031	DP		
		8999386	Aug 14, 2033	DP		
		9295642	Aug 14, 2033	DP	U-1827	
		9545399	Aug 14, 2033	DP	U-1827	
		9844544	Aug 14, 2033	DP	U-2203	
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311	001	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	

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<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311	001	10617651	Mar 23, 2032		U-2357	
		10881618	Mar 23, 2032		U-2357	
		10905652	Mar 23, 2032	DP		
		11241391	Mar 23, 2032		U-2357	
		11241392	Mar 23, 2032	DP		
		8916588	Mar 23, 2032		U-2357	
		8927010	Mar 23, 2032	DP		
		9023389	Mar 23, 2032	DP		
		9028868	Mar 23, 2032		U-2357	
		9034902	Mar 23, 2032		U-2357	
		9283214	Mar 23, 2032	DP		
		9498447	Mar 23, 2032	DP		
		9603809	Mar 23, 2032		U-2357	
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311	002	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	
		10617651	Mar 23, 2032		U-2357	
		10881618	Mar 23, 2032		U-2357	
		10905652	Mar 23, 2032	DP		
		11241391	Mar 23, 2032		U-2357	
		11241392	Mar 23, 2032	DP		
		8916588	Mar 23, 2032		U-2357	
		8927010	Mar 23, 2032	DP		
		9023389	Mar 23, 2032	DP		
		9028868	Mar 23, 2032		U-2357	
		9034902	Mar 23, 2032		U-2357	
		9283214	Mar 23, 2032	DP		
		9498447	Mar 23, 2032	DP		
		9603809	Mar 23, 2032		U-2357	
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311	003	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	
		10617651	Mar 23, 2032		U-2357	
		10881618	Mar 23, 2032		U-2357	
		10905652	Mar 23, 2032	DP		
		11241391	Mar 23, 2032		U-2357	
		11241392	Mar 23, 2032	DP		
		8916588	Mar 23, 2032		U-2357	
		8927010	Mar 23, 2032	DP		
		9023389	Mar 23, 2032	DP		
		9028868	Mar 23, 2032		U-2357	
		9034902	Mar 23, 2032		U-2357	
		9283214	Mar 23, 2032	DP		
		9498447	Mar 23, 2032	DP		
		9603809	Mar 23, 2032		U-2357	
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311	004	10182995	Mar 23, 2032	DP		
		10292937	Mar 23, 2032		U-2357	

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<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 004	10617651	Mar 23, 2032	U-2357			
	10881618	Mar 23, 2032	U-2357			
	10905652	Mar 23, 2032	DP			
	11241391	Mar 23, 2032	U-2357			
	11241392	Mar 23, 2032	DP			
	8916588	Mar 23, 2032	U-2357			
	8927010	Mar 23, 2032	DP			
	9023389	Mar 23, 2032	DP			
	9028868	Mar 23, 2032	U-2357			
	9034902	Mar 23, 2032	U-2357			
	9283214	Mar 23, 2032	DP			
	9498447	Mar 23, 2032	DP			
	9603809	Mar 23, 2032	U-2357			
<u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u>						
N 209311 005	10182995	Mar 23, 2032	DP			
	10292937	Mar 23, 2032	U-2357			
	10617651	Mar 23, 2032	U-2357			
	10881618	Mar 23, 2032	U-2357			
	10905652	Mar 23, 2032	DP			
	11241391	Mar 23, 2032	U-2357			
	11241392	Mar 23, 2032	DP			
	8916588	Mar 23, 2032	U-2357			
	8927010	Mar 23, 2032	DP			
	9023389	Mar 23, 2032	DP			
	9028868	Mar 23, 2032	U-2357			
	9034902	Mar 23, 2032	U-2357			
	9283214	Mar 23, 2032	DP			
	9498447	Mar 23, 2032	DP			
	9603809	Mar 23, 2032	U-2357			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 001	10111839	Oct 30, 2035	U-2357		M-82	Jun 28, 2024
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP U-2357			
	10449159	Oct 30, 2035	U-2357			
	10500162	Oct 30, 2035	U-2357			
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035	U-2357			
	10568841	Oct 30, 2035	DP U-2357			
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038	U-2357			
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 002	10111839	Oct 30, 2035	U-2357		M-82	Jun 28, 2024
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP U-2357			
	10449159	Oct 30, 2035	U-2357			
	10500162	Oct 30, 2035	U-2357			

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<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 002	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 003	10111839	Oct 30, 2035		U-2357	M-82	Jun 28, 2024
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 004	10111839	Oct 30, 2035		U-2357	M-82	Jun 28, 2024
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038 005	10111839	Oct 30, 2035		U-2357	M-82	Jun 28, 2024
	10292938	Oct 30, 2035	DP			
	10292939	Oct 30, 2035	DP	U-2357		
	10449159	Oct 30, 2035		U-2357		
	10500162	Oct 30, 2035		U-2357		
	10507186	Oct 30, 2035	DP			
	10512612	Oct 30, 2035	DP			
	10512613	Oct 30, 2035		U-2357		
	10568841	Oct 30, 2035	DP	U-2357		
	10688060	Oct 30, 2035	DP			
	10722473	Nov 19, 2038		U-2357		
	9974752	Oct 30, 2035	DP			

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<u>METHYLPHENIDATE HYDROCHLORIDE - ADHANSIA XR</u>						
N 212038	006	10111839	Oct 30, 2035	U-2357	M-82	Jun 28, 2024
		10292938	Oct 30, 2035	DP		
		10292939	Oct 30, 2035	DP U-2357		
		10449159	Oct 30, 2035	U-2357		
		10500162	Oct 30, 2035	U-2357		
		10507186	Oct 30, 2035	DP		
		10512612	Oct 30, 2035	DP		
		10512613	Oct 30, 2035	U-2357		
		10568841	Oct 30, 2035	DP U-2357		
		10688060	Oct 30, 2035	DP		
		10722473	Nov 19, 2038	U-2357		
		9974752	Oct 30, 2035	DP		
<u>METOCLOPRAMIDE HYDROCHLORIDE - GIMOTI</u>						
N 209388	001	11020361	Dec 22, 2029	U-2843		
		11628150	Dec 22, 2029	DP U-2843		
		11813231	Dec 22, 2029	DP U-2843		
		8334281	May 16, 2030	DP U-2843		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	001	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	002	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	003	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u>						
N 210428	004	9504655	Jul 09, 2035	DP		
		9700530	Jul 09, 2035	DP		
<u>METRONIDAZOLE - VANDAZOLE</u>						
N 021806	001	7456207	Sep 22, 2024	DP		
<u>METRONIDAZOLE - NUVESSA</u>						
N 205223	001	10238634	Jun 28, 2032	DP		
		10596155	Jun 28, 2032	DP		
		7893097	Feb 19, 2028	DP		
		8658678	Jun 27, 2028	U-1682		
		8877792	Feb 02, 2028	DP		
		8946276	Jun 28, 2032	U-1664		
		9198858	Jun 28, 2032	U-1664		
<u>METRONIDAZOLE - LIKMEZ</u>						
N 216755	001	11541035	Oct 04, 2039	DP		
<u>MICONAZOLE NITRATE; WHITE PETROLATUM; ZINC OXIDE - VUSION</u>						
N 021026	001	8147852	Mar 30, 2028	U-1426		
<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
A 216159	001				PC	Jan 02, 2024

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<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
A 216159	002				PC	Jan 02, 2024
<u>MIDAZOLAM - NAYZILAM</u>						
N 211321	001	8217033	Jan 18, 2028	DP U-2526	ODE-243	May 17, 2026
		8809322	Jan 18, 2028	DP		
		9289432	Jan 18, 2028	DP U-2526		
		9687495	Jan 18, 2028	DP U-2526		
<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
N 211844	001	10966990	Jun 20, 2038	DP		
<u>MIDAZOLAM - MIDAZOLAM IN 0.9% SODIUM CHLORIDE</u>						
N 211844	002	10966990	Jun 20, 2038	DP		
<u>MIDAZOLAM HYDROCHLORIDE - SEIZALAM</u>						
N 209566	001				ODE-207	Sep 14, 2025
<u>MIDOSTAURIN - RYDAPT</u>						
N 207997	001	7973031	Oct 09, 2028	U-2007	ODE-140	Apr 28, 2024
		8575146	Dec 02, 2030	U-2008	ODE-141	Apr 28, 2024
<u>MIFEPRISTONE - KORLYM</u>						
N 202107	001	10006924	Aug 12, 2036	U-1643		
		10151763	Jan 18, 2037	U-1643		
		10166242	Apr 20, 2036	U-1643		
		10166243	Apr 20, 2036	U-1643		
		10195214	Jun 19, 2037	U-1643		
		10231983	Aug 22, 2038	U-1643		
		10314850	Aug 22, 2038	U-1643		
		10495650	Aug 12, 2036	U-1643		
		10500216	Mar 05, 2033	U-1643		
		10660904	Apr 20, 2036	U-1643		
		10780097	Aug 22, 2038	U-1643		
		10842800	Jun 19, 2037	U-1643		
		10842801	Nov 15, 2032	U-1643		
		8921348	Aug 27, 2028	U-1643		
		9829495	Aug 15, 2036	U-1643		
		9943526	Apr 20, 2036	U-1643		
<u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u>						
N 208623	001	10076514	Mar 15, 2037	U-2371	ODE-205	Aug 10, 2025
		10251873	May 30, 2038	U-2371		
		10383864	May 16, 2027	U-2371		
		10406143	May 16, 2027	U-2371		
		10471053	May 30, 2038	U-2371		
		10525045	Apr 28, 2028	U-2371		
		10792278	May 30, 2038	U-2371		
		10792279	May 30, 2038	U-2371		
		10799491	May 30, 2038	U-2371		
		10806727	May 30, 2038	U-2371		
		10813921	Feb 12, 2029	U-2371		
		10849889	May 30, 2038	U-2371		
		10849890	May 30, 2038	U-2371		
		10857141	May 30, 2038	U-2371		

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<u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u>						
N 208623 001	10857142	May 30, 2038				
	10874655	May 30, 2038				
	10874656	May 30, 2038				
	10874657	May 30, 2038				
	10925866	Apr 28, 2028				
	11033538	Apr 28, 2028				
	11234972	Mar 15, 2037				
	11241422	May 16, 2027				
	11278536	May 30, 2038				
	11278537	May 30, 2038				
	11278538	May 30, 2038				
	11278539	May 30, 2038				
	11278540	May 30, 2038				
	11304940	May 30, 2038	DS			
	11357761	May 30, 2038				
	11357762	May 30, 2038				
	11357763	May 30, 2038				
	11357764	May 30, 2038	DS			
	11357765	May 30, 2038	DS			
	11357784	Feb 06, 2039				
	11376244	May 30, 2038	DS			
	11389436	May 30, 2038				
	11389437	May 30, 2038				
	11426396	May 30, 2038	DS			
	11458128	May 30, 2038				
	11612593	May 30, 2038	DS			
	11612594	May 30, 2038	DS			
	11622962	Mar 17, 2039	DS			
	11633387	May 30, 2038	DS			
	11633388	Mar 25, 2039				
	11642334	Feb 20, 2039				
	11666564	May 30, 2038				
	11786516	May 30, 2038	DS			
	11813255	May 30, 2038				
	11826360	Feb 16, 2039	DS			
	11833164	Jan 11, 2042				
	8592362	Feb 12, 2029			Y	
	9000011	May 16, 2027				
	9095584	Feb 12, 2029				
	9480682	May 16, 2027				
	9987263	May 16, 2027				
	9999618	Apr 28, 2028				
	9999618	Apr 28, 2028				
	RE48608	Feb 12, 2029				
<u>MIGLUSTAT - OPFOLDA</u>						
N 215211 001	10208299	Sep 30, 2035	DS DP		NP	Sep 28, 2026
	10512677	Mar 07, 2033				
	10857212	Aug 12, 2037				
	10961522	Sep 30, 2035				

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<u>MIGLUSTAT - OPFOLDA</u>						
N 215211	001	11278599	Mar 07, 2033			U-3726
		11278601	Dec 29, 2036	DP		U-3726
		11753632	Sep 30, 2035			U-3726
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	001	7994220	Sep 19, 2029			U-819
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	002	7994220	Sep 19, 2029			U-819
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	003	7994220	Sep 19, 2029			U-819
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
N 022256	004	7994220	Sep 19, 2029			U-819
<u>MINOCYCLINE HYDROCHLORIDE - MINOCIN</u>						
N 050444	001	9084802	May 12, 2031			U-282
		9278105	May 12, 2031			U-282
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	001	7790705	Jun 24, 2025			U-1078
		7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	002	7541347	Apr 02, 2027			U-917
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025			U-1078
		7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	003	7790705	Jun 24, 2025			U-1078
		7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	004	7790705	Jun 24, 2025			U-1078
		7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
		9192615	Nov 17, 2031	DP		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	005	7790705	Jun 24, 2025			U-1078
		7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
		9192615	Nov 17, 2031	DP		
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	006	7790705	Jun 24, 2025			U-1078

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<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	006	7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
		8722650	Jun 24, 2025			U-1078
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	007	7790705	Jun 24, 2025			U-1078
		7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
		8722650	Jun 24, 2025			U-1078
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
N 050808	008	7790705	Jun 24, 2025			U-1078
		7919483	Mar 07, 2027			U-1078
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-1078
		8722650	Jun 24, 2025			U-1078
<u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u>						
N 201922	001	7541347	Apr 02, 2027			U-917
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025			U-124
		7919483	Mar 07, 2027			U-124
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-124
<u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u>						
N 201922	003	7541347	Apr 02, 2027			U-917
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025			U-124
		7919483	Mar 07, 2027			U-124
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-124
<u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u>						
N 201922	005	7541347	Apr 02, 2027			U-917
		7544373	Apr 02, 2027	DP		
		7790705	Jun 24, 2025			U-124
		7919483	Mar 07, 2027			U-124
		8252776	Jun 24, 2025			U-124
		8268804	Jun 24, 2025			U-124
<u>MINOCYCLINE HYDROCHLORIDE - MINOLIRA</u>						
N 209269	001	11103517	Apr 07, 2036	DP		
<u>MINOCYCLINE HYDROCHLORIDE - MINOLIRA</u>						
N 209269	002	11103517	Apr 07, 2036	DP		
<u>MINOCYCLINE HYDROCHLORIDE - AMZEEQ</u>						
N 212379	001	10086080	Oct 01, 2030			U-2647
		10137200	Oct 01, 2030			U-2647
		10213512	Oct 01, 2030	DP		U-2647
		10265404	Oct 01, 2030	DP		

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<u>MINOCYCLINE HYDROCHLORIDE - AMZEEQ</u>						
N 212379	001 10398641	Sep 08, 2037	U-2647			
	10517882	Oct 01, 2030	U-2647			
	10821187	Oct 01, 2030	U-2647			
	10849847	Sep 08, 2037	U-2647			
	8865139	Oct 01, 2030	DP U-2647			
	8945516	Oct 01, 2030	DP			
	8992896	Oct 01, 2030	DP U-2647			
	9675700	Oct 01, 2030	DP U-2647			
<u>MINOCYCLINE HYDROCHLORIDE - ZILXI</u>						
N 213690	001 10213512	Oct 01, 2030	DP U-1631			
	10265404	Oct 01, 2030	DP			
	10322186	Oct 01, 2030	U-1631			
	10946101	Oct 01, 2030	U-1631			
	8865139	Oct 01, 2030	DP U-1631			
	8945516	Oct 01, 2030	DP			
	8992896	Oct 01, 2030	DP U-1631			
	9675700	Oct 01, 2030	DP U-1631			
<u>MIPOMERSEN SODIUM - KYNAMRO</u>						
N 203568	001 7511131	Jan 29, 2027	DS			
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	001 10842780	Sep 28, 2029	DP U-2996		I-855	Mar 25, 2024
	10842780	Sep 28, 2029	DP U-3670		PED	Sep 25, 2024
	10842780*PED	Mar 28, 2030				
	11707451	Sep 28, 2029	U-2996			
	11707451	Sep 28, 2029	U-3670			
	11707451*PED	Mar 28, 2030				
	8772315	Oct 30, 2028	U-2300			
	8772315*PED	Apr 30, 2029				
	8835474*PED	May 04, 2024				
	RE44872*PED	May 04, 2024				
<u>MIRABEGRON - MYRBETRIO</u>						
N 202611	002 10842780	Sep 28, 2029	DP U-2996		I-855	Mar 25, 2024
	10842780	Sep 28, 2029	DP U-3670		PED	Sep 25, 2024
	10842780*PED	Mar 28, 2030				
	11707451	Sep 28, 2029	U-2996			
	11707451	Sep 28, 2029	U-3670			
	11707451*PED	Mar 28, 2030				
	8772315	Oct 30, 2028	U-2300			
	8772315*PED	Apr 30, 2029				
	8835474*PED	May 04, 2024				
	RE44872*PED	May 04, 2024				
<u>MIRABEGRON - MYRBETRIO GRANULES</u>						
N 213801	001 10058536	Mar 31, 2036	DP U-3108		NP	Mar 25, 2024
	10058536*PED	Oct 01, 2036			PED	Sep 25, 2024
	7342117*PED	May 04, 2024				
	7982049*PED	May 04, 2024				

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<u>MITAPIVAT SULFATE - PYRUKYND</u>						
N 216196 001	10632114	May 03, 2032	U-3320		NCE	Feb 17, 2027
	11234976	Oct 11, 2038	U-3321		ODE-392	Feb 17, 2029
	11254652	Nov 21, 2038	DS DP			
	11793806	Apr 12, 2033	U-3320			
	9193701	Oct 26, 2032	U-3319			
	9682080	May 03, 2032	U-3319			
	9980961	May 03, 2032	U-3319			
	RE49582	Feb 24, 2031	DS DP			
<u>MITAPIVAT SULFATE - PYRUKYND</u>						
N 216196 002	10632114	May 03, 2032	U-3320		NCE	Feb 17, 2027
	11234976	Oct 11, 2038	U-3321		ODE-392	Feb 17, 2029
	11254652	Nov 21, 2038	DS DP			
	11793806	Apr 12, 2033	U-3320			
	9193701	Oct 26, 2032	U-3319			
	9682080	May 03, 2032	U-3319			
	9980961	May 03, 2032	U-3319			
	RE49582	Feb 24, 2031	DS DP			
<u>MITAPIVAT SULFATE - PYRUKYND</u>						
N 216196 003	10632114	May 03, 2032	U-3320		NCE	Feb 17, 2027
	11234976	Oct 11, 2038	U-3321		ODE-392	Feb 17, 2029
	11254652	Nov 21, 2038	DS DP			
	11793806	Apr 12, 2033	U-3320			
	9193701	Oct 26, 2032	U-3319			
	9682080	May 03, 2032	U-3319			
	9980961	May 03, 2032	U-3319			
	RE49582	Feb 24, 2031	DS DP			
<u>MITOMYCIN - MITOSOL</u>						
N 022572 001	7806265	Feb 01, 2029	DP			
	8186511	Jul 19, 2026	DP			
	9205075	Jul 19, 2026	DP			
	9539241	Jan 02, 2028	DS DP U-2095			
	9649428	May 21, 2029	U-2095			
<u>MITOMYCIN - JELMYTO</u>						
N 211728 001	9040074	Jan 20, 2031	DP		ODE-289	Apr 15, 2027
	9950069	Jan 20, 2031	DP			
<u>MOBOCERTINIB SUCCINATE - EXKIVITY</u>						
N 215310 001	10227342	May 13, 2035	DS DP U-3220		NCE	Sep 15, 2026
	9796712	May 13, 2035	DS DP		ODE-374	Sep 15, 2028
<u>MOMELOTINIB DIHYDROCHLORIDE - OJJAARA</u>						
N 216873 001	8486941	Jan 03, 2030	DS DP U-1201		NCE	Sep 15, 2028
	9809559	Jun 11, 2035	U-1201		ODE-441	Sep 15, 2030
	RE48285	Jun 11, 2035	DS DP U-1201			
<u>MOMELOTINIB DIHYDROCHLORIDE - OJJAARA</u>						
N 216873 002	8486941	Jan 03, 2030	DS DP U-1201		NCE	Sep 15, 2028
	9809559	Jun 11, 2035	U-1201		ODE-441	Sep 15, 2030
	RE48285	Jun 11, 2035	DS DP U-1201			

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<u>MOMELOTINIB DIHYDROCHLORIDE - OJJAARA</u>						
N 216873	003 8486941	Jan 03, 2030	DS DP U-1201		NCE	Sep 15, 2028
	9809559	Jun 11, 2035	U-1201		ODE-441	Sep 15, 2030
	RE48285	Jun 11, 2035	DS DP U-1201			
<u>MOMETASONE FUROATE - SINUVA</u>						
N 209310	001 10232152	Nov 24, 2034	DP U-2272			
	10357640	Oct 03, 2031	U-2272			
	10406332	Mar 13, 2034	DP			
	7544192	Nov 29, 2026	U-2272			
	7662141	Mar 12, 2024	U-2272			
	7713255	Mar 12, 2024	U-2272			
	7951130	Mar 12, 2024	U-2272			
	7951131	Mar 12, 2024	U-2272			
	7951133	Mar 12, 2024	U-2272			
	8025635	Jun 12, 2027	DP U-2272			
	8109918	Mar 12, 2024	U-2272			
	8763222	Feb 08, 2032	DP			
	9585681	Apr 04, 2026	U-2272			
<u>MOMETASONE FUROATE; OLOPATADINE HYDROCHLORIDE - RYALTRIS</u>						
N 211746	001 10016443	Sep 04, 2034	U-3297		NP	Jan 13, 2025
	10376526	Sep 04, 2034	DP			
	10517880	Sep 04, 2034	DP U-3297			
	10548907	Sep 04, 2034	U-3297			
	10561672	Sep 04, 2034	DP			
	10646500	Sep 04, 2034	U-3297			
	10758550	Sep 04, 2034	U-3296			
	10765686	Sep 04, 2034	U-3295			
	11400101	Sep 04, 2034	U-3297			
	11679210	Sep 03, 2038	DP			
	9078923	Sep 04, 2034	DP U-3297			
	9370483	Sep 04, 2034	DP			
	9750754	Sep 04, 2034	DP U-3297			
	9937189	Sep 04, 2034	DP			
<u>MONOMETHYL FUMARATE - BAFIERTAM</u>						
N 210296	001 10098863	Feb 27, 2035	DP U-1384			
	10105335	Feb 27, 2035	DP			
	10105336	Feb 27, 2035	DP U-1384			
	10105337	Feb 27, 2035	DP			
	10918615	Aug 12, 2035	DP U-1384			
	10918616	Jun 03, 2035	U-1384			
	10918617	Aug 10, 2035	DP			
	10945985	Aug 14, 2035	DP			
	11590095	Mar 18, 2036	U-1384			
	9326947	Feb 27, 2035	DP			
	9326965	Feb 27, 2035	U-1384			
	9511043	Feb 27, 2035	U-1384			
	9517209	Feb 27, 2035	DP			
	9566259	Feb 27, 2035	DP			
	9636318	Feb 27, 2035	U-1384			

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<u>MONOMETHYL FUMARATE - BAFIERTAM</u>						
N 210296	001 9636319	Feb 27, 2035	DP			
	9814691	Feb 27, 2035	U-1384			
	9814692	Feb 27, 2035	U-1384			
	9820960	Feb 27, 2035	DP			
	9820961	Feb 27, 2035	DP U-1384			
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022195	001				NPP	Jun 02, 2024
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022195	002				NPP	Jun 02, 2024
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022207	001				NPP	Jun 02, 2024
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 022207	002				NPP	Jun 02, 2024
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	001 9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034	U-43			
	9192608	Mar 12, 2034	U-55			
	9248229	Mar 12, 2034	DP			
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	002 9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034	U-43			
	9192608	Mar 12, 2034	U-55			
	9248229	Mar 12, 2034	DP			
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	003 9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034	U-43			
	9192608	Mar 12, 2034	U-55			
	9248229	Mar 12, 2034	DP			
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	004 9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034	U-43			
	9192608	Mar 12, 2034	U-55			
	9248229	Mar 12, 2034	DP			
<u>MORPHINE SULFATE - MORPHINE SULFATE</u>						
N 204223	005 9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034	U-43			
	9192608	Mar 12, 2034	U-55			
	9248229	Mar 12, 2034	DP			
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	001 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	002 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			

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<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	003	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - MORPHABOND ER</u>						
N 206544	004	10314788	Aug 12, 2028	DP		
		7955619	Aug 12, 2028	DP		
<u>MORPHINE SULFATE - ARYMO ER</u>						
N 208603	001	9044402	Jul 01, 2033	DP U-1556		
		9549899	Jul 01, 2033	DP U-1556		
<u>MORPHINE SULFATE - ARYMO ER</u>						
N 208603	002	9044402	Jul 01, 2033	DP U-1556		
		9549899	Jul 01, 2033	DP U-1556		
<u>MORPHINE SULFATE - ARYMO ER</u>						
N 208603	003	9044402	Jul 01, 2033	DP U-1556		
		9549899	Jul 01, 2033	DP U-1556		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	001	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	002	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	003	7682633	Jun 19, 2027	U-1510		
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027	U-1510		
		8623418	Nov 07, 2029	U-1640		
		8685443	Jul 03, 2025	U-1508		
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	004	7682633	Jun 19, 2027	U-1510		

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<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	004	7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027		U-1510	
		8623418	Nov 07, 2029		U-1640	
		8685443	Jul 03, 2025		U-1508	
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	005	7682633	Jun 19, 2027		U-1510	
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027		U-1510	
		8623418	Nov 07, 2029		U-1640	
		8685443	Jul 03, 2025		U-1508	
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N 022321	006	7682633	Jun 19, 2027		U-1510	
		7682634	Jun 19, 2027	DP		
		7815934	Dec 12, 2027	DP		
		8158156	Jun 19, 2027		U-1510	
		8623418	Nov 07, 2029		U-1640	
		8685443	Jul 03, 2025		U-1508	
		8685444	Jul 03, 2025	DP		
		8846104	Jun 19, 2027	DP		
		8877247	Jun 19, 2027	DP		
<u>MOTIXAFORTIDE ACETATE - APHEXDA</u>						
N 217159	001				NCE	Sep 08, 2028
					ODE-442	Sep 08, 2030
<u>MOXIDECTIN - MOXIDECTIN</u>						
N 210867	001				ODE-193	Jun 13, 2025
<u>MOXIFLOXACIN HYDROCHLORIDE - MOXEZA</u>						
N 022428	001	8450311	May 29, 2029	DP		
		9114168	May 29, 2029	DP		
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N 204286	001	10166205	Jan 31, 2033	DP		
		10166206	Jan 31, 2033	DP		
		10695303	Jan 31, 2033	DP		
		10729667	Jan 31, 2033	DP		
		8778365	Jan 31, 2033	DP		
		9161914	Jan 31, 2033		U-540	
<u>NALDEMEDINE TOSYLATE - SYMPROIC</u>						
N 208854	001	10952968	May 13, 2033	DS DP		
		9108975	Nov 11, 2031	DS DP		
		RE46365	Jan 11, 2028	DS DP		

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<u>NALDEMEDINE TOSYLATE - SYMPROIC</u>						
N 208854	001 RE46375	Oct 05, 2026	DS DP U-1185			
<u>NALMEFENE HYDROCHLORIDE - OPVEE</u>						
N 217470	001 11458091	Jul 10, 2038	DP U-3630		NP	May 22, 2026
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760	001 7056500	Jun 29, 2024	DP U-1185			
	7786133	Sep 16, 2028	DS DP			
	8067431	Dec 16, 2024	U-1185			
	9012469	Apr 02, 2032	DS DP			
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760	002 7056500	Jun 29, 2024	DP U-1185			
	7786133	Sep 16, 2028	DS DP			
	8067431	Dec 16, 2024	U-1185			
	9012469	Apr 02, 2032	DS DP			
<u>NALOXONE HYDROCHLORIDE - EVZIO</u>						
N 205787	001 10143972	May 24, 2031	U-2476			
	10220158	Mar 20, 2035	DP U-2500			
	10314977	Nov 23, 2024	DP			
	10322239	Feb 28, 2031	U-1907			
	10335549	Apr 30, 2025	DP			
	10737028	Nov 23, 2024	DP			
	10960155	Jun 25, 2026	DP			
	11590286	Dec 12, 2026	DP			
	7731686	Jun 10, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8627816	Feb 04, 2032	DP			
	8926594	Mar 31, 2026	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9056170	Nov 23, 2024	DP			
	9238108	Feb 20, 2027	DP			
	9278182	Feb 01, 2026	DP			
	9474869	Feb 28, 2031	DP U-1907			
	9517307	Jul 18, 2034	DP U-1925			
	9724471	May 23, 2027	DP U-2092			
	9737669	Nov 23, 2024	DP			

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<u>NALOXONE HYDROCHLORIDE - NARCAN</u>						
N 208411	002 9480644	Mar 16, 2035	DP U-1903	Y		
	9707226	Mar 16, 2035	DP U-1903	Y		
<u>NALOXONE HYDROCHLORIDE - REXTOVY</u>						
N 208969	001				NP	Mar 07, 2026
<u>NALOXONE HYDROCHLORIDE - EVZIO (AUTOINJECTOR)</u>						
N 209862	001 10143792	May 24, 2031	U-2476			
	10220158	Mar 20, 2035	DP U-2500			
	10314977	Nov 23, 2024	DP			
	10322239	Feb 28, 2031	U-1907			
	10335549	Apr 30, 2025	DP			
	10737028	Nov 23, 2024	DP			
	10960155	Jun 25, 2026	DP			
	11590286	Dec 12, 2026	DP			
	7731686	Jun 01, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8627816	Feb 04, 2032	DP			
	8926594	Mar 31, 2026	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9056170	Nov 23, 2024	DP			
	9238108	Feb 20, 2027	DP			
	9278182	Feb 01, 2026	DP			
	9474869	Feb 28, 2031	DP U-1907			
	9517307	Jul 18, 2034	DP U-1925			
	9724471	May 23, 2027	DP U-2092			
	9737669	Nov 23, 2024	DP			
<u>NALOXONE HYDROCHLORIDE - KLOXXADO</u>						
N 212045	001 10722510	Aug 26, 2034	DP U-3110			
	10973814	Aug 26, 2034	DP U-3110			
	11135155	Aug 26, 2034	DP			
	11617713	Aug 26, 2034	DP U-3110			
	11628139	Aug 26, 2034	DP U-3110			
<u>NALOXONE HYDROCHLORIDE - ZIMHI</u>						
N 212854	001 11027072	May 24, 2039	DP			
	11571518	Jun 14, 2041	DP U-3515			
	11571518	Jun 14, 2041	DP U-3516			

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<u>NALOXONE HYDROCHLORIDE - ZIMHI</u>						
N 212854	001 11571518	Jun 14, 2041	DP U-3517			
<u>NALOXONE HYDROCHLORIDE - NALOXONE HYDROCHLORIDE (AUTOINJECTOR)</u>						
N 215457	001 10143792	May 24, 2031	U-2476			
	10314977	Nov 23, 2024	DP			
	10322239	Feb 28, 2031	U-1907			
	10335549	Apr 30, 2025	DP			
	10737028	Nov 23, 2024	DP			
	11590286	Dec 12, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9056170	Nov 23, 2024	DP			
	9474869	Feb 28, 2031	DP U-1907			
	9737669	Nov 23, 2024	DP			
	9814838	Feb 28, 2031	DP			
<u>NALOXONE HYDROCHLORIDE - RIVIVE</u>						
N 217722	001 11020343	May 11, 2032	U-3671			
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	001 9073933	Mar 30, 2025	DS			
	9522919	Mar 30, 2025	DS DP			
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	002 9073933	Mar 30, 2025	DS			
	9522919	Mar 30, 2025	DS DP			
<u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u>						
N 205777	003 9073933	Mar 30, 2025	DS			
	9522919	Mar 30, 2025	DS DP			
<u>NALTREXONE - VIVITROL</u>						
N 021897	001 7919499	Oct 15, 2029	U-1123			
	7919499	Oct 15, 2029	U-1124			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	001 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	002 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	003 7815934	Dec 12, 2027	DP			
	8685443	Jul 03, 2025	U-1508			

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<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	004	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	005	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u>						
N 207621	006	7815934	Dec 12, 2027	DP		
		8685443	Jul 03, 2025		U-1508	
<u>NAPROXEN SODIUM - NAPROXEN SODIUM</u>						
N 021920	001	10022344	Mar 03, 2026	DP	U-1731	
		10022344	Mar 03, 2026	DP	U-1732	
		10028925	Mar 03, 2026	DP	U-1731	
		10028925	Mar 03, 2026	DP	U-1732	
		11090280	Mar 03, 2026	DP	U-1731	
		11090280	Mar 03, 2026	DP	U-1732	
		9693978	Mar 03, 2026	DP		
		9693979	Mar 03, 2026	DP		
<u>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</u>						
N 021926	001	7332183	Oct 02, 2025	DP	U-867	
		7332183*PED	Apr 02, 2026			
<u>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</u>						
N 021926	002	7332183	Oct 02, 2025	DP	U-1719	
		7332183*PED	Apr 02, 2026			
<u>NEBIVOLOL HYDROCHLORIDE; VALSARTAN - BYVALSON</u>						
N 206302	001	7803838	Aug 29, 2026	DP		
		7838552	Oct 04, 2027		U-185	
<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	001	10351854	Oct 09, 2035	DS DP		NCE Sep 29, 2028
		10738311	Oct 09, 2035	DS DP	U-3709	ODE-443 Sep 29, 2030
		11053502	Oct 29, 2035	DS DP		
		11286488	Oct 12, 2038	DS DP	U-3709	
		11359203	Oct 09, 2035	DS DP	U-3709	
		11661604	Oct 12, 2038	DS DP	U-3709	
<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	002	10351854	Oct 09, 2035	DS DP		NCE Sep 29, 2028
		10738311	Oct 09, 2035	DS DP	U-3709	ODE-443 Sep 29, 2030
		11053502	Oct 29, 2035	DS DP		
		11286488	Oct 12, 2038	DS DP	U-3709	
		11359203	Oct 09, 2035	DS DP	U-3709	
		11661604	Oct 12, 2038	DS DP	U-3709	
<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	003	10351854	Oct 09, 2035	DS DP		NCE Sep 29, 2028
		10738311	Oct 09, 2035	DS DP	U-3709	ODE-443 Sep 29, 2030
		11053502	Oct 29, 2035	DS DP		
		11286488	Oct 12, 2038	DS DP	U-3709	
		11359203	Oct 09, 2035	DS DP	U-3709	

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<u>NEDOSIRAN SODIUM - RIVFLOZA</u>						
N 215842	003 11661604	Oct 12, 2038	DS DP U-3709			
<u>NEPAFENAC - NEVANAC</u>						
N 021862	001 7834059	Jan 31, 2027		U-1095		
	8071648	Dec 02, 2025	DP			
	8324281	Dec 02, 2025	DP			
<u>NEPAFENAC - ILEVRO</u>						
N 203491	001 7947295	Jun 08, 2024	DP			
	8921337	Mar 31, 2032	DP			
	9662398	Dec 01, 2030	DP			
<u>NERATINIB MALEATE - NERLYNX</u>						
N 208051	001 10035788	Oct 15, 2028		U-2043	D-182	Jun 28, 2024
	10035788	Oct 15, 2028		U-3047		
	10035788	Oct 15, 2028		U-3097		
	7399865	Dec 29, 2030	DS DP			
	7982043	Oct 08, 2025		U-2043		
	7982043	Oct 08, 2025		U-3047		
	7982043	Oct 08, 2025		U-3097		
	8518446	Nov 20, 2030	DP	U-2043		
	8518446	Nov 20, 2030	DP	U-3047		
	8518446	Nov 20, 2030	DP	U-3097		
	8669273	Jul 18, 2031		U-3047		
	8790708	Nov 05, 2030	DP	U-2043		
	8790708	Nov 05, 2030	DP	U-3047		
	8790708	Nov 05, 2030	DP	U-3097		
	9139558	Oct 15, 2028		U-2043		
	9139558	Oct 15, 2028		U-3047		
	9139558	Oct 15, 2028		U-3097		
	9211291	Mar 24, 2030		U-2043		
	9211291	Mar 24, 2030		U-3097		
	9265784	Aug 04, 2029		U-3047		
	9630946	Oct 15, 2028		U-2043		
	9630946	Oct 15, 2028		U-3047		
	9630946	Oct 15, 2028		U-3097		
<u>NETARSUDIL MESYLATE - RHOPRESSA</u>						
N 208254	001 10174017	Jan 27, 2030	DS DP U-1524			
	10532993	Jul 11, 2026		U-1524		
	10588901	Mar 14, 2034	DS DP U-1524			
	10654844	Jan 27, 2030	DS DP U-1524			
	10882840	Jul 11, 2026		U-1524		
	11021456	Jul 11, 2026		U-1524		
	11028081	Jan 27, 2030		U-1524		
	11185538	Mar 14, 2034	DP			
	11618748	Jan 27, 2030		U-1524		
	8394826	Nov 10, 2030	DS DP U-1524			
	8450344	Jul 11, 2026	DS DP U-1524			
	9096569	Jul 11, 2026	DS DP U-1524			
	9415043	Mar 14, 2034	DS			
	9931336	Mar 14, 2034	DS DP U-1524			

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<u>NETARSUDIL MESYLATE - RHOPRESSA</u>						
N 208254	001	10174017	Jan 27, 2030	DS DP	U-1524	
		10532993	Jul 11, 2026		U-1524	
		10588901	Mar 14, 2034	DS DP	U-1524	
		10654844	Jan 27, 2030	DS DP	U-1524	
		10882840	Jul 11, 2026		U-1524	
		11021456	Jul 11, 2026		U-1524	
		11028081	Jan 27, 2030		U-1524	
		11185538	Mar 14, 2034	DP		
		11618748	Jan 27, 2030		U-1524	
		8394826	Nov 10, 2030	DS DP	U-1524	
		8450344	Jul 11, 2026	DS DP	U-1524	
		9096569	Jul 11, 2026	DS DP	U-1524	
		9415043	Mar 14, 2034	DS		
		9931336	Mar 14, 2034	DS DP	U-1524	
<u>NETUPITANT; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u>						
N 205718	001	10233154	Sep 25, 2035	DS		
		10676440	Sep 25, 2035	DS DP		
		10828297	Dec 17, 2030		U-2293	
		10961195	Sep 25, 2035	DS DP		
		11559523	Nov 18, 2030	DP	U-3522	
		8623826	Nov 18, 2030		U-2293	
		8951969	Nov 18, 2030	DP		
		9186357	Nov 18, 2030		U-2293	
		9271975	Sep 09, 2031		U-2293	
		9943515	Nov 18, 2030		U-2293	
		9951016	Sep 25, 2035	DS DP		
<u>NEVIRAPINE - VIRAMUNE XR</u>						
N 201152	001	8460704	Mar 12, 2029		U-1409	
<u>NICARDIPINE HYDROCHLORIDE - NICARDIPINE HYDROCHLORIDE</u>						
A 215377	001				CGT	Mar 20, 2024
<u>NICARDIPINE HYDROCHLORIDE - NICARDIPINE HYDROCHLORIDE</u>						
A 215377	002				CGT	Mar 20, 2024
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER</u>						
N 019734	002	10758616	Apr 18, 2027	DP		
		11547758	Apr 18, 2027		U-1029	
		7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027		U-1029	
		8455524	Apr 18, 2027		U-1029	
		9364564	Dec 26, 2027	DP		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N 019734	003	10758616	Apr 18, 2027	DP		
		11547758	Apr 18, 2027		U-1029	
		7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027		U-1029	
		8455524	Apr 18, 2027		U-1029	
		9364564	Dec 26, 2027	DP		

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<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N 019734	004	10758616	Apr 18, 2027	DP		
		11547758	Apr 18, 2027	U-1029		
		7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027	U-1029		
		8455524	Apr 18, 2027	U-1029		
		9364564	Dec 26, 2027	DP		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER</u>						
N 019734	005	7612102	Dec 26, 2027	DP		
		7659291	Apr 18, 2027	U-1029		
		8455524	Apr 18, 2027	U-1029		
		9364564	Dec 26, 2027	DP		
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 018612	002	8323683	Apr 30, 2028			
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 020066	002	8323683	Apr 30, 2028	DP		
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 022360	001	8501164	Jun 14, 2029	DP		
		8940772	Apr 30, 2029	DP		
<u>NICOTINE POLACRILEX - NICORETTE</u>						
N 022360	002	8501164	Jun 14, 2029	DP		
		8940772	Apr 30, 2029	DP		
<u>NIFURTIMOX - LAMPIT</u>						
N 213464	001				NCE	Aug 06, 2025
					ODE-319	Aug 06, 2027
<u>NIFURTIMOX - LAMPIT</u>						
N 213464	002				NCE	Aug 06, 2025
					ODE-319	Aug 06, 2027
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	001	7169791*PED	Jan 04, 2024		ODE-171	Mar 22, 2025
		8163904	Aug 23, 2028	DS DP	ODE-172	Mar 22, 2025
		8163904*PED	Feb 23, 2029		ODE-380	Sep 23, 2028
		8293756	Sep 25, 2027	DP	PED	Sep 22, 2025
		8293756*PED	Mar 25, 2028		PED	Sep 22, 2025
		8389537	Jul 18, 2026	U-1374	PED	Mar 23, 2029
		8389537*PED	Jan 18, 2027			
		8415363	Jul 18, 2026	DS DP U-1374		
		8415363	Jul 18, 2026	DS DP U-1407		
		8415363*PED	Jan 18, 2027			
		8501760	Jul 18, 2026	DP		
		8501760*PED	Jan 18, 2027			
		9061029	Apr 07, 2032	DS U-1374		
		9061029	Apr 07, 2032	DS U-3231		
		9061029*PED	Oct 07, 2032			
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	002	7169791*PED	Jan 04, 2024		ODE-171	Mar 22, 2025
		8163904	Aug 23, 2028	DS DP	ODE-172	Mar 22, 2025

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<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	002	8163904*PED			ODE-380	Sep 23, 2028
		8293756		DP	PED	Sep 22, 2025
		8293756*PED			PED	Sep 22, 2025
		8389537		U-1374	PED	Mar 23, 2029
		8389537*PED				
		8415363	DS DP	U-1374		
		8415363	DS DP	U-1407		
		8415363*PED				
		8501760		DP		
		8501760*PED				
		9061029	DS	U-1374		
		9061029	DS	U-3231		
		9061029*PED				
<u>NILOTINIB HYDROCHLORIDE - TASIGNA</u>						
N 022068	003	7169791*PED			ODE-171	Mar 22, 2025
		8163904	DS DP		ODE-172	Mar 22, 2025
		8163904*PED			ODE-380	Sep 23, 2028
		8293756		DP	PED	Sep 22, 2025
		8293756*PED			PED	Sep 22, 2025
		8389537		U-1374	PED	Mar 23, 2029
		8389537*PED				
		8415363	DS DP	U-1374		
		8415363	DS DP	U-1407		
		8415363*PED				
		8501760		DP		
		8501760*PED				
		9061029	DS	U-1374		
		9061029	DS	U-3231		
		9061029*PED				
<u>NIMODIPINE - NYMALIZE</u>						
N 203340	002	10342787		DP	U-2804	
		10576070		DP	U-2804	
		11207306			U-2804	
		11413277		DP	U-2804	
		11806338		DP		
		8517997		DP		
<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832	001	10105323		DP	ODE-261	Sep 06, 2026
		10105323*PED			PED	Mar 06, 2027
		10154990			U-2620	
		10154990*PED				
		6762180	DS DP			
		6762180*PED				
		7119093	DS DP			
		9907756		DP		
		9907756*PED				

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<u>NINTEDANIB ESYLATE - OFEV</u>						
N 205832 002	10105323	Jun 04, 2029	DP		ODE-261	Sep 06, 2026
	10105323*PED	Dec 04, 2029			PED	Mar 06, 2027
	10154990	Jan 08, 2026		U-2620		
	10154990*PED	Jul 08, 2026				
	6762180	Oct 01, 2025	DS DP			
	6762180*PED	Apr 01, 2026				
	7119093	Feb 21, 2024	DS DP			
	9907756	Jun 07, 2029	DP			
	9907756*PED	Dec 07, 2029				
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 208447 001	11091459	Mar 27, 2038	DP		ODE-133	Mar 27, 2024
	11673877	Mar 27, 2038	DP U-3646		ODE-277	Oct 23, 2026
	11673877	Mar 27, 2038	DP U-3647		ODE-295	Apr 29, 2027
	8071579	Aug 12, 2027		U-2655		
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027		U-2655		
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031		U-2655		
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876 001	11091459	Mar 27, 2038	DP			
	11673877	Mar 27, 2038	DP U-3646			
	11673877	Mar 27, 2038	DP U-3647			
	11730725	Jan 04, 2039	DP			
	8071579	Aug 12, 2027		U-2655		
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027		U-2655		
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031		U-2655		
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876 002	11091459	Mar 27, 2038	DP			
	11673877	Mar 27, 2038	DP U-3646			
	11673877	Mar 27, 2038	DP U-3647			
	11730725	Jan 04, 2039	DP			
	8071579	Aug 12, 2027		U-2655		
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027		U-2655		
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031		U-2655		
<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876 003	11091459	Mar 27, 2038	DP			
	11673877	Mar 27, 2038	DP U-3646			
	11673877	Mar 27, 2038	DP U-3647			
	11730725	Jan 04, 2039	DP			
	8071579	Aug 12, 2027		U-2655		
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027		U-2655		
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031		U-2655		

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<u>NIRAPARIB TOSYLATE - ZEJULA</u>						
N 214876	003 11091459	Mar 27, 2038	DP			
	11673877	Mar 27, 2038	DP U-3646			
	11673877	Mar 27, 2038	DP U-3647			
	11730725	Jan 04, 2039	DP			
	8071579	Aug 12, 2027	U-2655			
	8071623	Mar 27, 2031	DS DP			
	8143241	Aug 12, 2027	U-2655			
	8436185	Apr 24, 2029	DS			
	8859562	Aug 04, 2031	U-2655			
<u>NIRMATRELVIR; RITONAVIR - PAXLOVID (COPACKAGED)</u>						
N 217188	001 11351149	Aug 05, 2041	DS DP U-3629		NCE	May 25, 2028
	11541034	Oct 31, 2041	U-3629			
<u>NIRMATRELVIR; RITONAVIR - PAXLOVID (COPACKAGED)</u>						
N 217188	002				NCE	May 25, 2028
<u>NIROGACESTAT HYDROBROMIDE - OGSIVEO</u>						
N 217677	001 10590087	Aug 09, 2039	DS			
	10710966	Aug 09, 2039	DS U-3754			
	10941118	Aug 09, 2039	DS U-3754			
	11504354	Jul 08, 2042	DP			
	11612588	Jul 08, 2042	DP U-3754			
	11807611	Sep 08, 2042	DP U-3754			
	11820748	Aug 09, 2039	DP			
	11844780	Sep 08, 2042	DP U-3754			
	11845732	Aug 09, 2039	DS U-3754			
	7342118	Aug 18, 2025	DS			
	7795447	Aug 18, 2025	DS			
	7951958	Mar 11, 2025	DS			
<u>NITISINONE - ORFADIN</u>						
N 206356	001 9301932	Feb 28, 2033	DP U-1836			
<u>NITISINONE - NITYR</u>						
N 209449	001 10328029	Jan 05, 2035	DP U-1836			
<u>NITISINONE - NITYR</u>						
N 209449	002 10328029	Jan 05, 2035	DP U-1836			
<u>NITISINONE - NITYR</u>						
N 209449	003 10328029	Jan 05, 2035	DP U-1836			
<u>NITRIC OXIDE - INOMAX</u>						
N 020845	002 8282966	Jun 30, 2029	U-1286			
	8291904	Jan 06, 2031	DP U-1226			
	8293284	Jun 30, 2029	U-1286			
	8431163	Jun 30, 2029	U-1286			
	8431163*PED	Dec 30, 2029				
	8573209	Jan 06, 2031	DP			
	8573209*PED	Jul 06, 2031				
	8573210	Jan 06, 2031	DP U-1453			
	8573210*PED	Jul 06, 2031				
	8776794	Jan 06, 2031	DP U-1226			

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<u>NITRIC OXIDE - INOMAX</u>						
N 020845 002	8776794*PED	Jul 06, 2031				
	8776795	Jan 06, 2031	DP U-1226			
	8776795*PED	Jul 06, 2031				
	8795741	Jun 30, 2029	U-1286			
	8795741*PED	Dec 30, 2029				
	8846112	Jun 30, 2029	U-1286			
	8846112*PED	Dec 30, 2029				
<u>NITRIC OXIDE - INOMAX</u>						
N 020845 003	8282966	Jun 30, 2029	U-1286			
	8291904	Jan 06, 2031	DP U-1226			
	8293284	Jun 30, 2029	U-1286			
	8431163	Jun 30, 2029	U-1286			
	8431163*PED	Dec 30, 2029				
	8573209	Jan 06, 2031	DP			
	8573209*PED	Jul 06, 2031				
	8573210	Jan 06, 2031	DP U-1453			
	8573210*PED	Jul 06, 2031				
	8776794	Jan 06, 2031	DP U-1226			
	8776794*PED	Jul 06, 2031				
	8776795	Jan 06, 2031	DP U-1226			
	8776795*PED	Jul 06, 2031				
	8795741	Jun 30, 2029	U-1286			
	8795741*PED	Dec 30, 2029				
	8846112	Jun 30, 2029	U-1286			
	8846112*PED	Dec 30, 2029				
	9265911	Jan 06, 2031	DP U-1824			
	9265911*PED	Jul 06, 2031				
	9279794	Feb 19, 2034	DP U-1823			
	9279794*PED	Aug 19, 2034				
	9295802	Jan 06, 2031	DP U-1226			
	9295802*PED	Jul 06, 2031				
	9408993	Jan 06, 2031	DP U-1824			
	9408993*PED	Jul 06, 2031				
	9770570	May 03, 2036	U-2148			
	9770570*PED	Nov 03, 2036				
<u>NITRIC OXIDE - GENOSYL</u>						
N 202860 001	10124142	Aug 18, 2025	U-3037			
	10213572	Feb 12, 2036	DP			
	10737051	Oct 20, 2035	DP			
	10814092	Oct 17, 2025	U-3037			
	10926054	Aug 13, 2029	DP			
	11103669	Jun 21, 2030	DP			
	11291793	Aug 18, 2025	DP			
	11383059	Aug 18, 2025	U-3037			
	11511252	Sep 21, 2029	DP			
	11554241	Aug 18, 2025	U-3037			
	11672938	Jul 22, 2040	U-3037			
	7560076	Apr 21, 2027	DP			
	7618594	Oct 17, 2026	DP			

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<u>NITRIC OXIDE - GENOSYL</u>						
N 202860	001	7947227	Oct 17, 2026	U-3037		
		8057742	Jan 18, 2026	U-3037		
		8226916	Aug 18, 2025	U-3037		
		8607785	Jul 14, 2030	DP		
		8609028	Aug 18, 2025	U-3037		
		8821801	Aug 18, 2025	DP		
		8944049	Aug 13, 2029	DP		
		9522249	Aug 18, 2025	DP		
		9604028	Aug 13, 2029	U-2793		
		9701538	Jan 28, 2029	DP		
		9956373	Aug 18, 2025	U-3037		
<u>NITROGLYCERIN - NITROLINGUAL PUMPSPRAY</u>						
N 018705	002	7872049	Mar 12, 2029	DP	U-2223	
<u>NITROGLYCERIN - GONITRO</u>						
N 208424	001	9101592	Mar 11, 2032	DP		
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 214628	001	10159657	Jan 30, 2038	DP		
		10226436	Jan 30, 2038	DP	U-3461	
		10420735	Jan 30, 2038	DP	U-3461	
		10568850	Jan 30, 2038	DP		
		11413259	Jan 30, 2038	DP	U-3461	
		11602508	Jan 30, 2038	DP		
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 214628	002	10159657	Jan 30, 2038	DP		
		10226436	Jan 30, 2038	DP	U-3461	
		10420735	Jan 30, 2038	DP	U-3461	
		10568850	Jan 30, 2038	DP		
		11413259	Jan 30, 2038	DP	U-3461	
		11602508	Jan 30, 2038	DP		
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 214628	003	10159657	Jan 30, 2038	DP		
		10226436	Jan 30, 2038	DP	U-3461	
		10420735	Jan 30, 2038	DP	U-3461	
		10568850	Jan 30, 2038	DP		
		11413259	Jan 30, 2038	DP	U-3461	
		11602508	Jan 30, 2038	DP		
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 215700	001	10888534	Apr 26, 2039	DP		
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 215700	002	10888534	Apr 26, 2039	DP		
<u>NOREPINEPHRINE BITARTRATE - NOREPINEPHRINE BITARTRATE IN 0.9% SODIUM CHLORIDE</u>						
N 215700	003	10888534	Apr 26, 2039	DP		
<u>NORGESTREL - OPILL</u>						
N 017031	001				RTO	Jul 13, 2026

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<u>NUSINERSEN SODIUM - SPINRAZA</u>						
N 209531	001	10266822	Dec 05, 2025			U-1942
		10266822	Dec 05, 2025			U-1943
		10266822	Dec 05, 2025			U-1944
		10436802	Sep 11, 2035			U-1941
		10436802	Sep 11, 2035			U-1942
		10436802	Sep 11, 2035			U-1943
		10436802	Sep 11, 2035			U-1944
		10436802	Sep 11, 2035			U-2093
		10436802	Sep 11, 2035			U-2094
		7838657	Jul 11, 2027	DS		
		8110560	Dec 05, 2025			U-1942
		8110560	Dec 05, 2025			U-1943
		8110560	Dec 05, 2025			U-1944
		8361977	May 27, 2030	DS DP		
		8980853	Nov 24, 2030			U-1941
		9717750	Jun 17, 2030			U-1942
		9717750	Jun 17, 2030			U-1943
		9717750	Jun 17, 2030			U-2093
		9717750	Jun 17, 2030			U-2094
		9926559	Jan 09, 2034			U-1943
<u>OBETICHOLIC ACID - OCALIVA</u>						
N 207999	001	10047117	Sep 06, 2033			U-1854
		10052337	Apr 26, 2036	DP		
		10174073	Jun 17, 2033	DS		
		10751349	Apr 26, 2036	DP		
		10758549	Apr 26, 2036			U-2945
		9238673	Jun 17, 2033	DP		
		RE48286	Feb 21, 2027	DS DP		
<u>OBETICHOLIC ACID - OCALIVA</u>						
N 207999	002	10047117	Sep 06, 2033			U-1854
		10052337	Apr 26, 2036	DP		
		10174073	Jun 17, 2033	DS		
		10751349	Apr 26, 2036	DP		
		10758549	Apr 26, 2036			U-2945
		9238673	Jun 17, 2033	DP		
		RE48286	Feb 21, 2027	DS DP		
<u>OCTREOTIDE ACETATE - MYCAPSSA</u>						
N 208232	001	10238709	Feb 03, 2036			U-2857
		10695397	Feb 03, 2036			U-2857
		11052126	Feb 03, 2036			U-2857
		11141457	Dec 28, 2040			U-3232
		11338011	Feb 03, 2036			U-2857
		11510963	Feb 03, 2036			U-2857
		8329198	Sep 17, 2029	DP		
		8535695	Sep 17, 2029			U-2857
		9265812	Sep 17, 2029	DP		
		9566246	Sep 17, 2029	DP		

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<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 001	10011633	Nov 08, 2031	U-3186		I-918	Jun 13, 2026
	10011633	Nov 08, 2031	U-3187		NCE	Jul 20, 2026
	10011633	Nov 08, 2031	U-3648		ODE-363	Jul 20, 2028
	10011633	Nov 08, 2031	U-3649		ODE-436	Jun 13, 2030
	10093697	Nov 08, 2031	U-3186			
	10093697	Nov 08, 2031	U-3187			
	10093697	Nov 08, 2031	U-3648			
	10093697	Nov 08, 2031	U-3649			
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10487111	Nov 08, 2031	U-3648			
	10487111	Nov 08, 2031	U-3649			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	10981952	Nov 08, 2031	U-3648			
	10981952	Nov 08, 2031	U-3649			
	11365182	Jun 20, 2039	U-3186			
	11365182	Jun 20, 2039	U-3187			
	11365182	Jun 20, 2039	U-3648			
	11365182	Jun 20, 2039	U-3649			
	11583539	Nov 12, 2041	U-3186			
	11732006	Nov 08, 2031	U-3186			
	11732006	Nov 08, 2031	U-3187			
	11732006	Nov 08, 2031	U-3648			
	11732006	Nov 08, 2031	U-3649			
	11801226	Jun 20, 2039	DP			
	11802115	Jun 20, 2039	DP			
	9694018	Nov 08, 2031	U-3186			
	9694018	Nov 08, 2031	U-3187			
	9694018	Nov 08, 2031	U-3648			
	9694018	Nov 08, 2031	U-3649			
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 002	10011633	Nov 08, 2031	U-3186		I-918	Jun 13, 2026
	10011633	Nov 08, 2031	U-3187		NCE	Jul 20, 2026
	10011633	Nov 08, 2031	U-3648		ODE-363	Jul 20, 2028
	10011633	Nov 08, 2031	U-3649		ODE-436	Jun 13, 2030
	10093697	Nov 08, 2031	U-3186			
	10093697	Nov 08, 2031	U-3187			
	10093697	Nov 08, 2031	U-3648			
	10093697	Nov 08, 2031	U-3649			
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10487111	Nov 08, 2031	U-3648			
	10487111	Nov 08, 2031	U-3649			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	10981952	Nov 08, 2031	U-3648			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 002	10981952	Nov 08, 2031	U-3649			
	11365182	Jun 20, 2039	U-3186			
	11365182	Jun 20, 2039	U-3187			
	11365182	Jun 20, 2039	U-3648			
	11365182	Jun 20, 2039	U-3649			
	11583539	Nov 12, 2041	U-3186			
	11732006	Nov 08, 2031	U-3186			
	11732006	Nov 08, 2031	U-3187			
	11732006	Nov 08, 2031	U-3648			
	11732006	Nov 08, 2031	U-3649			
	11801226	Jun 20, 2039	DP			
	11802115	Jun 20, 2039	DP			
	9694018	Nov 08, 2031	U-3186			
	9694018	Nov 08, 2031	U-3187			
	9694018	Nov 08, 2031	U-3648			
	9694018	Nov 08, 2031	U-3649			
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 003	10011633	Nov 08, 2031	U-3186		I-918	Jun 13, 2026
	10011633	Nov 08, 2031	U-3187		NCE	Jul 20, 2026
	10011633	Nov 08, 2031	U-3648		ODE-363	Jul 20, 2028
	10011633	Nov 08, 2031	U-3649		ODE-436	Jun 13, 2030
	10093697	Nov 08, 2031	U-3186			
	10093697	Nov 08, 2031	U-3187			
	10093697	Nov 08, 2031	U-3648			
	10093697	Nov 08, 2031	U-3649			
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10487111	Nov 08, 2031	U-3648			
	10487111	Nov 08, 2031	U-3649			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	10981952	Nov 08, 2031	U-3648			
	10981952	Nov 08, 2031	U-3649			
	11365182	Jun 20, 2039	U-3186			
	11365182	Jun 20, 2039	U-3187			
	11365182	Jun 20, 2039	U-3648			
	11365182	Jun 20, 2039	U-3649			
	11583539	Nov 12, 2041	U-3186			
	11732006	Nov 08, 2031	U-3186			
	11732006	Nov 08, 2031	U-3187			
	11732006	Nov 08, 2031	U-3648			
	11732006	Nov 08, 2031	U-3649			
	11801226	Jun 20, 2039	DP			
	11802115	Jun 20, 2039	DP			
	9694018	Nov 08, 2031	U-3186			
	9694018	Nov 08, 2031	U-3187			
	9694018	Nov 08, 2031	U-3648			
	9694018	Nov 08, 2031	U-3649			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 003	10011633	Nov 08, 2031	U-3186		I-918	Jun 13, 2026
	10011633	Nov 08, 2031	U-3187		NCE	Jul 20, 2026
	10011633	Nov 08, 2031	U-3648		ODE-363	Jul 20, 2028
	10011633	Nov 08, 2031	U-3649		ODE-436	Jun 13, 2030
	10093697	Nov 08, 2031	U-3186			
	10093697	Nov 08, 2031	U-3187			
	10093697	Nov 08, 2031	U-3648			
	10093697	Nov 08, 2031	U-3649			
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10487111	Nov 08, 2031	U-3648			
	10487111	Nov 08, 2031	U-3649			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	10981952	Nov 08, 2031	U-3648			
	10981952	Nov 08, 2031	U-3649			
	11365182	Jun 20, 2039	U-3186			
	11365182	Jun 20, 2039	U-3187			
	11365182	Jun 20, 2039	U-3648			
	11365182	Jun 20, 2039	U-3649			
	11583539	Nov 12, 2041	U-3186			
	11732006	Nov 08, 2031	U-3186			
	11732006	Nov 08, 2031	U-3187			
	11732006	Nov 08, 2031	U-3648			
	11732006	Nov 08, 2031	U-3649			
	11801226	Jun 20, 2039	DP			
	11802115	Jun 20, 2039	DP			
	9694018	Nov 08, 2031	U-3186			
	9694018	Nov 08, 2031	U-3187			
	9694018	Nov 08, 2031	U-3648			
	9694018	Nov 08, 2031	U-3649			
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498 004	10011633	Nov 08, 2031	U-3186		I-918	Jun 13, 2026
	10011633	Nov 08, 2031	U-3187		NCE	Jul 20, 2026
	10011633	Nov 08, 2031	U-3648		ODE-363	Jul 20, 2028
	10011633	Nov 08, 2031	U-3649		ODE-436	Jun 13, 2030
	10093697	Nov 08, 2031	U-3186			
	10093697	Nov 08, 2031	U-3187			
	10093697	Nov 08, 2031	U-3648			
	10093697	Nov 08, 2031	U-3649			
	10487111	Nov 08, 2031	U-3186			
	10487111	Nov 08, 2031	U-3187			
	10487111	Nov 08, 2031	U-3648			
	10487111	Nov 08, 2031	U-3649			
	10975046	Jun 20, 2039	DS			
	10981952	Nov 08, 2031	U-3186			
	10981952	Nov 08, 2031	U-3187			
	10981952	Nov 08, 2031	U-3648			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ODEVIXIBAT - BYLVAY</u>						
N 215498	004	10981952	Nov 08, 2031	U-3649		
		11365182	Jun 20, 2039	U-3186		
		11365182	Jun 20, 2039	U-3187		
		11365182	Jun 20, 2039	U-3648		
		11365182	Jun 20, 2039	U-3649		
		11583539	Nov 12, 2041	U-3186		
		11732006	Nov 08, 2031	U-3186		
		11732006	Nov 08, 2031	U-3187		
		11732006	Nov 08, 2031	U-3648		
		11732006	Nov 08, 2031	U-3649		
		11801226	Jun 20, 2039	DP		
		11802115	Jun 20, 2039	DP		
		9694018	Nov 08, 2031	U-3186		
		9694018	Nov 08, 2031	U-3187		
		9694018	Nov 08, 2031	U-3648		
		9694018	Nov 08, 2031	U-3649		
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	001	10300054	Aug 23, 2031	DP U-3140	NCE	May 28, 2026
		10300054	Aug 23, 2031	DP U-3141		
		10716785	Aug 23, 2031	U-3136		
		10716785	Aug 23, 2031	U-3137		
		11185541	Aug 23, 2031	U-3140		
		11241425	Aug 23, 2031	U-3137		
		11351166	Aug 23, 2031	U-3140		
		11351166	Aug 23, 2031	U-3141		
		11707466	Nov 12, 2041	DP		
		11793805	Aug 23, 2031	U-3734		
		7262298	Nov 23, 2025	DS		
		8778960	Feb 13, 2032	U-3136		
		8778960	Feb 13, 2032	U-3137		
		9119848	Aug 30, 2031	DS		
		9126977	Aug 23, 2031	DP U-3136		
		9126977	Aug 23, 2031	DP U-3137		
		9517235	Aug 23, 2031	U-3138		
		9517235	Aug 23, 2031	U-3139		
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	002	10300054	Aug 23, 2031	DP U-3140	NCE	May 28, 2026
		10300054	Aug 23, 2031	DP U-3141		
		10716785	Aug 23, 2031	U-3136		
		10716785	Aug 23, 2031	U-3137		
		11185541	Aug 23, 2031	U-3140		
		11241425	Aug 23, 2031	U-3137		
		11351166	Aug 23, 2031	U-3140		
		11351166	Aug 23, 2031	U-3141		
		11707466	Nov 12, 2041	DP		
		11793805	Aug 23, 2031	U-3734		
		7262298	Nov 23, 2025	DS		
		8778960	Feb 13, 2032	U-3136		
		8778960	Feb 13, 2032	U-3137		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	002	9119848	Aug 30, 2031	DS		
		9126977	Aug 23, 2031	DP U-3136		
		9126977	Aug 23, 2031	DP U-3137		
		9517235	Aug 23, 2031	U-3138		
		9517235	Aug 23, 2031	U-3139		
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	003	10300054	Aug 23, 2031	DP U-3140	NCE	May 28, 2026
		10300054	Aug 23, 2031	DP U-3141		
		10716785	Aug 23, 2031	U-3136		
		10716785	Aug 23, 2031	U-3137		
		11185541	Aug 23, 2031	U-3140		
		11241425	Aug 23, 2031	U-3137		
		11351166	Aug 23, 2031	U-3140		
		11351166	Aug 23, 2031	U-3141		
		11707466	Nov 12, 2041	DP		
		11793805	Aug 23, 2031	U-3734		
		7262298	Nov 23, 2025	DS		
		8778960	Feb 13, 2032	U-3136		
		8778960	Feb 13, 2032	U-3137		
		9119848	Aug 30, 2031	DS		
		9126977	Aug 23, 2031	DP U-3136		
		9126977	Aug 23, 2031	DP U-3137		
		9517235	Aug 23, 2031	U-3138		
		9517235	Aug 23, 2031	U-3139		
<u>OLANZAPINE; SAMIDORPHAN L-MALATE - LYBALVI</u>						
N 213378	004	10300054	Aug 23, 2031	DP U-3140	NCE	May 28, 2026
		10300054	Aug 23, 2031	DP U-3141		
		10716785	Aug 23, 2031	U-3136		
		10716785	Aug 23, 2031	U-3137		
		11185541	Aug 23, 2031	U-3140		
		11241425	Aug 23, 2031	U-3137		
		11351166	Aug 23, 2031	U-3140		
		11351166	Aug 23, 2031	U-3141		
		11707466	Nov 12, 2041	DP		
		11793805	Aug 23, 2031	U-3734		
		7262298	Nov 23, 2025	DS		
		8778960	Feb 13, 2032	U-3136		
		8778960	Feb 13, 2032	U-3137		
		9119848	Aug 30, 2031	DS		
		9126977	Aug 23, 2031	DP U-3136		
		9126977	Aug 23, 2031	DP U-3137		
		9517235	Aug 23, 2031	U-3138		
		9517235	Aug 23, 2031	U-3139		
<u>OLAPARIB - LYNPARZA</u>						
N 206162	001	7449464	Oct 11, 2024	DS DP		
		7981889	Oct 11, 2024	DS DP		
		8143241	Aug 12, 2027	U-1634		
		8247416	Sep 24, 2028	DS		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 206162 001	8859562	Aug 04, 2031	U-1634			
	8912187	Mar 12, 2024	U-1634			
<u>OLAPARIB - LYNPARZA</u>						
N 208558 001	11633396	Oct 07, 2029	DP		I-885	Mar 11, 2025
	7449464	Oct 11, 2024	DS DP		I-914	May 31, 2026
	7981889	Oct 11, 2024	DS DP		ODE-180	Aug 17, 2024
	8071579	Aug 12, 2027	U-2480		ODE-181	Aug 17, 2024
	8071579	Aug 12, 2027	U-2482		ODE-226	Dec 19, 2025
	8071579	Aug 12, 2027	U-2483		ODE-283	Dec 27, 2026
	8071579	Aug 12, 2027	U-2716		ODE-306	May 08, 2027
	8071579	Aug 12, 2027	U-2819			
	8071579	Aug 12, 2027	U-2820			
	8071579	Aug 12, 2027	U-2821			
	8071579	Aug 12, 2027	U-2822			
	8071579	Aug 12, 2027	U-2823			
	8071579	Aug 12, 2027	U-2824			
	8071579	Aug 12, 2027	U-2832			
	8071579	Aug 12, 2027	U-2833			
	8071579	Aug 12, 2027	U-3333			
	8071579	Aug 12, 2027	U-3631			
	8071579	Aug 12, 2027	U-3695			
	8143241	Aug 12, 2027	U-2480			
	8143241	Aug 12, 2027	U-2482			
	8143241	Aug 12, 2027	U-2483			
	8143241	Aug 12, 2027	U-2716			
	8143241	Aug 12, 2027	U-2819			
	8143241	Aug 12, 2027	U-2820			
	8143241	Aug 12, 2027	U-2821			
	8143241	Aug 12, 2027	U-2822			
	8143241	Aug 12, 2027	U-2823			
	8143241	Aug 12, 2027	U-2824			
	8143241	Aug 12, 2027	U-2832			
	8143241	Aug 12, 2027	U-2833			
	8143241	Aug 12, 2027	U-3333			
	8143241	Aug 12, 2027	U-3631			
	8143241	Aug 12, 2027	U-3695			
	8475842	Dec 31, 2029	DP			
	8859562	Aug 04, 2031	U-2101			
	8859562	Aug 04, 2031	U-2103			
	8859562	Aug 04, 2031	U-2480			
	8859562	Aug 04, 2031	U-2482			
	8859562	Aug 04, 2031	U-2483			
	8859562	Aug 04, 2031	U-2716			
	8859562	Aug 04, 2031	U-2819			
	8859562	Aug 04, 2031	U-2820			
	8859562	Aug 04, 2031	U-2821			
	8859562	Aug 04, 2031	U-2822			
	8859562	Aug 04, 2031	U-2823			
	8859562	Aug 04, 2031	U-2824			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 208558 001	8859562	Aug 04, 2031	U-2832			
	8859562	Aug 04, 2031	U-2833			
	8859562	Aug 04, 2031	U-3333			
	8859562	Aug 04, 2031	U-3631			
	8859562	Aug 04, 2031	U-3695			
	8912187	Mar 12, 2024	U-2480			
	8912187	Mar 12, 2024	U-2482			
	8912187	Mar 12, 2024	U-2483			
	8912187	Mar 12, 2024	U-2819			
	8912187	Mar 12, 2024	U-2820			
	8912187	Mar 12, 2024	U-2821			
	8912187	Mar 12, 2024	U-2822			
	8912187	Mar 12, 2024	U-2823			
	8912187	Mar 12, 2024	U-2824			
	8912187	Mar 12, 2024	U-3333			
	8912187	Mar 12, 2024	U-3695			
	9169235	Mar 12, 2024	U-2832			
	9169235	Mar 12, 2024	U-2833			
	9169235	Mar 12, 2024	U-3631			
	9566276	Mar 12, 2024	U-2716			
<u>OLAPARIB - LYNPARZA</u>						
N 208558 002	11633396	Oct 07, 2029	DP		I-885	Mar 11, 2025
	7449464	Oct 11, 2024	DS DP		I-914	May 31, 2026
	7981889	Oct 11, 2024	DS DP		ODE-180	Aug 17, 2024
	8071579	Aug 12, 2027	U-2480		ODE-181	Aug 17, 2024
	8071579	Aug 12, 2027	U-2482		ODE-226	Dec 19, 2025
	8071579	Aug 12, 2027	U-2483		ODE-283	Dec 27, 2026
	8071579	Aug 12, 2027	U-2716		ODE-306	May 08, 2027
	8071579	Aug 12, 2027	U-2819			
	8071579	Aug 12, 2027	U-2820			
	8071579	Aug 12, 2027	U-2821			
	8071579	Aug 12, 2027	U-2822			
	8071579	Aug 12, 2027	U-2823			
	8071579	Aug 12, 2027	U-2824			
	8071579	Aug 12, 2027	U-2832			
	8071579	Aug 12, 2027	U-2833			
	8071579	Aug 12, 2027	U-3333			
	8071579	Aug 12, 2027	U-3631			
	8071579	Aug 12, 2027	U-3695			
	8143241	Aug 12, 2027	U-2480			
	8143241	Aug 12, 2027	U-2482			
	8143241	Aug 12, 2027	U-2483			
	8143241	Aug 12, 2027	U-2716			
	8143241	Aug 12, 2027	U-2819			
	8143241	Aug 12, 2027	U-2820			
	8143241	Aug 12, 2027	U-2821			
	8143241	Aug 12, 2027	U-2822			
	8143241	Aug 12, 2027	U-2823			
	8143241	Aug 12, 2027	U-2824			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OLAPARIB - LYNPARZA</u>						
N 208558 002	8143241	Aug 12, 2027	U-2832			
	8143241	Aug 12, 2027	U-2833			
	8143241	Aug 12, 2027	U-3333			
	8143241	Aug 12, 2027	U-3631			
	8143241	Aug 12, 2027	U-3695			
	8475842	Dec 31, 2029	DP			
	8859562	Aug 04, 2031	U-2101			
	8859562	Aug 04, 2031	U-2103			
	8859562	Aug 04, 2031	U-2480			
	8859562	Aug 04, 2031	U-2482			
	8859562	Aug 04, 2031	U-2483			
	8859562	Aug 04, 2031	U-2716			
	8859562	Aug 04, 2031	U-2819			
	8859562	Aug 04, 2031	U-2820			
	8859562	Aug 04, 2031	U-2821			
	8859562	Aug 04, 2031	U-2822			
	8859562	Aug 04, 2031	U-2823			
	8859562	Aug 04, 2031	U-2824			
	8859562	Aug 04, 2031	U-2832			
	8859562	Aug 04, 2031	U-2833			
	8859562	Aug 04, 2031	U-3333			
	8859562	Aug 04, 2031	U-3631			
	8859562	Aug 04, 2031	U-3695			
	8912187	Mar 12, 2024	U-2480			
	8912187	Mar 12, 2024	U-2482			
	8912187	Mar 12, 2024	U-2483			
	8912187	Mar 12, 2024	U-2819			
	8912187	Mar 12, 2024	U-2820			
	8912187	Mar 12, 2024	U-2821			
	8912187	Mar 12, 2024	U-2822			
	8912187	Mar 12, 2024	U-2823			
	8912187	Mar 12, 2024	U-2824			
	8912187	Mar 12, 2024	U-3333			
	8912187	Mar 12, 2024	U-3695			
	9169235	Mar 12, 2024	U-2832			
	9169235	Mar 12, 2024	U-2833			
	9169235	Mar 12, 2024	U-3631			
	9566276	Mar 12, 2024	U-2716			
<u>OLICERIDINE - OLINVKY</u>						
N 210730 001	11077098	Mar 23, 2032	DS DP U-2986		NCE	Oct 30, 2025
	8835488	Mar 23, 2032	DS DP U-2986			
	9309234	Mar 23, 2032	DS DP U-2986			
	9642842	Mar 23, 2032	DP U-2986			
<u>OLICERIDINE - OLINVKY</u>						
N 210730 002	11077098	Mar 23, 2032	DS DP U-2986		NCE	Oct 30, 2025
	8835488	Mar 23, 2032	DS DP U-2986			
	9309234	Mar 23, 2032	DS DP U-2986			
	9642842	Mar 23, 2032	DP U-2986			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLICERIDINE - OLINVYK</u>						
N 210730	003 11077098	Mar 23, 2032	DS DP U-2986		NCE	Oct 30, 2025
	8835488	Mar 23, 2032	DS DP U-2986			
	9309234	Mar 23, 2032	DS DP U-2986			
	9642842	Mar 23, 2032	DP U-2986			
<u>OLODATEROL HYDROCHLORIDE - STRIVERDI RESPIMAT</u>						
N 203108	001 7220742	May 12, 2025	DS DP U-1547			
	7284474	Aug 26, 2024	DP			
	7396341	Oct 10, 2026	DP U-1547			
	7727984	Jan 19, 2027	DS			
	7837235	Mar 13, 2028	DP			
	7896264	May 26, 2025	DP			
	8034809	May 12, 2025	U-1547			
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
<u>OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u>						
N 206756	001 7220742	May 12, 2025	DS DP U-1703			
	7284474	Aug 26, 2024	DP			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7727984	Jan 19, 2027	DS			
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	8034809	May 12, 2025	U-1702			
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
<u>OLOPATADINE HYDROCHLORIDE - PATADAY ONCE DAILY RELIEF</u>						
N 206276	001 8791154	May 19, 2032	DP U-1680			
	9533053	May 19, 2032	DP			
<u>OLUTASIDENIB - REZLIDHIA</u>						
N 215814	001 10414752	Sep 18, 2035	DP		NCE	Dec 01, 2027
	10532047	May 16, 2039	DS		ODE-413	Dec 01, 2029
	10550098	Sep 18, 2035	DP			
	10959994	May 16, 2039	DP			
	11013733	May 16, 2039	U-3496			
	11013734	May 16, 2039	U-3495			
	11376246	May 16, 2039	U-3495			
	11497743	May 16, 2039	U-3495			
	11498913	Sep 18, 2035	DP			
	11723905	Nov 12, 2039	DP			
	11738018	Jul 17, 2039	U-3684			
	9834539	Sep 18, 2035	DS DP U-3497			
<u>OMACETAXINE MEPESUCCINATE - SYNRIBO</u>						
N 203585	001 6987103	Oct 26, 2026	U-1300			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>OMADACYCLINE TOSYLATE - NUZYRA</u>						
N 209816 001	10111890	Aug 03, 2037	U-2444		NCE	Oct 02, 2023
	10124014	Mar 05, 2029	U-2449		GAIN	Oct 02, 2028
	10383884	Oct 31, 2037	U-2576			
	10835542	Oct 31, 2037	U-2576			
	7553828	Jun 02, 2024	DS			
	8383610	Sep 23, 2030	DS			
	9265740	Mar 05, 2029	U-1569			
	9314475	Mar 18, 2031	DP			
	9724358	Mar 05, 2029	U-1569			
<u>OMADACYCLINE TOSYLATE - NUZYRA</u>						
N 209817 001	10124014	Mar 05, 2029	U-2449		NCE	Oct 02, 2023
	10383884	Oct 31, 2037	U-2576		GAIN	Oct 02, 2028
	10835542	Oct 31, 2037	U-2576			
	7553828	Jun 02, 2024	DS			
	9265740	Mar 05, 2029	DP			
	9724358	Mar 05, 2029	U-1569			
<u>OMAVELOXOLONE - SKYCLARYS</u>						
N 216718 001	11091430	Apr 20, 2029	U-3552		NCE	Feb 28, 2028
	8124799	Dec 03, 2029	DS		ODE-427	Feb 28, 2030
	8440854	Apr 20, 2029	DP			
	8993640	Apr 24, 2033	DS DP			
	9670147	Apr 20, 2029	DS			
	9701709	Apr 24, 2033	DS DP			
<u>OMBITASVIR; PARITAPREVR; RITONAVIR - TECHNIVIE</u>						
N 207931 001	8268349	Aug 25, 2024	DP			
	8268349*PED	Feb 25, 2025				
	8399015	Aug 25, 2024	DP			
	8399015*PED	Feb 25, 2025				
	8420596	Apr 10, 2031	DS DP			
	8420596*PED	Oct 10, 2031				
	8642538	Sep 10, 2029	DS DP	U-1638		
	8686026	Jun 09, 2031	DP			
	8691938	Apr 13, 2032	DS DP			
	9006387	Jun 10, 2030	U-1687			
	9044480	Apr 10, 2031	U-1638			
<u>OMEGA-3-CARBOXYLIC ACIDS - EPANOVA</u>						
N 205060 001	10117844	Jan 04, 2033	U-2447			
	7960370	Dec 20, 2026	DP			
	8383678	Feb 07, 2025	DP	U-1511		
	9012501	Feb 07, 2025	DP	U-1511		
	9050308	Jan 04, 2033	U-1511			
	9050309	Jan 04, 2033	DS			
	9132112	Feb 07, 2025	DP	U-1511		
<u>OMEPRAZOLE - OMEPRAZOLE</u>						
N 022032 001	9023391	Aug 16, 2025	DP			

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<u>OMEPRAZOLE - OMEPRAZOLE</u>						
N 209400	001	10076494	Dec 08, 2036	DP		
		10835488	Dec 08, 2036	DP		
<u>OMEPRAZOLE; SODIUM BICARBONATE - KONVOMEF</u>						
N 213593	001	10751333	Jul 16, 2039	DP		
		11103492	Jul 16, 2039	DP		
		11633478	Jul 16, 2039	DP		
		11771686	Mar 01, 2040	U-623		
<u>OMIDENEPAG ISOPROPYL - OMLONTI</u>						
N 215092	001	10179127	Jan 08, 2035	DP U-3454	NCE	Sep 22, 2027
		10702511	Jan 08, 2035	DP U-3454		
		10765750	Jan 08, 2035	DP		
		10774072	Jun 10, 2035	DS		
		11197849	Jan 08, 2035	DP U-3454		
		11666563	Jul 16, 2039	U-3454		
		11793798	Jan 08, 2035	DP U-3454		
		8648097	Oct 13, 2029	DS DP		
		8685986	Oct 13, 2029	DP		
		9415038	Jan 08, 2035	DP U-3454		
		RE48183	Jan 08, 2035	DP U-3454		
<u>ONDANSETRON - ZUPLENZ</u>						
N 022524	001	8580830	Nov 23, 2029	DP		
		9095577	Jul 13, 2030	DP		
<u>ONDANSETRON - ZUPLENZ</u>						
N 022524	002	8580830	Nov 23, 2029	DP		
		9095577	Jul 13, 2030	DP		
<u>OPICAPONE - ONGENTYS</u>						
N 212489	001	10071085	Mar 31, 2030	DP	NCE	Apr 24, 2025
		10357468	May 27, 2035	U-2812		
		10583130	Mar 31, 2030	U-2811		
		8168793	Apr 02, 2029	DS DP U-2811		
		8524746	Jul 14, 2029	U-2811		
		8907099	May 12, 2027	DS		
		9550759	Jul 26, 2026	U-2811		
		9550759	Jul 26, 2026	U-2817		
		9550759	Jul 26, 2026	U-2818		
		9630955	Dec 12, 2032	DS DP U-2811		
		9745290	Oct 10, 2027	DP U-2811		
<u>OPICAPONE - ONGENTYS</u>						
N 212489	002	10071085	Mar 31, 2030	DP	NCE	Apr 24, 2025
		10357468	May 27, 2035	U-2812		
		10583130	Mar 31, 2030	U-2811		
		8168793	Apr 02, 2029	DS DP U-2811		
		8524746	Jul 14, 2029	U-2811		
		8907099	May 12, 2027	DS		
		9550759	Jul 26, 2026	U-2811		
		9550759	Jul 26, 2026	U-2817		
		9550759	Jul 26, 2026	U-2818		

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<u>OPICAPONE - ONGENTYS</u>						
N 212489	002 9630955	Dec 12, 2032	DS DP U-2811			
	9745290	Oct 10, 2027	DP U-2811			
<u>ORITAVANCIN DIPHOSPHATE - ORBACTIV</u>						
N 206334	001 8420592	Aug 29, 2029	U-1570		NCE	Aug 06, 2019
	9649352	Jul 16, 2035	DP		GAIN	Aug 06, 2024
	9682061	Apr 26, 2030	U-1569			
<u>ORITAVANCIN DIPHOSPHATE - KIMYRSA</u>						
N 214155	001 8420592	Aug 29, 2029	U-3101		NCE	Aug 06, 2019
	9649352	Jul 16, 2035	DS DP		NP	Mar 12, 2024
	9682061	Apr 26, 2030	U-3101		GAIN	Aug 06, 2024
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801	001 10143680	Jul 06, 2035	DP		NCE	Mar 06, 2025
	10709691	Oct 12, 2035	U-2770		ODE-286	Mar 06, 2027
	8314097	Mar 27, 2029	DS DP			
	8609862	Jan 13, 2031	U-2770			
	8835646	Aug 23, 2026	DS DP			
	9434754	Jan 13, 2031	DS			
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801	002 10143680	Jul 06, 2035	DP		NCE	Mar 06, 2025
	10709691	Oct 12, 2035	U-2770		ODE-286	Mar 06, 2027
	8314097	Mar 27, 2029	DS DP			
	8609862	Jan 13, 2031	U-2770			
	8835646	Aug 23, 2026	DS DP			
	9434754	Jan 13, 2031	DS			
<u>OSILODROSTAT PHOSPHATE - ISTURISA</u>						
N 212801	003 10143680	Jul 06, 2035	DP		NCE	Mar 06, 2025
	10709691	Oct 12, 2035	U-2770		ODE-286	Mar 06, 2027
	8314097	Mar 27, 2029	DS DP			
	8609862	Jan 13, 2031	U-2770			
	8835646	Aug 23, 2026	DS DP			
	9434754	Jan 13, 2031	DS			
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065	001 10183020	Jan 02, 2035	DP U-1777		ODE-176	Apr 18, 2025
	10183020	Jan 02, 2035	DP U-2289		ODE-337	Dec 18, 2027
	10183020	Jan 02, 2035	DP U-3016			
	11524951	Jul 25, 2032	DS DP			
	8946235	Aug 08, 2032	DS DP U-1777			
	8946235	Aug 08, 2032	DS DP U-2289			
	8946235	Aug 08, 2032	DS DP U-3016			
	9732058	Jul 25, 2032	DS DP U-1777			
	9732058	Jul 25, 2032	DS DP U-2289			
	9732058	Jul 25, 2032	DS DP U-3016			
<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065	002 10183020	Jan 02, 2035	DP U-1777		ODE-176	Apr 18, 2025
	10183020	Jan 02, 2035	DP U-2289		ODE-337	Dec 18, 2027
	10183020	Jan 02, 2035	DP U-3016			
	11524951	Jul 25, 2032	DS DP			

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<u>OSIMERTINIB MESYLATE - TAGRISSO</u>						
N 208065	002	8946235	Aug 08, 2032	DS DP U-1777		
		8946235	Aug 08, 2032	DS DP U-2289		
		8946235	Aug 08, 2032	DS DP U-3016		
		9732058	Jul 25, 2032	DS DP U-1777		
		9732058	Jul 25, 2032	DS DP U-2289		
		9732058	Jul 25, 2032	DS DP U-3016		
<u>OSPHEMIFENE - OSPHENA</u>						
N 203505	001	6245819	Jul 21, 2025	U-1370		
		6245819	Jul 21, 2025	U-905		
		8236861	Aug 11, 2026	U-1369		
		8236861	Aug 11, 2026	U-1370		
		8236861	Aug 11, 2026	U-905		
		8470890	Feb 13, 2024	U-1369		
		8470890	Feb 13, 2024	U-1370		
		8470890	Feb 13, 2024	U-905		
		8642079	Jul 09, 2028	DP		
		8772353	Feb 13, 2024	U-1369		
		8772353	Feb 13, 2024	U-1370		
		8772353	Feb 13, 2024	U-905		
		9241915	Feb 13, 2024	U-1369		
		9241915	Feb 13, 2024	U-1370		
		9241915	Feb 13, 2024	U-905		
		9855224	Feb 13, 2024	U-1369		
		9855224	Feb 13, 2024	U-1370		
		9855224	Feb 13, 2024	U-905		
<u>OTESECONAZOLE - VIVJOA</u>						
N 215888	001	10414751	Mar 17, 2036	DS DP	NCE	Apr 26, 2027
		11247981	May 09, 2033	U-3366	GAIN	Apr 26, 2032
		8236962	Apr 22, 2031	DS DP		
		8754227	Apr 22, 2031	U-3366		
		9840492	Mar 17, 2036	DS DP		
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	001	10220042	Apr 13, 2027	U-2501		
		11166960	Apr 13, 2027	DP		
		7722898	Apr 13, 2027	DP		
		7910131	Apr 13, 2027	U-2041		
		8617600	Apr 13, 2027	DP		
		8821930	Apr 13, 2027	DP		
		9119791	Apr 13, 2027	U-2041		
		9351975	Apr 13, 2027	DP		
		9370525	Apr 13, 2027	DP		
		9855278	Apr 13, 2027	DP		
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810	002	10220042	Apr 13, 2027	U-2501		
		11166960	Apr 13, 2027	DP		
		7722898	Apr 13, 2027	DP		
		7910131	Apr 13, 2027	U-2041		
		8617600	Apr 13, 2027	DP		

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<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810 002	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027		U-2041		
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<u>OXCARBAZEPINE - OXTELLAR XR</u>						
N 202810 003	10220042	Apr 13, 2027		U-2501		
	11166960	Apr 13, 2027	DP			
	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027		U-2041		
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027		U-2041		
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<u>OXYBUTYNIN CHLORIDE - GELNIQUE</u>						
N 022204 001	10449173	Nov 06, 2029	DP	U-2637		
	8920392	Mar 26, 2031		U-1644		
	9259388	Nov 06, 2029		U-1644		
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090 001	10004729	Dec 10, 2030	DP	U-1556		
	10188644	Sep 02, 2036	DP	U-1556		
	10646485	Sep 02, 2036	DP	U-1556		
	10668060	Dec 10, 2030	DP	U-1556		
	7399488	Mar 24, 2025	DP			
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025		U-1556		
	9682075	Dec 10, 2030	DP	U-1556		
	9737530	Sep 02, 2036	DP	U-1556		
	9968598	Sep 02, 2036	DP	U-1556		
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090 002	10004729	Dec 10, 2030	DP	U-1556		
	10188644	Sep 02, 2036	DP	U-1556		
	10646485	Sep 02, 2036	DP	U-1556		
	10668060	Dec 10, 2030	DP	U-1556		
	7399488	Mar 24, 2025	DP			
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025		U-1556		
	9682075	Dec 10, 2030	DP	U-1556		
	9737530	Sep 02, 2036	DP	U-1556		
	9968598	Sep 02, 2036	DP	U-1556		

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<u>OXYCODONE - XTAMPZA ER</u>						
N 208090	003	10004729	Dec 10, 2030	DP U-1556		
		10188644	Sep 02, 2036	DP U-1556		
		10646485	Sep 02, 2036	DP U-1556		
		10668060	Dec 10, 2030	DP U-1556		
		7399488	Mar 24, 2025	DP		
		7771707	Mar 24, 2025	DP		
		8449909	Mar 24, 2025	DP		
		8557291	Mar 21, 2025	DP		
		8758813	Jun 10, 2025	U-1556		
		9682075	Dec 10, 2030	DP U-1556		
		9737530	Sep 02, 2036	DP U-1556		
		9968598	Sep 02, 2036	DP U-1556		
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090	004	10004729	Dec 10, 2030	DP U-1556		
		10188644	Sep 02, 2036	DP U-1556		
		10646485	Sep 02, 2036	DP U-1556		
		10668060	Dec 10, 2030	DP U-1556		
		7399488	Mar 24, 2025	DP		
		7771707	Mar 24, 2025	DP		
		8449909	Mar 24, 2025	DP		
		8557291	Mar 21, 2025	DP		
		8758813	Jun 10, 2025	U-1556		
		9682075	Dec 10, 2030	DP U-1556		
		9737530	Sep 02, 2036	DP U-1556		
		9968598	Sep 02, 2036	DP U-1556		
<u>OXYCODONE - XTAMPZA ER</u>						
N 208090	005	10004729	Dec 10, 2030	DP U-1556		
		10188644	Sep 02, 2036	DP U-1556		
		10646485	Sep 02, 2036	DP U-1556		
		10668060	Dec 10, 2030	DP U-1556		
		7399488	Mar 24, 2025	DP		
		7771707	Mar 24, 2025	DP		
		8449909	Mar 24, 2025	DP		
		8557291	Mar 21, 2025	DP		
		8758813	Jun 10, 2025	U-1556		
		9682075	Dec 10, 2030	DP U-1556		
		9737530	Sep 02, 2036	DP U-1556		
		9968598	Sep 02, 2036	DP U-1556		
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	001	10407434	Mar 30, 2025	DS		
		10696684	Mar 30, 2025	DS		
		11304908	Aug 24, 2027	DP		
		11304909	Aug 24, 2027	U-1556		
		8808741	Aug 24, 2027	U-1556		
		8894987	Mar 29, 2030	DP		
		8894988	Aug 24, 2027	DP		
		9073933	Mar 30, 2025	DS		
		9492389	Aug 24, 2027	DP		

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<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 001	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 002	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027	U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 003	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027	U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 004	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027	U-1556			
	8808741	Aug 24, 2027	U-1556			
	8894987	Mar 29, 2030	DP			

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<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 004	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 005	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	8808741	Aug 24, 2027		U-1556		
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 006	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	8808741	Aug 24, 2027		U-1556		
	8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272 007	10407434	Mar 30, 2025	DS			
	10696684	Mar 30, 2025	DS			
	11304908	Aug 24, 2027	DP			
	11304909	Aug 24, 2027		U-1556		
	8808741	Aug 24, 2027		U-1556		

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<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N 022272	007 8894988	Aug 24, 2027	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<u>OXYCODONE HYDROCHLORIDE - OXAYDO</u>						
N 202080	001 7201920	Mar 16, 2025	DP			
	9492443	May 26, 2024	DP			
<u>OXYCODONE HYDROCHLORIDE - OXAYDO</u>						
N 202080	002 7201920	Mar 16, 2025	DP			
	9492443	May 26, 2024	DP			
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	001 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	002 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u>						
N 209777	003 10314788	Aug 12, 2028	DP			
	7955619	Aug 12, 2028	DP			
<u>OXYMETAZOLINE HYDROCHLORIDE - RHOFADÉ</u>						
N 208552	001 10335391	Jun 11, 2035		U-2567		
	10751325	Jun 11, 2035		U-2921		
	11517560	Jun 11, 2035		U-3494		
	7812049	May 02, 2028		U-1959		
	8420688	Aug 02, 2024		U-1959		
	8815929	Jan 22, 2024		U-1959		
	8883838	Dec 01, 2031	DP			
	9974773	Jun 11, 2035		U-2306		
<u>OXYMETAZOLINE HYDROCHLORIDE - UPNEEQ</u>						
N 212520	001 10799481	Dec 16, 2039		U-2849		
	10814001	Dec 16, 2039	DP			
	10898573	Dec 16, 2039	DP			
	10912765	Aug 26, 2031		U-2849		
	10940138	Dec 16, 2039		U-2849		
	11103482	Dec 16, 2039	DP			
	11311515	Dec 16, 2039	DP			
	11324722	Dec 16, 2039		U-2849		
	11541036	Dec 16, 2039	DP			
	11701343	Dec 16, 2039		U-2849		
	8357714	Aug 26, 2031		U-2849		
	9867808	Aug 26, 2031		U-2849		

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<u>OXYMETAZOLINE HYDROCHLORIDE - UPNEEQ</u>						
N 212520	001	10799481	Dec 16, 2039			U-2849
		10814001	Dec 16, 2039			DP
		10898573	Dec 16, 2039			DP
		10912765	Aug 26, 2031			U-2849
		10940138	Dec 16, 2039			U-2849
		11103482	Dec 16, 2039			DP
		11311515	Dec 16, 2039			DP
		11324722	Dec 16, 2039			U-2849
		11541036	Dec 16, 2039			DP
		11701343	Dec 16, 2039			U-2849
		8357714	Aug 26, 2031			U-2849
		9867808	Aug 26, 2031			U-2849
<u>OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE - KOVANAZE</u>						
N 208032	001	8580282	Apr 02, 2030			DP U-1876
		9308191	Apr 02, 2030			DP U-1876
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	001	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	002	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	003	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	004	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	005	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	006	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 021610	007	8808737	Jun 21, 2027			U-3085
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	001	7851482	Jul 10, 2029			DS
		8114383	Aug 08, 2024			DP
		8192722	Sep 15, 2025			DP
		8808737	Jun 21, 2027			U-1598
		8871779	Nov 22, 2029			DS
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	002	7851482	Jul 10, 2029			DS
		8114383	Aug 08, 2024			DP
		8192722	Sep 15, 2025			DP

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<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	002	8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	003	7851482	Jul 10, 2029	DS		
		8114383	Aug 08, 2024	DP		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	004	7851482	Jul 10, 2029	DS		
		8114383	Aug 08, 2024	DP		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	005	7851482	Jul 10, 2029	DS		
		8114383	Aug 08, 2024	DP		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	006	7851482	Jul 10, 2029	DS		
		8114383	Aug 08, 2024	DP		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N 201655	007	7851482	Jul 10, 2029	DS		
		8114383	Aug 08, 2024	DP		
		8192722	Sep 15, 2025	DP		
		8808737	Jun 21, 2027		U-1598	
		8871779	Nov 22, 2029	DS		
<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	001	10239846	Nov 15, 2030		I-860	May 27, 2024
		11680050	Sep 30, 2038	DS DP	U-2774	NCE Mar 25, 2025
		11680050	Sep 30, 2038	DS DP	U-3740	
		8481573	May 14, 2029	DS DP	U-2774	
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	
<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	002	10239846	Nov 15, 2030		I-860	May 27, 2024
		11680050	Sep 30, 2038	DS DP	U-2774	NCE Mar 25, 2025
		11680050	Sep 30, 2038	DS DP	U-3740	
		8481573	May 14, 2029	DS DP	U-2774	
		8796318	May 14, 2029	DS DP		
		9382217	May 14, 2029		U-2774	

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<u>OZANIMOD HYDROCHLORIDE - ZEPOSIA</u>						
N 209899	003 10239846	Nov 15, 2030	U-3132		I-860	May 27, 2024
	11680050	Sep 30, 2038	DS DP U-2774		NCE	Mar 25, 2025
	11680050	Sep 30, 2038	DS DP U-3740			
	8481573	May 14, 2029	DS DP U-2774			
	8796318	May 14, 2029	DS DP			
	9382217	May 14, 2029	U-2774			
<u>OZENOXACIN - XEPI</u>						
N 208945	001 9180200	Jan 29, 2032	DP U-805			
	9399014	Dec 15, 2029	U-805			
<u>PACLITAXEL - ABRAXANE</u>						
N 021660	001 7758891	Feb 21, 2026	U-1434			
	7758891*PED	Aug 21, 2026				
	7820788	Oct 27, 2024	DP U-1092			
	7820788	Oct 27, 2024	DP U-1290			
	7820788	Oct 27, 2024	DP U-1434			
	7820788*PED	Apr 27, 2025				
	7923536*PED	Jun 09, 2024				
	8034375	Aug 13, 2026	U-1290			
	8138229*PED	Jun 09, 2024				
	8268348	Feb 21, 2026	U-1290			
	8314156*PED	Jun 09, 2024				
	9101543	Feb 21, 2026	U-1434			
	9101543*PED	Aug 21, 2026				
	9393318	Mar 04, 2032	U-1290			
	9393318*PED	Sep 04, 2032				
	9511046	Jan 12, 2034	U-1434			
	9511046*PED	Jul 12, 2034				
	9597409	Mar 04, 2032	U-1290			
	9597409*PED	Sep 04, 2032				
<u>PACRITINIB CITRATE - VONJO</u>						
N 208712	001 8153632	Jan 17, 2029	DS DP U-3331		NCE	Feb 28, 2027
	8153632	Jan 17, 2029	DS DP U-3332		ODE-397	Feb 28, 2029
	8980873	Mar 25, 2030	DS DP U-3331			
	8980873	Mar 25, 2030	DS DP U-3332			
	9573964	May 05, 2028	U-3331			
	9573964	May 05, 2028	U-3332			
<u>PAFOLACIANINE SODIUM - CYTALUX</u>						
N 214907	001 10881747	Aug 26, 2033	DS DP U-3291		I-905	Dec 16, 2025
	9061057	Aug 26, 2033	DS DP U-3291		NCE	Nov 29, 2026
	9254341	Oct 04, 2033	DS DP		ODE-390	Nov 29, 2028
	9333270	Aug 26, 2033	DS DP U-3291			
	9341629	Aug 26, 2033	DS DP			
	9789208	Aug 26, 2033	DS DP U-3291			
<u>PALBOCICLIB - IBRANCE</u>						
N 207103	001 10723730	Feb 08, 2034	DS DP			
	RE47739	Mar 05, 2027	DS DP			

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<u>PALBOCICLIB - IBRANCE</u>						
N 207103 002	10723730	Feb 08, 2034	DS DP			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 207103 003	10723730	Feb 08, 2034	DS DP			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 212436 001	10723730	Feb 08, 2034	DS DP			
	11065250	Aug 19, 2036	DP			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 212436 002	10723730	Feb 08, 2034	DS DP			
	11065250	Aug 19, 2036	DP			
	RE47739	Mar 05, 2027	DS DP			
<u>PALBOCICLIB - IBRANCE</u>						
N 212436 003	10723730	Feb 08, 2034	DS DP			
	11065250	Aug 19, 2036	DP			
	RE47739	Mar 05, 2027	DS DP			
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 001	9439906	Jan 26, 2031		U-1901		
	9439906	Jan 26, 2031		U-2757		
	9439906	Jan 26, 2031		U-2758		
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 002	9439906	Jan 26, 2031		U-1901		
	9439906	Jan 26, 2031		U-2757		
	9439906	Jan 26, 2031		U-2758		
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 003	9439906	Jan 26, 2031		U-1901		
	9439906	Jan 26, 2031		U-2757		
	9439906	Jan 26, 2031		U-2758		
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 004	9439906	Jan 26, 2031		U-1901		
	9439906	Jan 26, 2031		U-2757		
	9439906	Jan 26, 2031		U-2758		
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N 022264 005	9439906	Jan 26, 2031		U-1901		
	9439906	Jan 26, 2031		U-2757		
	9439906	Jan 26, 2031		U-2758		
	9439906	Jan 26, 2031		U-543		
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 001	10143693	Apr 05, 2036		U-2457		
	10143693	Apr 05, 2036		U-2458		

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<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 002	10143693	Apr 05, 2036	U-2457			
	10143693	Apr 05, 2036	U-2458			
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 003	10143693	Apr 05, 2036	U-2457			
	10143693	Apr 05, 2036	U-2458			
<u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u>						
N 207946 004	10143693	Apr 05, 2036	U-2457			
	10143693	Apr 05, 2036	U-2458			
<u>PALIPERIDONE PALMITATE - INVEGA HAFYERA</u>						
N 207946 005	11304951	May 07, 2041	U-3349		NS	Aug 30, 2024
	11324751	May 07, 2041	U-3359			
	11666697	Nov 24, 2041	U-3626			
<u>PALIPERIDONE PALMITATE - INVEGA HAFYERA</u>						
N 207946 006	11304951	May 07, 2041	U-3349		NS	Aug 30, 2024
	11324751	May 07, 2041	U-3359			
	11666697	Nov 24, 2041	U-3626			
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>						
N 021372 001	7947724	Jan 30, 2024	DP			
	7947724*PED	Jul 30, 2024				
	7947725	Jan 30, 2024	DP			
	7947725*PED	Jul 30, 2024				
	7960424	Jan 30, 2024	DP			
	7960424*PED	Jul 30, 2024				
	8518981	Jan 30, 2024	DP			
	8518981*PED	Jul 30, 2024				
	8598218	Jan 30, 2024	DP			
	8598218*PED	Jul 30, 2024				
	8598219	Jan 30, 2024	DP			
	8598219*PED	Jul 30, 2024				
	8729094	Jan 30, 2024	DP U-528			
	8729094*PED	Jul 30, 2024				
	9066980	Jan 30, 2024	DP U-528			
	9066980*PED	Jul 30, 2024				
	9125905	Jan 30, 2024	DP			
	9125905*PED	Jul 30, 2024				
	9173942	Jan 30, 2024	DP			
	9173942*PED	Jul 30, 2024				
	9439854	Jan 30, 2024	DP			
	9439854*PED	Jul 30, 2024				
	9457020	Jan 30, 2024	DP			
	9457020*PED	Jul 30, 2024				
	9457021	Jan 30, 2024	DP			
	9457021*PED	Jul 30, 2024				
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>						
N 021372 002	7947724	Jan 30, 2024	DP			
	7947724*PED	Jul 30, 2024				
	7947725	Jan 30, 2024	DP			

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<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>						
N 021372	002 7947725*PED	Jul 30, 2024				
	7960424	Jan 30, 2024	DP			
	7960424*PED	Jul 30, 2024				
	8518981	Jan 30, 2024	DP			
	8518981*PED	Jul 30, 2024				
	8598218	Jan 30, 2024	DP			
	8598218*PED	Jul 30, 2024				
	9173942	Jan 30, 2024	DP			
	9173942*PED	Jul 30, 2024				
	9439854	Jan 30, 2024	DP			
	9439854*PED	Jul 30, 2024				
	9457020	Jan 30, 2024	DP			
	9457020*PED	Jul 30, 2024				
<u>PALOVAROTENE - SOHONOS</u>						
N 215559	001 10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559	002 10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559	003 10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559	004 10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PALOVAROTENE - SOHONOS</u>						
N 215559	005 10292954	Aug 31, 2031	U-3676		NCE	Aug 16, 2028
	10864194	Jun 08, 2037	U-3676		ODE-439	Aug 16, 2030
	11622959	Jun 08, 2037	U-3676			
	9314439	Aug 31, 2031	U-3676			
	9789074	Aug 31, 2031	U-3676			
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353	001 7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028	U-1669			

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<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353	002 7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028			U-1669	
<u>PANOBINOSTAT LACTATE - FARYDAK</u>						
N 205353	003 7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028			U-1669	
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
N 022020	001 7544370	Jun 07, 2026	DP			
	7550153	Sep 30, 2024			U-859	
	7553498	Sep 30, 2024			U-859	
	7838027	Sep 30, 2024	DP		U-859	
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	001 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	002 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	003 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - PEXEVA</u>						
N 021299	004 7598271	May 04, 2025	DS			
<u>PAROXETINE MESYLATE - BRISDELLE</u>						
N 204516	001 7598271	May 04, 2025	DS			
	8658663	Apr 06, 2029	DS DP		U-904	
	8946251	Aug 04, 2026	DS DP		U-904	
	9393237	Aug 04, 2026			U-904	
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	001 7473761	Dec 14, 2026	DS DP			
	8299209	Dec 27, 2025	DS DP			
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	002 7473761	Dec 14, 2026	DS DP			
	8299209	Dec 27, 2025	DS DP			
<u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u>						
N 200677	003 7473761	Dec 14, 2026	DS DP			
	8299209	Dec 27, 2025	DS DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	001 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	002 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255	003 7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			

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<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255 004	7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u>						
N 203255 005	7473761	Dec 14, 2026	DS DP		ODE-268	Jun 29, 2025
	7759308	Oct 25, 2026	DP			
	9351923	May 23, 2028	DP			
<u>PATIOMER SORBITE X CALCIUM - VELTASSA</u>						
N 205739 001	10485821	Mar 30, 2024		U-1766	NPP	Oct 02, 2026
	11123363	Oct 08, 2033		U-1766		
	7556799	Feb 27, 2025		U-1766		
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027		U-1766		
	8282913	May 29, 2027	DP			
	8287847	Mar 30, 2024		U-1766		
	8337824	May 29, 2030	DS	U-1766		
	8475780	Mar 30, 2024		U-1766		
	8778324	Mar 30, 2024		U-1766		
	8889115	Mar 30, 2024		U-1766		
	9492476	Oct 08, 2033		U-1766		
	9925212	Oct 08, 2033		U-1766		
<u>PATIOMER SORBITE X CALCIUM - VELTASSA</u>						
N 205739 002	10485821	Mar 30, 2024		U-1766	NPP	Oct 02, 2026
	11123363	Oct 08, 2033		U-1766		
	7556799	Feb 27, 2025		U-1766		
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027		U-1766		
	8282913	May 29, 2027	DP			
	8287847	Mar 30, 2024		U-1766		
	8337824	May 29, 2030	DS	U-1766		
	8475780	Mar 30, 2024		U-1766		
	8778324	Mar 30, 2024		U-1766		
	8889115	Mar 30, 2024		U-1766		
	9492476	Oct 08, 2033		U-1766		
	9925212	Oct 08, 2033		U-1766		
<u>PATIOMER SORBITE X CALCIUM - VELTASSA</u>						
N 205739 003	10485821	Mar 30, 2024		U-1766	NPP	Oct 02, 2026
	11123363	Oct 08, 2033		U-1766		
	7556799	Feb 27, 2025		U-1766		
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027		U-1766		
	8282913	May 29, 2027	DP			
	8287847	Mar 30, 2024		U-1766		
	8337824	May 29, 2030	DS	U-1766		
	8475780	Mar 30, 2024		U-1766		
	8778324	Mar 30, 2024		U-1766		
	8889115	Mar 30, 2024		U-1766		
	9492476	Oct 08, 2033		U-1766		

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<u>PATIROMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	003 9925212	Oct 08, 2033	U-1766			
<u>PATIROMER SORBITEX CALCIUM - VELTASSA</u>						
N 205739	004 10485821	Mar 30, 2024	U-1766		NS	Oct 02, 2026
	11123363	Oct 08, 2033	U-1766			
	7556799	Feb 27, 2025	U-1766			
	8147873	Jun 20, 2028	DP			
	8216560	Mar 14, 2027	U-1766			
	8282913	May 29, 2027	DP			
	8287847	Mar 30, 2024	U-1766			
	8337824	May 29, 2030	DS U-1766			
	8475780	Mar 30, 2024	U-1766			
	8778324	Mar 30, 2024	U-1766			
	8889115	Mar 30, 2024	U-1766			
	9492476	Oct 08, 2033	U-1766			
	9925212	Oct 08, 2033	U-1766			
<u>PATISIRAN SODIUM - ONPATTRO</u>						
N 210922	001 10240152	Oct 20, 2029	DS DP U-2378		ODE-197	Aug 10, 2025
	11079379	Aug 27, 2035	DS DP U-2378			
	11141378	Apr 15, 2029	DP			
	8058069	Apr 15, 2029	DP			
	8158601	Nov 10, 2030	DP U-2378			
	8168775	Oct 20, 2029	DS DP U-2378			
	8334373	May 27, 2025	DS DP			
	8492359	Apr 15, 2029	DP			
	8642076	Oct 03, 2027	DP			
	8741866	Oct 20, 2029	U-2378			
	8802644	Oct 21, 2030	DP U-2378			
	8822668	Apr 15, 2029	DP U-2378			
	9234196	Oct 20, 2029	DP U-2378			
	9364435	Apr 15, 2029	DP U-2378			
<u>PEGCETACOPLAN - EMPAVELI</u>						
N 215014	001 10035822	Nov 15, 2033	DS		M-288	Feb 08, 2026
	10125171	Aug 02, 2033	DS		NCE	May 14, 2026
	10875893	Nov 15, 2033	DS U-3124		ODE-351	May 14, 2028
	11040107	Apr 09, 2038	DP U-3172			
	11040107	Apr 09, 2038	DP U-3173			
	11040107	Apr 09, 2038	DP U-3174			
	11292815	Nov 15, 2033	DS DP U-3124			
	11292815	Nov 15, 2033	DS DP U-3354			
	11661441	Jan 13, 2033	DS U-3124			
	7888323	Dec 04, 2027	DS			
	7989589	Dec 04, 2027	DS			
	9169307	Nov 18, 2027	DS U-3123			
<u>PEGCETACOPLAN - SYFOVRE</u>						
N 217171	001 10035822	Nov 15, 2033	DS		NCE	May 14, 2026
	10125171	Aug 02, 2033	DS		NP	Feb 22, 2026
	10875893	Nov 15, 2033	DS U-3540			
	11292815	Nov 15, 2033	DS DP U-3540			

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<u>PEGCETACOPLAN - SYFOVRE</u>						
N 217171	001	11661441	Jan 13, 2033	DS	U-3540	
		7888323	Dec 04, 2027	DS		
		7989589	Dec 04, 2027	DS		
		8168584	Apr 07, 2027		U-3540	
		8168584	Apr 07, 2027		U-3542	
		9056076	Oct 25, 2026		U-3540	
		9169307	Nov 18, 2027	DS	U-3541	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	001	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	002	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	003	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	004	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	005	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	
		7919118	May 12, 2024	DS DP		
		7919461	Jun 02, 2026		U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799	006	7084245	May 12, 2024	DS DP	U-1238	
		7414105	May 12, 2024	DS DP	U-1238	
		7528104	May 12, 2024	DS DP		
		7550433	Jun 02, 2026		U-1238	

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<u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u>						
N 202799 006	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026			U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N 202799 007	7084245	May 12, 2024	DS DP		U-1238	
	7414105	May 12, 2024	DS DP		U-1238	
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026			U-1238	
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026			U-1238	
<u>PEGINESATIDE ACETATE - OMONTYS</u>						
N 202799 008	7084245	May 12, 2024	DS DP		U-1238	
	7414105	May 12, 2024	DS DP		U-1238	
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026			U-1238	
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026			U-1238	
<u>PEMETREXED - PEMFEXY</u>						
N 209472 001	11793813	Feb 19, 2036		DP		
	9604990	Oct 28, 2035	DS			
<u>PEMETREXED DISODIUM - PEMETREXED</u>						
N 215179 001	11147817	Mar 26, 2035		DP		
<u>PEMETREXED DISODIUM - PEMETREXED</u>						
N 215179 002	11147817	Mar 26, 2035		DP		
<u>PEMETREXED DISODIUM - PEMETREXED</u>						
N 215179 003	11147817	Mar 26, 2035		DP		
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736 001	10131667	Jun 12, 2033		U-2809	I-899	Aug 26, 2025
	11466004	May 03, 2039		U-3464	NCE	Apr 17, 2025
	11466004	May 03, 2039		U-3465	ODE-292	Apr 17, 2027
	11466004	May 03, 2039		U-3466	ODE-404	Aug 26, 2029
	11628162	Aug 30, 2040		U-3568		
	11628162	Aug 30, 2040		U-3569		
	11628162	Aug 30, 2040		U-3570		
	11628162	Aug 30, 2040		U-3571		
	9611267	Jan 30, 2035	DS DP			
<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736 002	10131667	Jun 12, 2033		U-2809	I-899	Aug 26, 2025
	11466004	May 03, 2039		U-3464	NCE	Apr 17, 2025
	11466004	May 03, 2039		U-3465	ODE-292	Apr 17, 2027
	11466004	May 03, 2039		U-3466	ODE-404	Aug 26, 2029
	11628162	Aug 30, 2040		U-3568		
	11628162	Aug 30, 2040		U-3569		
	11628162	Aug 30, 2040		U-3570		
	11628162	Aug 30, 2040		U-3571		
	9611267	Jan 30, 2035	DS DP			

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<u>PEMIGATINIB - PEMAZYRE</u>						
N 213736 003	10131667	Jun 12, 2033	U-2809		I-899	Aug 26, 2025
	11466004	May 03, 2039	U-3464		NCE	Apr 17, 2025
	11466004	May 03, 2039	U-3465		ODE-292	Apr 17, 2027
	11466004	May 03, 2039	U-3466		ODE-404	Aug 26, 2029
	11628162	Aug 30, 2040	U-3568			
	11628162	Aug 30, 2040	U-3569			
	11628162	Aug 30, 2040	U-3570			
	11628162	Aug 30, 2040	U-3571			
	9611267	Jan 30, 2035	DS DP			
<u>PENCICLOVIR - PENCICLOVIR</u>						
A 216981 001					CGT	Feb 28, 2024
<u>PERAMIVIR - RAPIVAB</u>						
N 206426 001	10391075	Feb 12, 2027	U-2622			
	10391075	Feb 12, 2027	U-3069			
	8778997	May 07, 2027	U-1627			
	8778997	May 07, 2027	U-2622			
	8778997	May 07, 2027	U-3069			
<u>PERAMPANEL - FYCOMPA</u>						
N 202834 001	6949571	Jun 08, 2024	DS DP U-106			
	6949571	Jun 08, 2024	DS DP U-2088			
	6949571	Jun 08, 2024	DS DP U-2089			
	6949571	Jun 08, 2024	DS DP U-2428			
	6949571	Jun 08, 2024	DS DP U-2429			
	8772497	Jul 01, 2026	DS			
<u>PERAMPANEL - FYCOMPA</u>						
N 202834 002	6949571	Jun 08, 2024	DS DP U-106			
	6949571	Jun 08, 2024	DS DP U-2088			
	6949571	Jun 08, 2024	DS DP U-2089			
	6949571	Jun 08, 2024	DS DP U-2428			
	6949571	Jun 08, 2024	DS DP U-2429			
	8772497	Jul 01, 2026	DS			
<u>PERAMPANEL - FYCOMPA</u>						
N 202834 003	6949571	Jun 08, 2024	DS DP U-106			
	6949571	Jun 08, 2024	DS DP U-2088			
	6949571	Jun 08, 2024	DS DP U-2089			
	6949571	Jun 08, 2024	DS DP U-2428			
	6949571	Jun 08, 2024	DS DP U-2429			
	8772497	Jul 01, 2026	DS			
<u>PERAMPANEL - FYCOMPA</u>						
N 202834 004	6949571	Jun 08, 2024	DS DP U-106			
	6949571	Jun 08, 2024	DS DP U-2088			
	6949571	Jun 08, 2024	DS DP U-2089			
	6949571	Jun 08, 2024	DS DP U-2428			
	6949571	Jun 08, 2024	DS DP U-2429			
	8772497	Jul 01, 2026	DS			

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<u>PERAMPANEL - FYCOMPA</u>						
N 202834	005	6949571	Jun 08, 2024	DS DP U-106		
		6949571	Jun 08, 2024	DS DP U-2088		
		6949571	Jun 08, 2024	DS DP U-2089		
		6949571	Jun 08, 2024	DS DP U-2428		
		6949571	Jun 08, 2024	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 202834	006	6949571	Jun 08, 2024	DS DP U-106		
		6949571	Jun 08, 2024	DS DP U-2088		
		6949571	Jun 08, 2024	DS DP U-2089		
		6949571	Jun 08, 2024	DS DP U-2428		
		6949571	Jun 08, 2024	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERAMPANEL - FYCOMPA</u>						
N 208277	001	6949571	Jun 08, 2024	DS DP U-106		
		6949571	Jun 08, 2024	DS DP U-2088		
		6949571	Jun 08, 2024	DS DP U-2089		
		6949571	Jun 08, 2024	DS DP U-2428		
		6949571	Jun 08, 2024	DS DP U-2429		
		8772497	Jul 01, 2026	DS		
<u>PERFLUOROHEXYLOCTANE - MIEBO</u>						
N 216675	001	10058615	Sep 12, 2033	U-1900	NCE	May 18, 2028
		10369117	Sep 12, 2033	U-1900		
		10449164	Sep 12, 2033	U-1900		
		10507132	Jun 21, 2037	U-1900		
		10576154	Sep 12, 2033	U-1900		
		11357738	Sep 29, 2036	DP		
<u>PERFLUTREN - DEFINITY</u>						
N 021064	001	10583207	Dec 28, 2035	U-665		
		10583208	Mar 16, 2037	U-665		
		10588988	May 04, 2037	U-665		
		11266750	Mar 16, 2037	U-665		
		11529431	Mar 16, 2037	U-665		
		9789210	Mar 16, 2037	U-665		
<u>PERFLUTREN - DEFINITY RT</u>						
N 021064	002	10022460	Dec 28, 2035	DS DP		
		10583207	Dec 28, 2035	U-665		
		10583208	Mar 16, 2037	U-665		
		10588988	May 04, 2037	U-665		
		11266750	Mar 16, 2037	U-665		
		11395856	Dec 28, 2035	DS DP U-665		
		11529431	Mar 16, 2037	U-665		
		9789210	Mar 16, 2037	U-665		
<u>PEXIDARTINIB HYDROCHLORIDE - TURALIO</u>						
N 211810	001	10189833	May 05, 2036	U-2606	NCE	Aug 02, 2024
		10435404	Jul 24, 2038	DP	ODE-250	Aug 02, 2026
		10730876	May 05, 2036	DS		

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<u>PEXIDARTINIB HYDROCHLORIDE - TURALIO</u>						
N 211810 001	10941142	Jul 24, 2038	DP			
	10961240	Jul 24, 2038	U-2606			
	7893075	May 04, 2033	DS			
	8404700	Nov 21, 2027	DS			
	8461169	Apr 19, 2028	U-2606			
	8722702	Nov 21, 2027	DS			
	9169250	Nov 21, 2027	DS			
	9358235	Jun 08, 2033	U-2606			
	9802932	May 05, 2036	DS			
<u>PEXIDARTINIB HYDROCHLORIDE - TURALIO</u>						
N 211810 002	10189833	May 05, 2036	U-2606		NCE	Aug 02, 2024
	10435404	Jul 24, 2038	DP		ODE*	Aug 02, 2026
	10730876	May 05, 2036	DS			
	10941142	Jul 24, 2038	DP			
	10961240	Jul 24, 2038	U-2606			
	7893075	May 04, 2033	DS			
	8404700	Nov 21, 2027	DS			
	8461169	Apr 19, 2028	U-2606			
	8722702	Nov 21, 2027	DS			
	9169250	Nov 21, 2027	DS			
	9358235	Jun 08, 2033	U-2606			
	9802932	May 05, 2036	DS			
<u>PHENOBARBITAL SODIUM - SEZABY</u>						
N 215910 001	11857683	Apr 07, 2042	DP U-3779		ODE-414	Nov 17, 2029
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N 202088 001	8440170	Mar 14, 2029	DP			
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N 202088 002	8440170	Mar 14, 2029	DP			
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N 202088 003	8440170	Mar 14, 2029	DP			
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580 001	8580298	May 15, 2029	DP		NPP	Jun 24, 2025
	8580299	Jun 14, 2029	U-3399			
	8895057	Jun 09, 2028	U-3398			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028	U-3398			
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580 002	8580298	May 15, 2029	DP		NPP	Jun 24, 2025
	8580299	Jun 14, 2029	U-3399			
	8895057	Jun 09, 2028	U-3398			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028	U-3398			
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMIA</u>						
N 022580 003	8580298	May 15, 2029	DP		NPP	Jun 24, 2025

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<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N 022580	003	8580299	Jun 14, 2029	U-3399		
		8895057	Jun 09, 2028	U-3398		
		8895058	Jun 09, 2028	DP		
		9011905	Jun 09, 2028	DP		
		9011906	Jun 09, 2028	U-3398		
<u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u>						
N 022580	004	8580298	May 15, 2029	DP	NPP	Jun 24, 2025
		8580299	Jun 14, 2029	U-3399		
		8895057	Jun 09, 2028	U-3398		
		8895058	Jun 09, 2028	DP		
		9011905	Jun 09, 2028	DP		
		9011906	Jun 09, 2028	U-3398		
<u>PHENTOLAMINE MESYLATE - RYZUMVI</u>						
N 217064	001				NP	Sep 25, 2026
<u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u>						
N 203510	001	8859623	Nov 14, 2033	U-1594		
<u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u>						
N 203510	002	8859623	Nov 14, 2033	U-1594		
<u>PHENYLEPHRINE HYDROCHLORIDE - IMPHENTIV</u>						
N 203826	004	11213480	Sep 26, 2036	DP		
		11471400	Aug 05, 2036	DP		
<u>PHENYLEPHRINE HYDROCHLORIDE - IMPHENTIV</u>						
N 203826	005	11213480	Sep 26, 2036	DP		
		11471400	Aug 05, 2036	DP		
<u>PHENYLEPHRINE HYDROCHLORIDE; TROPICAMIDE - MYDCOMBI</u>						
N 215352	001	10839960	Jul 15, 2031	U-3685	NP	May 05, 2026
		11398306	Jul 15, 2031	U-3685		
<u>PIFLUFOLASTAT F-18 - PYLARIFY</u>						
N 214793	001	10947197	Jun 09, 2037	DS DP U-3130	NCE	May 26, 2026
		8487129	Nov 07, 2027	DS DP		
		8778305	Sep 21, 2030	DS DP U-3130		
		9861713	Jul 31, 2029	DS DP U-3130		
<u>PILOCARPINE HYDROCHLORIDE - VUITY</u>						
N 214028	001	10610518	Apr 24, 2039	U-3252	D-187	Mar 28, 2026
		10610518	Apr 24, 2039	U-3561	NP	Oct 28, 2024
		10610518	Apr 24, 2039	U-3562		
		11285134	Apr 24, 2039	U-3252		
		11285134	Apr 24, 2039	U-3561		
		11285134	Apr 24, 2039	U-3562		
<u>PILOCARPINE HYDROCHLORIDE - OLOSI</u>						
N 217836	001	10639297	Aug 18, 2037	DP U-3741	NP	Oct 17, 2026
		11129812	Aug 18, 2037	U-3741		
		9867810	Aug 18, 2037	DP U-3741		
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318	001	10028944	Jan 15, 2024	U-1974		

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<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318	001	7601740	Apr 29, 2030	DS DP		
		7659285	Aug 24, 2026		U-1844	
		7732615	Jun 03, 2028	DS DP		
		7923564	Sep 26, 2025	DS DP		
		8618130	Jan 15, 2024		U-1845	
		8921393	Jan 15, 2024		U-1846	
		9566271	Jan 15, 2024		U-1974	
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 207318	002	10028944	Jan 15, 2024		U-1974	
		10517860	Mar 23, 2037		U-1974	
		10953000	Mar 23, 2037		U-1974	
		7601740	Apr 29, 2030	DS DP		
		7659285	Aug 24, 2026		U-1844	
		7732615	Jun 03, 2028	DS DP		
		7923564	Sep 26, 2025	DS DP		
		8618130	Jan 15, 2024		U-1845	
		8921393	Jan 15, 2024		U-1846	
		9566271	Jan 15, 2024		U-1974	
<u>PIMAVANSERIN TARTRATE - NUPLAZID</u>						
N 210793	001	10028944	Jan 15, 2024		U-1974	
		10449185	Aug 27, 2038	DP		
		10646480	Aug 27, 2038	DP		
		10849891	Aug 27, 2038	DP	U-1974	
		11452721	Aug 27, 2038	DP		
		7601740	Apr 29, 2030	DS DP		
		7659285	Aug 24, 2026		U-1844	
		7732615	Jun 03, 2028	DS DP		
		7923564	Sep 26, 2025	DS DP		
		8618130	Jan 15, 2024		U-1845	
		8921393	Jan 15, 2024		U-1846	
		9566271	Jan 15, 2024		U-1974	
<u>PIRFENIDONE - ESBRIET</u>						
N 022535	001	7566729	Apr 22, 2029		U-1600	
		7635707	Apr 22, 2029		U-1609	
		7696236	Dec 18, 2027		U-1601	
		7767225	Sep 22, 2026	DP	U-1602	
		7767700	Dec 18, 2027		U-1601	
		7816383	Jan 08, 2030		U-1603	
		7910610	Jan 08, 2030		U-1604	
		7988994	Sep 22, 2026	DP	U-1602	
		8013002	Jan 08, 2030		U-1603	
		8084475	Jan 08, 2030		U-1605	
		8318780	Jan 08, 2030		U-1606	
		8383150	May 10, 2028	DP	U-2361	
		8420674	Dec 18, 2027	DP	U-1608	
		8592462	Apr 22, 2029		U-1609	
		8609701	Apr 22, 2029		U-1610	
		8648098	Jan 08, 2030		U-1611	

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<u>PIRFENIDONE - ESBRIET</u>						
N 022535	001 8753679	Sep 22, 2026	DP U-1602			
	8754109	Jan 08, 2030	U-1612			
	8778947	Aug 30, 2033	U-1613			
<u>PIRFENIDONE - ESBRIET</u>						
N 208780	001 10188637	Mar 28, 2037	DP			
	7566729	Apr 22, 2029	U-2077			
	7566729	Apr 22, 2029	U-2078			
	7635707	Apr 22, 2029	U-2072			
	7635707	Apr 22, 2029	U-2073			
	7635707	Apr 22, 2029	U-2074			
	7635707	Apr 22, 2029	U-2075			
	7635707	Apr 22, 2029	U-2076			
	7635707	Apr 22, 2029	U-2083			
	7767700	Dec 18, 2027	U-2080			
	7816383	Jan 08, 2030	U-2042			
	7816383	Jan 08, 2030	U-2050			
	7910610	Jan 08, 2030	U-2048			
	7910610	Jan 08, 2030	U-2049			
	8013002	Jan 08, 2030	U-2047			
	8013002	Jan 08, 2030	U-2082			
	8084475	Jan 08, 2030	U-2052			
	8084475	Jan 08, 2030	U-2054			
	8318780	Jan 08, 2030	U-2046			
	8318780	Jan 08, 2030	U-2081			
	8383150	May 10, 2028	DP U-2361			
	8420674	Dec 18, 2027	U-2079			
	8592462	Apr 22, 2029	U-2055			
	8592462	Apr 22, 2029	U-2056			
	8592462	Apr 22, 2029	U-2057			
	8592462	Apr 22, 2029	U-2058			
	8592462	Apr 22, 2029	U-2059			
	8592462	Apr 22, 2029	U-2060			
	8592462	Apr 22, 2029	U-2061			
	8592462	Apr 22, 2029	U-2062			
	8592462	Apr 22, 2029	U-2063			
	8609701	Apr 22, 2029	U-2064			
	8609701	Apr 22, 2029	U-2065			
	8609701	Apr 22, 2029	U-2066			
	8609701	Apr 22, 2029	U-2067			
	8609701	Apr 22, 2029	U-2068			
	8609701	Apr 22, 2029	U-2069			
	8609701	Apr 22, 2029	U-2070			
	8648098	Jan 08, 2030	U-2051			
	8648098	Jan 08, 2030	U-2052			
	8754109	Jan 08, 2030	U-2053			
	8778947	Aug 30, 2033	U-2044			
	8778947	Aug 30, 2033	U-2045			

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<u>PIRFENIDONE - ESBRIET</u>						
N 208780	002 10188637	Mar 28, 2037	DP			
	7566729	Apr 22, 2029			U-2269	
	7566729	Apr 22, 2029			U-2270	
	7635707	Apr 22, 2029			U-2072	
	7635707	Apr 22, 2029			U-2073	
	7635707	Apr 22, 2029			U-2074	
	7635707	Apr 22, 2029			U-2075	
	7635707	Apr 22, 2029			U-2076	
	7635707	Apr 22, 2029			U-2083	
	7767700	Dec 18, 2027			U-2080	
	7816383	Jan 08, 2030			U-2042	
	7816383	Jan 08, 2030			U-2050	
	7910610	Jan 08, 2030			U-2048	
	7910610	Jan 08, 2030			U-2049	
	8013002	Jan 08, 2030			U-2047	
	8013002	Jan 08, 2030			U-2082	
	8084475	Jan 08, 2030			U-2054	
	8084475	Jan 08, 2030			U-2268	
	8318780	Jan 08, 2030			U-2046	
	8318780	Jan 08, 2030			U-2081	
	8383150	May 10, 2028	DP		U-2361	
	8420674	Dec 18, 2027			U-2079	
	8592462	Apr 22, 2029			U-2055	
	8592462	Apr 22, 2029			U-2056	
	8592462	Apr 22, 2029			U-2057	
	8592462	Apr 22, 2029			U-2058	
	8592462	Apr 22, 2029			U-2059	
	8592462	Apr 22, 2029			U-2060	
	8592462	Apr 22, 2029			U-2061	
	8592462	Apr 22, 2029			U-2062	
	8592462	Apr 22, 2029			U-2063	
	8609701	Apr 22, 2029			U-2064	
	8609701	Apr 22, 2029			U-2065	
	8609701	Apr 22, 2029			U-2066	
	8609701	Apr 22, 2029			U-2067	
	8609701	Apr 22, 2029			U-2068	
	8609701	Apr 22, 2029			U-2069	
	8609701	Apr 22, 2029			U-2070	
	8648098	Jan 08, 2030			U-2051	
	8648098	Jan 08, 2030			U-2052	
	8754109	Jan 08, 2030			U-2053	
	8778947	Aug 30, 2033			U-2044	
	8778947	Aug 30, 2033			U-2045	
<u>PIRFENIDONE - ESBRIET</u>						
N 208780	003 10188637	Mar 28, 2037	DP			
	7566729	Apr 22, 2029			U-2077	
	7566729	Apr 22, 2029			U-2078	
	7635707	Apr 22, 2029			U-2072	
	7635707	Apr 22, 2029			U-2073	

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<u>PIRFENIDONE - ESBRIET</u>						
N 208780 003	7635707	Apr 22, 2029	U-2074			
	7635707	Apr 22, 2029	U-2075			
	7635707	Apr 22, 2029	U-2076			
	7635707	Apr 22, 2029	U-2083			
	7767700	Dec 18, 2027	U-2080			
	7816383	Jan 08, 2030	U-2042			
	7816383	Jan 08, 2030	U-2050			
	7910610	Jan 08, 2030	U-2048			
	7910610	Jan 08, 2030	U-2049			
	8013002	Jan 08, 2030	U-2047			
	8013002	Jan 08, 2030	U-2082			
	8084475	Jan 08, 2030	U-2052			
	8084475	Jan 08, 2030	U-2054			
	8318780	Jan 08, 2030	U-2046			
	8318780	Jan 08, 2030	U-2081			
	8383150	May 10, 2028	DP U-2361			
	8420674	Dec 18, 2027	U-2079			
	8592462	Apr 22, 2029	U-2055			
	8592462	Apr 22, 2029	U-2056			
	8592462	Apr 22, 2029	U-2057			
	8592462	Apr 22, 2029	U-2058			
	8592462	Apr 22, 2029	U-2059			
	8592462	Apr 22, 2029	U-2060			
	8592462	Apr 22, 2029	U-2061			
	8592462	Apr 22, 2029	U-2062			
	8592462	Apr 22, 2029	U-2063			
	8609701	Apr 22, 2029	U-2064			
	8609701	Apr 22, 2029	U-2065			
	8609701	Apr 22, 2029	U-2066			
	8609701	Apr 22, 2029	U-2067			
	8609701	Apr 22, 2029	U-2068			
	8609701	Apr 22, 2029	U-2069			
	8609701	Apr 22, 2029	U-2070			
	8648098	Jan 08, 2030	U-2051			
	8648098	Jan 08, 2030	U-2052			
	8754109	Jan 08, 2030	U-2053			
	8778947	Aug 30, 2033	U-2044			
	8778947	Aug 30, 2033	U-2045			
<u>PIRTOBRUTINIB - JAYPIRCA</u>						
N 216059 001	10342780	Dec 16, 2036	DS DP		NCE	Jan 27, 2028
	10464905	Dec 16, 2036	U-3518		ODE-424	Jan 27, 2030
	10464905	Dec 16, 2036	U-3761			
	10695323	Dec 16, 2036	DS DP U-3518			
	10695323	Dec 16, 2036	DS DP U-3761			
	10918622	Dec 16, 2036	U-3518			
	10918622	Dec 16, 2036	U-3761			
<u>PIRTOBRUTINIB - JAYPIRCA</u>						
N 216059 002	10342780	Dec 16, 2036	DS DP		NCE	Jan 27, 2028
	10464905	Dec 16, 2036	U-3518		ODE-424	Jan 27, 2030

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<u>PIRTOBRUTINIB - JAYPIRCA</u>						
N 216059	002 10464905	Dec 16, 2036	U-3761			
	10695323	Dec 16, 2036	DS DP U-3518			
	10695323	Dec 16, 2036	DS DP U-3761			
	10918622	Dec 16, 2036	U-3518			
	10918622	Dec 16, 2036	U-3761			
<u>PITAVASTATIN CALCIUM - PITAVASTATIN CALCIUM</u>						
A 206015	001				PC	Feb 19, 2024
<u>PITAVASTATIN CALCIUM - PITAVASTATIN CALCIUM</u>						
A 206015	002				PC	Feb 19, 2024
<u>PITAVASTATIN CALCIUM - PITAVASTATIN CALCIUM</u>						
A 206015	003				PC	Feb 19, 2024
<u>PITAVASTATIN CALCIUM - LIVALO</u>						
N 022363	001 7022713	Feb 19, 2024	U-998			
	7022713*PED	Aug 19, 2024				
	8557993	Feb 02, 2024	DP			
	8557993*PED	Aug 02, 2024				
<u>PITAVASTATIN CALCIUM - LIVALO</u>						
N 022363	002 7022713	Feb 19, 2024	U-998			
	7022713*PED	Aug 19, 2024				
	8557993	Feb 02, 2024	DP			
	8557993*PED	Aug 02, 2024				
<u>PITAVASTATIN CALCIUM - LIVALO</u>						
N 022363	003 7022713	Feb 19, 2024	U-998			
	7022713*PED	Aug 19, 2024				
	8557993	Feb 02, 2024	DP			
	8557993*PED	Aug 02, 2024				
<u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u>						
N 208379	001 8829186	Jan 19, 2031	DS DP			
<u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u>						
N 208379	002 8829186	Jan 19, 2031	DS DP			
<u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u>						
N 208379	003 8829186	Jan 19, 2031	DS DP			
<u>PITOLISANT HYDROCHLORIDE - WAKIX</u>						
N 211150	001 8207197	Mar 07, 2030	DS DP		NCE	Aug 14, 2024
	8354430	Feb 06, 2026	U-1101		ODE-255	Aug 14, 2026
	8354430	Feb 06, 2026	U-1102		ODE-331	Oct 13, 2027
	8486947	Sep 26, 2029	U-1101			
	8486947	Sep 26, 2029	U-1102			
<u>PITOLISANT HYDROCHLORIDE - WAKIX</u>						
N 211150	002 8207197	Mar 07, 2030	DS DP		NCE	Aug 14, 2024
	8354430	Feb 06, 2026	U-1101		ODE-255	Aug 14, 2026
	8354430	Feb 06, 2026	U-1102		ODE-331	Oct 13, 2027
	8486947	Sep 26, 2029	U-1101			
	8486947	Sep 26, 2029	U-1102			

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<u>PLAZOMICIN SULFATE - ZEMDRI</u>						
N 210303 001	8383596	Jun 02, 2031	DS U-2328		NCE	Jun 25, 2023
	8822424	Nov 21, 2028	DP		GAIN	Jun 25, 2028
	9266919	Nov 21, 2028		U-2328		
	9688711	Nov 21, 2028	DS U-2328			
<u>PLECANATIDE - TRULANCE</u>						
N 208745 001	10011637	Jun 05, 2034	DS			
	11142549	Jun 05, 2034	DP			
	11319346	Mar 01, 2032	DP			
	11834521	Jun 05, 2034	DP			
	7041786	Jan 30, 2028	DS			
	9610321	Sep 15, 2031		U-1999		
	9610321	Sep 15, 2031		U-2230		
	9616097	Aug 20, 2032	DP			
	9919024	Sep 15, 2031		U-1999		
	9919024	Sep 15, 2031		U-2230		
	9925231	Sep 15, 2031	DP			
<u>PODOFILOX - PODOFILOX</u>						
A 211871 001					CGT	Jun 10, 2024
<u>POLIDOCANOL - VARITHENA</u>						
N 205098 001	7731986	Nov 17, 2024	DS DP U-1463			
	7814943	Nov 19, 2027	DP U-1461			
	8122917	Sep 09, 2024	DP			
	9480652	May 12, 2032	DP			
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 001	10555939	May 19, 2030	DP		M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025		U-1360	ODE-297	May 14, 2027
	8198262	Jun 17, 2025		U-2254	PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8828427	Jun 21, 2031	DS DP		PED	Nov 14, 2027
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 002	10555939	May 19, 2030	DP		M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025		U-1360	ODE-297	May 14, 2027
	8198262	Jun 17, 2025		U-2254	PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8828427	Jun 21, 2031	DS DP		PED	Nov 14, 2027
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 003	10555939	May 19, 2030	DP		M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025		U-1360	ODE-297	May 14, 2027

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<u>POMALIDOMIDE - POMALYST</u>						
N 204026 003	8198262	Jun 17, 2025	U-2254		PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8828427	Jun 21, 2031	DS DP		PED	Nov 14, 2027
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>POMALIDOMIDE - POMALYST</u>						
N 204026 004	10555939	May 19, 2030	DP		M-14	Nov 20, 2023
	10555939*PED	Nov 19, 2030			ODE-296	May 14, 2027
	8198262	Jun 17, 2025	U-1360		ODE-297	May 14, 2027
	8198262	Jun 17, 2025	U-2254		PED	May 20, 2024
	8198262*PED	Dec 17, 2025			PED	Nov 14, 2027
	8828427	Jun 21, 2031	DS DP		PED	Nov 14, 2027
	8828427*PED	Dec 21, 2031				
	9993467	May 19, 2030	DP			
	9993467*PED	Nov 19, 2030				
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469 001	11192895	Dec 12, 2033	U-1700			
	11192895	Dec 12, 2033	U-1701			
	11192895	Dec 12, 2033	U-1948			
	11192897	Dec 12, 2033	DS U-1700			
	11192897	Dec 12, 2033	DS U-1701			
	11192897	Dec 12, 2033	DS U-1948			
	11384086	Dec 12, 2033	DS DP U-1700			
	11384086	Dec 12, 2033	DS DP U-1701			
	11384086	Dec 12, 2033	DS DP U-1948			
	8114874	Jan 24, 2027	DS DP			
	9029533	Dec 22, 2026	U-1283			
	9029533	Dec 22, 2026	U-1699			
	9029533	Dec 22, 2026	U-1700			
	9029533	Dec 22, 2026	U-1701			
	9029533	Dec 22, 2026	U-836			
	9493470	Dec 12, 2033	DS DP U-1700			
	9493470	Dec 12, 2033	DS DP U-1948			
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469 002	11192895	Dec 12, 2033	U-1700			
	11192895	Dec 12, 2033	U-1701			
	11192895	Dec 12, 2033	U-1948			
	11192897	Dec 12, 2033	DS U-1700			
	11192897	Dec 12, 2033	DS U-1701			
	11192897	Dec 12, 2033	DS U-1948			
	11384086	Dec 12, 2033	DS DP U-1700			
	11384086	Dec 12, 2033	DS DP U-1701			
	11384086	Dec 12, 2033	DS DP U-1948			
	8114874	Jan 24, 2027	DS DP			
	9029533	Dec 22, 2026	U-1283			
	9029533	Dec 22, 2026	U-1699			
	9029533	Dec 22, 2026	U-1700			

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<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469	002	9029533	Dec 22, 2026		U-1701	
		9029533	Dec 22, 2026		U-836	
		9493470	Dec 12, 2033	DS DP	U-1700	
		9493470	Dec 12, 2033	DS DP	U-1948	
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469	003	11192895	Dec 12, 2033		U-1700	
		11192895	Dec 12, 2033		U-1701	
		11192895	Dec 12, 2033		U-1948	
		11192897	Dec 12, 2033	DS	U-1700	
		11192897	Dec 12, 2033	DS	U-1701	
		11192897	Dec 12, 2033	DS	U-1948	
		11384086	Dec 12, 2033	DS DP	U-1700	
		11384086	Dec 12, 2033	DS DP	U-1701	
		11384086	Dec 12, 2033	DS DP	U-1948	
		8114874	Jan 24, 2027	DS DP		
		9029533	Dec 22, 2026		U-1283	
		9029533	Dec 22, 2026		U-1699	
		9029533	Dec 22, 2026		U-1700	
		9029533	Dec 22, 2026		U-1701	
		9029533	Dec 22, 2026		U-836	
		9493470	Dec 12, 2033	DS DP	U-1700	
		9493470	Dec 12, 2033	DS DP	U-1948	
<u>PONATINIB HYDROCHLORIDE - ICLUSIG</u>						
N 203469	004	11192895	Dec 12, 2033		U-1700	
		11192895	Dec 12, 2033		U-1701	
		11192895	Dec 12, 2033		U-1948	
		11192897	Dec 12, 2033	DS	U-1700	
		11192897	Dec 12, 2033	DS	U-1701	
		11192897	Dec 12, 2033	DS	U-1948	
		11384086	Dec 12, 2033	DS DP	U-1700	
		11384086	Dec 12, 2033	DS DP	U-1701	
		11384086	Dec 12, 2033	DS DP	U-1948	
		8114874	Jan 24, 2027	DS DP		
		9029533	Dec 22, 2026		U-1283	
		9029533	Dec 22, 2026		U-1699	
		9029533	Dec 22, 2026		U-1700	
		9029533	Dec 22, 2026		U-1701	
		9029533	Dec 22, 2026		U-836	
		9493470	Dec 12, 2033	DS DP	U-1700	
		9493470	Dec 12, 2033	DS DP	U-1948	
<u>PONESIMOD - PONVORY</u>						
N 213498	001	10220023	Dec 10, 2035		U-3103	
		8273779	Dec 17, 2025		U-2774	
		9000018	Nov 16, 2024		U-3102	
		9062014	May 06, 2032	DS DP	U-2774	
		RE43728	Nov 16, 2024	DS DP		
					NCE	Mar 18, 2026

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<u>PONESIMOD - PONVORY</u>						
N 213498	002	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	003	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	004	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	005	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	006	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	007	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	008	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		
<u>PONESIMOD - PONVORY</u>						
N 213498	009	10220023	Dec 10, 2035	U-3103	NCE	Mar 18, 2026
		8273779	Dec 17, 2025	U-2774		
		9000018	Nov 16, 2024	U-3102		
		9062014	May 06, 2032	DS DP U-2774		
		RE43728	Nov 16, 2024	DS DP		

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<u>PONESIMOD - PONVORY</u>						
N 213498 009	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	8273779	Dec 17, 2025	U-2774			
	9000018	Nov 16, 2024	U-3102			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2024	DS DP			
<u>PONESIMOD - PONVORY</u>						
N 213498 010	10220023	Dec 10, 2035	U-3103		NCE	Mar 18, 2026
	8273779	Dec 17, 2025	U-2774			
	9000018	Nov 16, 2024	U-3102			
	9062014	May 06, 2032	DS DP U-2774			
	RE43728	Nov 16, 2024	DS DP			
<u>POSACONAZOLE - NOXAFIL</u>						
N 205053 001					I-881	Jun 17, 2024
					NPP	May 31, 2024
					ODE-355	Jun 17, 2028
<u>POSACONAZOLE - NOXAFIL</u>						
N 205596 001	10117951	Mar 13, 2029	DP		I-881	Jun 17, 2024
	8410077	Mar 13, 2029	DP		NPP	May 31, 2024
	9023790	Jul 04, 2031	DP U-1698		ODE-355	Jun 17, 2028
	9023790	Jul 04, 2031	DP U-3160			
	9023790	Jul 04, 2031	DP U-3171			
	9358297	Jun 24, 2031	DP U-3160			
	9358297	Jun 24, 2031	DP U-3171			
	9493582	Feb 27, 2033	DP			
	9750822	Mar 13, 2029	DP			
<u>POSACONAZOLE - NOXAFIL POWDERMIX KIT</u>						
N 214770 001					NP	May 31, 2024
<u>POTASSIUM CHLORIDE - POTASSIUM CHLORIDE</u>						
A 217704 001					CGT	Mar 11, 2024
<u>POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC - POTASSIUM PHOSPHATES</u>						
A 216274 001					CGT	Apr 13, 2024
<u>POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC - POTASSIUM PHOSPHATES</u>						
A 216274 002					CGT	Apr 10, 2024
<u>POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC - POTASSIUM PHOSPHATES</u>						
A 216274 003					CGT	Apr 10, 2024
<u>POTASSIUM PHOSPHATE, DIBASIC; POTASSIUM PHOSPHATE, MONOBASIC - POTASSIUM PHOSPHATES</u>						
N 212121 001	10632150	Apr 19, 2039	DP U-2789			
<u>PRALATREXATE - FOLOTYN</u>						
N 022468 001	7622470	May 31, 2025	U-1015			
	8299078	May 31, 2025	U-1004			
<u>PRALATREXATE - FOLOTYN</u>						
N 022468 002	7622470	May 31, 2025	U-1015			
	8299078	May 31, 2025	U-1004			

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<u>PRALSETINIB - GAVRETO</u>						
N 213721	001	10030005	Nov 01, 2036	DS DP U-2952	NCE	Sep 04, 2025
		10030005	Nov 01, 2036	DS DP U-3002	ODE-318	Sep 04, 2027
		11273160	Apr 03, 2039	U-2952	ODE-340	Dec 01, 2027
					ODE-341	Dec 01, 2027
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421	001	7695734	Apr 26, 2028	DP		
		8679533	Sep 08, 2029	DP U-219		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421	002	7695734	Apr 26, 2028	DP		
		8679533	Sep 08, 2029	DP U-219		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421	003	7695734	Apr 26, 2028	DP		
		8679533	Sep 08, 2029	DP U-219		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421	004	7695734	Apr 26, 2028	DP		
		8679533	Sep 08, 2029	DP U-219		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421	005	7695734	Apr 26, 2028	DP		
		8679533	Sep 08, 2029	DP U-219		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421	006	7695734	Apr 26, 2028	DP		
		8679533	Sep 08, 2029	DP U-219		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N 022421	007	7695734	Apr 26, 2028	DP		
		8679533	Sep 08, 2029	DP U-219		
<u>PRASTERONE - INTRAROSA</u>						
N 208470	001	8268806	Mar 19, 2031	DP		
		8629129	Aug 07, 2028	DP		
		8957054	Jan 08, 2030	U-1922		
<u>PREDNISOLONE ACETATE - FLO-PRED</u>						
N 022067	001	7799331	Oct 11, 2028	DP U-1068		
		7799331	Oct 11, 2028	DP U-139		
<u>PREDNISOLONE ACETATE - FLO-PRED</u>						
N 022067	002	7799331	Oct 11, 2028	DP U-1068		
		7799331	Oct 11, 2028	DP U-139		
<u>PREDNISONONE - RAYOS</u>						
N 202020	001	8309124	Apr 23, 2024	U-1292		
		8394407	Apr 23, 2024	DP U-1362		
		9040085	Apr 23, 2024	U-1362		
		9186332	Apr 23, 2024	U-1362		
		9504699	Aug 03, 2027	U-1362		
<u>PREDNISONONE - RAYOS</u>						
N 202020	002	8309124	Apr 23, 2024			
		8394407	Apr 23, 2024	DP U-1362		
		9040085	Apr 23, 2024	U-1362		

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<u>PREDNISONONE - RAYOS</u>						
N 202020 002	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<u>PREDNISONONE - RAYOS</u>						
N 202020 003	8168218	Jan 07, 2028	DP U-1269			
	8309124	Apr 23, 2024	U-1292			
	8394407	Apr 23, 2024	DP U-1362			
	9040085	Apr 23, 2024	U-1362			
	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<u>PREGABALIN - LYRICA CR</u>						
N 209501 001	10022447	Nov 02, 2026	U-2136			
	10022447	Nov 02, 2026	U-2137			
	10022447*PED	May 02, 2027				
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	8945620*PED	May 02, 2027				
	9144559	Nov 02, 2026	DP			
	9144559*PED	May 02, 2027				
<u>PREGABALIN - LYRICA CR</u>						
N 209501 002	10022447	Nov 02, 2026	U-2136			
	10022447	Nov 02, 2026	U-2137			
	10022447*PED	May 02, 2027				
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	8945620*PED	May 02, 2027				
	9144559	Nov 02, 2026	DP			
	9144559*PED	May 02, 2027				
<u>PREGABALIN - LYRICA CR</u>						
N 209501 003	10022447	Nov 02, 2026	U-2136			
	10022447	Nov 02, 2026	U-2137			
	10022447*PED	May 02, 2027				
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	8945620*PED	May 02, 2027				
	9144559	Nov 02, 2026	DP			
	9144559*PED	May 02, 2027				
<u>PRETOMANID - PRETOMANID</u>						
N 212862 001					NCE	Aug 14, 2024
					ODE-253	Aug 14, 2026
					GAIN	Aug 14, 2029
<u>PROGESTERONE - MILPROSA</u>						
N 201110 001	10537584	Feb 03, 2029	DP			
	10548904	Feb 03, 2029	U-2810			
	8580293	Jan 21, 2030	U-2810			
<u>PROPOFOL - DIPRIVAN</u>						
N 019627 002	8476010	Dec 01, 2024	DS DP			
	8476010*PED	Jun 01, 2025				

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<u>PROPOFOL - DIPRIVAN</u>						
N 019627	002 8476010	Dec 01, 2024	DS DP			
	8476010*PED	Jun 01, 2025				
<u>PROPRANOLOL HYDROCHLORIDE - HEMANGEOL</u>						
N 205410	001 8338489	Oct 16, 2028		U-1496		
	8987262	Oct 16, 2028		U-1988		
<u>QUAZEPAM - DORAL</u>						
N 018708	001 7608616	Jun 03, 2028		U-1012		
<u>QUAZEPAM - DORAL</u>						
N 018708	003 7608616	Jun 03, 2028		U-1012		
<u>QUIZARTINIB DIHYDROCHLORIDE - VANFLYTA</u>						
N 216993	001 7820657	Sep 26, 2028	DS		NCE	Jul 20, 2028
	7968543	Aug 15, 2029	DP U-3661		ODE-437	Jul 20, 2030
	8129374	Mar 16, 2027	U-3661			
	8357690	Feb 26, 2031	U-3661			
	8557810	Mar 16, 2027	DP			
	8836218	Mar 23, 2030	U-3661			
	8865710	Aug 15, 2029	DP			
	8883783	Mar 16, 2027	DS			
	9555040	May 14, 2030	U-3661			
	9585892	Mar 16, 2027	U-3661			
	9675549	Sep 30, 2033	DP			
<u>QUIZARTINIB DIHYDROCHLORIDE - VANFLYTA</u>						
N 216993	002 7820657	Sep 26, 2028	DS		NCE	Jul 20, 2028
	7968543	Aug 15, 2029	DP U-3661		ODE-437	Jul 20, 2030
	8129374	Mar 16, 2027	U-3661			
	8357690	Feb 26, 2031	U-3661			
	8557810	Mar 16, 2027	DP			
	8836218	Mar 23, 2030	U-3661			
	8865710	Aug 15, 2029	DP			
	8883783	Mar 16, 2027	DS			
	9555040	May 14, 2030	U-3661			
	9585892	Mar 16, 2027	U-3661			
	9675549	Sep 30, 2033	DP			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 022145	001 7169780*PED	Apr 03, 2024				
	7754731	Mar 11, 2029	DS DP U-257			
	7754731*PED	Sep 11, 2029				
	8771733	Jun 02, 2030	DS DP U-257			
	8852632	Jan 28, 2028	DS DP U-257			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS HD</u>						
N 022145	002 10772888	Mar 30, 2032		U-1663		
	7169780*PED	Apr 03, 2024				
	7754731	Mar 11, 2029	DS DP U-257			
	7754731*PED	Sep 11, 2029				
	8771733	Jun 02, 2030	DS DP U-257			
	9649311	Oct 21, 2030	DP			
	9649311*PED	Apr 21, 2031				

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<u>RALTEGRAVIR POTASSIUM - ISENTRESS HD</u>						
N 022145	002	10772888	Mar 30, 2032		U-1663	
		7169780*PED	Apr 03, 2024			
		7754731	Mar 11, 2029	DS DP	U-257	
		7754731*PED	Sep 11, 2029			
		8771733	Jun 02, 2030	DS DP	U-257	
		9649311	Oct 21, 2030	DP		
		9649311*PED	Apr 21, 2031			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045	001	7169780*PED	Apr 03, 2024			
		7754731	Mar 11, 2029	DS DP	U-257	
		7754731*PED	Sep 11, 2029			
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 203045	002	7169780*PED	Apr 03, 2024			
		7754731	Mar 11, 2029	DS DP	U-257	
		7754731*PED	Sep 11, 2029			
		8771733	Jun 02, 2030	DS DP	U-257	
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>						
N 205786	001	7169780*PED	Apr 03, 2024			
		7754731	Mar 11, 2029	DS DP	U-257	
		7754731*PED	Sep 11, 2029			
<u>RANOLAZINE - ASPRUZYO SPRINKLE</u>						
N 216018	001	11510878	Jan 24, 2038	DP		
<u>RANOLAZINE - ASPRUZYO SPRINKLE</u>						
N 216018	002	10898444	Jan 24, 2038	DP		
		11510878	Jan 24, 2038	DP		
<u>RASAGILINE MESYLATE - AZILECT</u>						
N 021641	001	7572834	Dec 05, 2026	DP		
		7815942	Aug 27, 2027	DS DP	U-219	
<u>RASAGILINE MESYLATE - AZILECT</u>						
N 021641	002	7572834	Dec 05, 2026	DP		
		7815942	Aug 27, 2027	DS DP	U-219	
<u>REGADENOSON - LEXISCAN</u>						
N 022161	001	8106183	Feb 02, 2027	DS		
		RE47301	Feb 02, 2027	DP		
<u>REGORAFENIB - STIVARGA</u>						
N 203085	001	8637553	Feb 16, 2031	DS DP		ODE-139
		8680124	Jun 02, 2030		U-1506	
		9458107	Apr 08, 2031	DP		
		9957232	Jul 09, 2032	DS		
<u>RELUGOLIX - ORGOVYX</u>						
N 214621	001	10350170	Feb 25, 2036	DP		NCE
		10449191	Sep 29, 2037		U-3020	Dec 18, 2025
		10786501	Sep 29, 2037		U-3020	
		11583526	Sep 29, 2037		U-3020	
		11795178	Sep 27, 2033	DS DP		
		7300935	Jan 28, 2025	DS		

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<u>RELUGOLIX - ORGOVYX</u>						
N 214621	001 8058280	Jan 28, 2025	DS DP			
	8735401	Feb 04, 2024		U-3019		
<u>REMEDESIVIR - VEKLURY</u>						
N 214787	001 10065958	Sep 16, 2031	DS		D-183	Jan 21, 2025
	10675296	Jul 10, 2038	DP		M-301	Jul 13, 2026
	10695361	Sep 16, 2036		U-2984	NCE	Oct 22, 2025
	10695361	Sep 16, 2036		U-3249	NPP	Apr 25, 2025
	10695361	Sep 16, 2036		U-3367		
	10695361	Sep 16, 2036		U-3368		
	11007208	Sep 16, 2036		U-2984		
	11007208	Sep 16, 2036		U-3249		
	11007208	Sep 16, 2036		U-3367		
	11007208	Sep 16, 2036		U-3368		
	11266681	Jul 10, 2038		U-2984		
	11266681	Jul 10, 2038		U-3249		
	11266681	Jul 10, 2038		U-3367		
	11266681	Jul 10, 2038		U-3368		
	11382926	Sep 16, 2036		U-3367		
	11382926	Sep 16, 2036		U-3368		
	11491169	May 28, 2041		U-3484		
	11491169	May 28, 2041		U-3485		
	11492353	Dec 08, 2031	DS			
	8008264	Sep 06, 2029	DS DP			
	8318682	Apr 22, 2029	DS DP			
	9724360	Oct 29, 2035	DS DP			
	9949994	Oct 29, 2035	DS			
	RE46762	Apr 22, 2029	DS DP			
<u>REMEDESIVIR - VEKLURY</u>						
N 214787	002 10065958	Sep 16, 2031	DS		D-183	Jan 21, 2025
	10675296	Jul 10, 2038	DP		M-301	Jul 13, 2026
	10695361	Sep 16, 2036		U-2984	NCE	Oct 22, 2025
	10695361	Sep 16, 2036		U-3249	NPP	Apr 25, 2025
	10695361	Sep 16, 2036		U-3367		
	10695361	Sep 16, 2036		U-3368		
	11007208	Sep 16, 2036		U-2984		
	11007208	Sep 16, 2036		U-3249		
	11007208	Sep 16, 2036		U-3367		
	11007208	Sep 16, 2036		U-3368		
	11266681	Jul 10, 2038		U-2984		
	11266681	Jul 10, 2038		U-3249		
	11266681	Jul 10, 2038		U-3367		
	11266681	Jul 10, 2038		U-3368		
	11382926	Sep 16, 2036		U-3367		
	11382926	Sep 16, 2036		U-3368		
	11491169	May 28, 2041		U-3484		
	11491169	May 28, 2041		U-3485		
	11492353	Dec 08, 2031	DS			
	8008264	Sep 06, 2029	DS DP			
	8318682	Apr 22, 2029	DS DP			

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<u>REMEDESIVIR - VEKLURY</u>						
N 214787	002 9724360	Oct 29, 2035	DS DP			
	9949994	Oct 29, 2035	DS			
	RE46762	Apr 22, 2029	DS DP			
<u>REMIMAZOLAM BESYLATE - BYFAVO</u>						
N 212295	001 10052334	Nov 07, 2031		U-2968	NCE	Oct 06, 2025
	10195210	Nov 07, 2031		U-2968		
	10342800	Nov 07, 2031		U-2968		
	10472365	Jul 10, 2027		U-2968		
	10722522	Nov 07, 2031		U-2968		
	10961250	Jul 10, 2027	DP	U-2968		
	9561236	Apr 30, 2033		U-2968		
	9737547	Nov 07, 2031		U-2968		
	9777007	Jul 10, 2027	DP			
	9827251	Nov 07, 2031		U-2968		
	9914738	Jul 10, 2027	DP			
<u>REPOTRECTINIB - AUGTYRO</u>						
N 218213	001 10294242	Jul 05, 2036	DS			
	11452725	Jul 24, 2036		U-3755		
	9714258	Jan 23, 2035	DS DP			
<u>RETAPAMULIN - ALTABAX</u>						
N 022055	001 7875630	Feb 14, 2027	DS			
	8207191	Aug 30, 2024		U-805		
<u>REVEFENACIN - YUPELRI</u>						
N 210598	001 10106503	Mar 10, 2025		U-2440		
	10343995	Mar 10, 2025		U-2440		
	10550081	Jul 14, 2030	DS			
	11008289	Jul 14, 2030		U-2440		
	11247969	Mar 10, 2025	DP			
	11484531	Oct 23, 2039		U-2440		
	11691948	Jul 14, 2030	DP			
	11858898	Jul 14, 2030	DS DP	U-2440		
	7288657	Dec 23, 2025	DS			
	7491736	Mar 10, 2025		U-2440		
	7521041	Mar 10, 2025		U-2440		
	7550595	Mar 10, 2025	DP			
	7585879	Mar 10, 2025	DS DP	U-2440		
	7910608	Mar 10, 2025	DS DP			
	8034946	Mar 10, 2025	DP			
	8053448	Mar 10, 2025		U-2440		
	8273894	Mar 10, 2025	DP			
	8541451	Aug 25, 2031	DS			
	9765028	Jul 14, 2030	DS			
<u>REZAFUNGIN ACETATE - REZZAYO</u>						
N 217417	001 10702573	Mar 14, 2033	DS DP	U-3566	NCE	Mar 22, 2028
	11197909	Jul 14, 2038	DS DP	U-3566	ODE-426	Mar 22, 2030
	11654196	Mar 02, 2032	DS DP	U-3566	GAIN	Mar 22, 2033
	11712459	Mar 15, 2037	DP	U-3566	GAIN	Mar 22, 2035
	11819533	Jul 11, 2038	DS DP	U-3566		

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<u>REZAFUNGIN ACETATE - REZZAYO</u>						
N 217417	001 8722619	Mar 02, 2032	DS DP U-3566			
	9526835	Mar 14, 2033	DS DP U-3566			
<u>RIBOCICLIB SUCCINATE - KISOALI</u>						
N 209092	001 10799506	Apr 14, 2036	DP		NPP	Dec 10, 2024
	8324225	Jun 17, 2028	DS DP			
	8415355	Mar 13, 2031	DS DP			
	8685980	May 25, 2030	DS DP			
	8962630	Dec 09, 2029	U-1981			
	8962630	Dec 09, 2029	U-2355			
	8962630	Dec 09, 2029	U-2356			
	8962630	Dec 09, 2029	U-3265			
	8962630	Dec 09, 2029	U-3266			
	9193732	Nov 09, 2031	DS DP			
	9416136	Aug 20, 2029	U-1981			
	9416136	Aug 20, 2029	U-2355			
	9416136	Aug 20, 2029	U-2356			
	9416136	Aug 20, 2029	U-3265			
	9416136	Aug 20, 2029	U-3266			
	9868739	Nov 09, 2031	U-1981			
	9868739	Nov 09, 2031	U-2355			
	9868739	Nov 09, 2031	U-2356			
	9868739	Nov 09, 2031	U-3265			
	9868739	Nov 09, 2031	U-3266			
<u>RIFAMYCIN SODIUM - AEMCOLO</u>						
N 210910	001 8263120	May 03, 2025	DP		NCE	Nov 16, 2023
	8486446	May 03, 2025	DP		GAIN	Nov 16, 2028
	8529945	May 03, 2025	DP			
	8741948	May 03, 2025	DP U-2448			
<u>RIFAXIMIN - XIFAXAN</u>						
N 021361	001 10703763	Feb 27, 2026	U-1708			
	10703763	Feb 27, 2026	U-2847			
	10703763	Feb 27, 2026	U-2848			
	7045620	Jun 19, 2024	DS DP			
	7612199	Jun 19, 2024	DS DP			
	7902206	Jun 19, 2024	DS DP			
	7906542	Jun 01, 2025	DS DP			
	7928115	Jul 24, 2029	U-1121			
	8158644	Jun 19, 2024	DP			
	8158781	Jun 19, 2024	DS			
	8193196	Sep 02, 2027	DS DP			
	8518949	Feb 27, 2026	DP			
	8741904	Feb 27, 2026	DS U-1526			
	8835452	Jun 19, 2024	DS DP			
	8853231	Jun 19, 2024	DP			
	9271968	Feb 27, 2026	DP			
<u>RIFAXIMIN - XIFAXAN</u>						
N 021361	002 10314828	Jul 24, 2029	U-1481			
	10335397	Jul 24, 2029	U-2579			

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<u>RIFAXIMIN - XIFAXAN</u>						
N 021361 002	10456384	Feb 26, 2029				U-2643
	10456384	Feb 26, 2029				U-2644
	10703763	Feb 27, 2026				U-1708
	10703763	Feb 27, 2026				U-2847
	10703763	Feb 27, 2026				U-2848
	10709694	Jul 24, 2029				U-2579
	10765667	Feb 26, 2029				U-2643
	10765667	Feb 26, 2029				U-2644
	11564912	Feb 26, 2029				U-3511
	11564912	Feb 26, 2029				U-3512
	11779571	Feb 26, 2029				U-3706
	7045620	Jun 19, 2024	DS			
	7612199	Jun 19, 2024	DS DP			
	7902206	Jun 19, 2024	DS DP			
	7906542	Jun 01, 2025	DS DP			
	7915275	Feb 23, 2025				U-1707
	7915275	Feb 23, 2025				U-1708
	8158644	Jun 19, 2024	DP			
	8158781	Jun 19, 2024	DS			
	8193196	Sep 02, 2027	DS DP			U-1707
	8193196	Sep 02, 2027	DS DP			U-1708
	8309569	Jul 18, 2029				U-1707
	8309569	Jul 18, 2029				U-1708
	8518949	Feb 27, 2026	DP			
	8642573	Oct 02, 2029				U-1481
	8741904	Feb 27, 2026	DS			U-1526
	8741904	Feb 27, 2026	DS			U-1707
	8741904	Feb 27, 2026	DS			U-1708
	8829017	Jul 24, 2029				U-1562
	8835452	Jun 19, 2024	DS DP			
	8853231	Jun 19, 2024	DP			
	8946252	Jul 24, 2029				U-1481
	8969398	Oct 02, 2029				U-1481
	9271968	Feb 27, 2026	DP			
	9421195	Jul 24, 2029				U-1481
	9629828	Jul 24, 2029				U-1994
<u>RILPIVIRINE HYDROCHLORIDE - EDURANT</u>						
N 202022 001	7125879	Apr 21, 2025	DS DP			U-1153
	7125879	Apr 21, 2025	DS DP			U-1307
	7125879	Apr 21, 2025	DS DP			U-1740
	7125879	Apr 21, 2025	DS DP			U-3353
<u>RILUZOLE - TIGLUTIK KIT</u>						
N 209080 001	8765150	Mar 12, 2029	DP			U-2401
<u>RILUZOLE - EXSERVAN</u>						
N 212640 001	8603514	Apr 03, 2024	DP			
	8765167	Feb 20, 2024	DP			

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<u>RIMEGEPANT SULFATE - NURTEC ODT</u>						
N 212728	001	11083724	Mar 25, 2039	DP U-2718	I-865	May 27, 2024
		11083724	Mar 25, 2039	DP U-3142	NCE	Feb 27, 2025
		8314117	Mar 09, 2030	DS DP U-2718		
		8314117	Mar 09, 2030	DS DP U-3142		
		8759372	Feb 25, 2033	DS DP		
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	001	10662188	Feb 18, 2034	DS DP U-2834		
		10662188	Feb 18, 2034	DS DP U-2835		
		11203593	Feb 18, 2034	DS DP U-2834		
		11203593	Feb 18, 2034	DS DP U-2835		
		7173037	Dec 04, 2026	DS DP		
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	002	10662188	Feb 18, 2034	DS DP U-2834		
		10662188	Feb 18, 2034	DS DP U-2835		
		11203593	Feb 18, 2034	DS DP U-2834		
		11203593	Feb 18, 2034	DS DP U-2835		
		7173037	Dec 04, 2026	DS DP		
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	003	10662188	Feb 18, 2034	DS DP U-2834		
		10662188	Feb 18, 2034	DS DP U-2835		
		11203593	Feb 18, 2034	DS DP U-2834		
		11203593	Feb 18, 2034	DS DP U-2835		
		7173037	Dec 04, 2026	DS DP		
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	004	10662188	Feb 18, 2034	DS DP U-2834		
		10662188	Feb 18, 2034	DS DP U-2835		
		11203593	Feb 18, 2034	DS DP U-2834		
		11203593	Feb 18, 2034	DS DP U-2835		
		7173037	Dec 04, 2026	DS DP		
<u>RIOCIGUAT - ADEMPAS</u>						
N 204819	005	10662188	Feb 18, 2034	DS DP U-2834		
		10662188	Feb 18, 2034	DS DP U-2835		
		11203593	Feb 18, 2034	DS DP U-2834		
		11203593	Feb 18, 2034	DS DP U-2835		
		7173037	Dec 04, 2026	DS DP		
<u>RIPRETINIB - QINLOCK</u>						
N 213973	001	10966966	Aug 12, 2040	U-3153	NCE	May 15, 2025
		11185535	Dec 30, 2040	DP	ODE-298	May 15, 2027
		11266635	Aug 12, 2040	U-3330		
		11344536	Aug 12, 2040	U-3381		
		11395818	Dec 30, 2040	DP		
		11426390	Aug 12, 2040	U-3416		
		11433056	Aug 12, 2040	U-3423		
		11529336	Aug 12, 2040	U-3382		
		11534432	Aug 12, 2040	U-3442		
		11576903	Dec 30, 2040	DP		
		11576904	Aug 12, 2040	U-3537		

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<u>RIPRETINIB - QINLOCK</u>						
N 213973	001	11612591	Dec 30, 2040	DP		
		11779572	Oct 06, 2042	U-3714		
		11793795	Dec 30, 2040	DP		
		11801237	Dec 30, 2040	DP U-3219		
		11813251	Aug 12, 2040	U-3750		
		11844788	Dec 30, 2040	DP		
		11850240	Dec 30, 2040	DP		
		11850241	Dec 30, 2040	DP		
		8188113	Jul 27, 2030	DS DP		
		8461179	May 15, 2034	DS DP		
		RE48731	Jun 07, 2032	U-3219		
<u>RISDIPLAM - EVRYSDI</u>						
N 213535	001	11534444	Oct 04, 2038	U-1943	M-270	Oct 03, 2026
		11827646	Jan 25, 2036	U-1943	NCE	Aug 07, 2025
		9586955	Feb 08, 2033	DS DP	NPP	May 27, 2025
		9969754	May 11, 2035	DS DP U-1943	ODE-334	Aug 07, 2027
					ODE-400	May 27, 2029
<u>RISEDRONATE SODIUM - ATELVIA</u>						
N 022560	001	7645459	Jan 09, 2028	DP U-662		
		7645460	Jan 09, 2028	DP U-662		
		8246989	Jan 16, 2026	DP		
<u>RISPERIDONE - PERSERIS KIT</u>						
N 210655	001	10010612	Feb 13, 2028	DP		
		10058554	Sep 26, 2026	U-2363		
		10376590	Feb 13, 2028	U-2608		
		10406160	Jun 26, 2026	DP U-2608		
		11013809	Feb 13, 2028	DP U-3135		
		11110093	Nov 05, 2026	DP U-3135		
		11712475	Feb 13, 2028	U-3135		
		9180197	Feb 13, 2028	DP		
		9186413	Feb 13, 2028	U-543		
		9597402	Sep 26, 2026	DP		
<u>RISPERIDONE - PERSERIS KIT</u>						
N 210655	002	10010612	Feb 13, 2028	DP		
		10058554	Sep 26, 2026	U-2363		
		10376590	Feb 13, 2028	U-2608		
		10406160	Jun 26, 2026	DP U-2608		
		11013809	Feb 13, 2028	DP U-3135		
		11110093	Nov 05, 2026	DP U-3135		
		11712475	Feb 13, 2028	U-3135		
		9180197	Feb 13, 2028	DP		
		9186413	Feb 13, 2028	U-543		
		9597402	Sep 26, 2026	DP		
<u>RISPERIDONE - RYKINDO</u>						
N 212849	001	10098882	Apr 10, 2032	DP U-3513		
		10406161	Apr 10, 2032	DP U-3513		
		11110094	Apr 10, 2032	DP		
		9446135	Apr 10, 2032	DP U-3513		

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<u>RISPERIDONE - RYKINDO</u>						
N 212849	001 9532991	Apr 10, 2032	DP U-3513			
<u>RISPERIDONE - RYKINDO</u>						
N 212849	002 10098882	Apr 10, 2032	DP U-3513			
	10406161	Apr 10, 2032	DP U-3513			
	11110094	Apr 10, 2032	DP			
	9446135	Apr 10, 2032	DP U-3513			
	9532991	Apr 10, 2032	DP U-3513			
<u>RISPERIDONE - RYKINDO</u>						
N 212849	003 10098882	Apr 10, 2032	DP U-3513			
	10406161	Apr 10, 2032	DP U-3513			
	11110094	Apr 10, 2032	DP			
	9446135	Apr 10, 2032	DP U-3513			
	9532991	Apr 10, 2032	DP U-3513			
<u>RISPERIDONE - RYKINDO</u>						
N 212849	004 10098882	Apr 10, 2032	DP U-3513			
	10406161	Apr 10, 2032	DP U-3513			
	11110094	Apr 10, 2032	DP			
	9446135	Apr 10, 2032	DP U-3513			
	9532991	Apr 10, 2032	DP U-3513			
<u>RISPERIDONE - UZEDY</u>						
N 213586	001 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	002 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	003 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			

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<u>RISPERIDONE - UZEDY</u>						
N 213586	003 9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	004 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	005 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	006 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RISPERIDONE - UZEDY</u>						
N 213586	007 10736965	Jan 12, 2025	DP		NP	Apr 28, 2026
	8221778	Nov 12, 2027	DP U-543			
	8741327	Nov 12, 2027	DP U-543			
	8802127	Jan 12, 2025	DP			
	9023897	Apr 05, 2033	DP			
	9439905	Jan 12, 2025	DP U-543			
	9717799	Jan 12, 2025	DP			
	9895447	Jan 12, 2025	DP			
	9925268	Jan 12, 2025	DP			
<u>RITLECITINIB TOSYLATE - LITFULO</u>						
N 215830	001 9617258	Dec 03, 2034	DS DP		NCE	Jun 23, 2028
<u>RIVAROXABAN - XARELTO</u>						
N 022406	001 7157456	Aug 28, 2024	DS DP U-1301		I-867	Aug 23, 2024
	7157456	Aug 28, 2024	DS DP U-1302		PED	Feb 23, 2025

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<u>RIVAROXABAN - XARELTO</u>						
N 022406 001	7157456*PED	Feb 28, 2025				
	9415053	Nov 13, 2024	DP U-1167			
	9415053	Nov 13, 2024	DP U-2142			
	9415053	Nov 13, 2024	DP U-2640			
	9415053	Nov 13, 2024	DP U-3284			
	9415053*PED	May 13, 2025				
	9539218	Feb 17, 2034	U-1957			
	9539218	Feb 17, 2034	U-2143			
	9539218	Feb 17, 2034	U-2641			
	9539218	Feb 17, 2034	U-3288			
	9539218*PED	Aug 17, 2034				
<u>RIVAROXABAN - XARELTO</u>						
N 022406 002	7157456	Aug 28, 2024	DS DP U-1301		I-867	Aug 23, 2024
	7157456	Aug 28, 2024	DS DP U-1302		PED	Feb 23, 2025
	7157456*PED	Feb 28, 2025				
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			
	9415053	Nov 13, 2024	DP U-1302			
	9415053	Nov 13, 2024	DP U-3286			
	9415053*PED	May 13, 2025				
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-3289			
	9539218*PED	Aug 17, 2034				
<u>RIVAROXABAN - XARELTO</u>						
N 022406 003	7157456	Aug 28, 2024	DS DP U-1301		I-867	Aug 23, 2024
	7157456	Aug 28, 2024	DS DP U-1302		PED	Feb 23, 2025
	7157456*PED	Feb 28, 2025				
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			
	9415053	Nov 13, 2024	DP U-1302			
	9415053	Nov 13, 2024	DP U-3287			
	9415053*PED	May 13, 2025				
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-3285			
	9539218*PED	Aug 17, 2034				
<u>RIVAROXABAN - XARELTO</u>						
N 022406 004	10828310	Jan 31, 2039	U-3207		I-867	Aug 23, 2024
	10828310	Jan 31, 2039	U-3208		PED	Feb 23, 2025
	10828310*PED	Jul 31, 2039				
	7157456	Aug 28, 2024	DS DP			
	7157456*PED	Feb 28, 2025				
	9415053	Nov 13, 2024	DP U-2435			
	9415053	Nov 13, 2024	DP U-3205			
	9415053	Nov 13, 2024	DP U-3206			
	9415053*PED	May 13, 2025				

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<u>RIVAROXABAN - XARELTO</u>						
N 215859	001 7157456	Aug 28, 2024	DS DP		NP	Dec 20, 2024
					PED	Jun 20, 2025
<u>RIZATRIPTAN BENZOATE - RIZAFILM</u>						
N 205394	001 9301948	Jul 30, 2034	DS DP			
<u>ROFLUMILAST - DALIRESP</u>						
N 022522	001 8536206	Mar 08, 2024		U-1115		
	8604064	Mar 08, 2024		U-1115		
	8618142	Mar 08, 2024	DP			
<u>ROFLUMILAST - DALIRESP</u>						
N 022522	002 8536206	Mar 08, 2024		U-1115		
	8604064	Mar 08, 2024		U-1115		
	8618142	Mar 08, 2024	DP			
<u>ROFLUMILAST - ZORYVE</u>						
N 215985	001 10940142	Jun 07, 2037	DP		NP	Jul 29, 2025
	11129818	Aug 25, 2037		U-3408	NPP	Oct 05, 2026
	11129818	Aug 25, 2037		U-3712		
	11793796	Jun 07, 2037	DP			
	11819496	Jun 07, 2037		U-3748		
	9884050	Jun 07, 2037	DP			
	9907788	Jun 07, 2037		U-3408		
	9907788	Jun 07, 2037		U-3712		
<u>ROFLUMILAST - ZORYVE</u>						
N 217242	001 10940142	Jun 07, 2037	DP			
	11129818	Aug 25, 2037		U-3773		
	11707454	Dec 03, 2041		U-3773		
	11793796	Jun 07, 2037	DP			
	11819496	Jun 07, 2037		U-3773		
	9884050	Jun 07, 2037	DP			
	9907788	Jun 07, 2037		U-3773		
<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 206500	001 7049320	Aug 19, 2028	DS DP	U-1741		
	7563801	Apr 04, 2027	DP			
	7981905	Apr 04, 2027		U-1741		
	8178550	Apr 04, 2027	DS DP			
	8361500	Oct 09, 2029	DP			
	8404702	Apr 04, 2027		U-1741		
	8470842	Jan 18, 2029		U-1741		
<u>ROLAPITANT HYDROCHLORIDE - VARUBI</u>						
N 208399	001 7049320	Aug 19, 2028	DS DP	U-1741		
	7981905	Apr 04, 2027		U-1741		
	8178550	Apr 04, 2027	DS DP			
	8404702	Apr 04, 2027		U-1741		
	8470842	Jan 18, 2029		U-1741		
	9101615	Jul 14, 2032		U-1741		
<u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u>						
N 020533	006 7828787	Oct 18, 2025	DP			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u>						
N 020533	006	7857802	Nov 28, 2026	DP		
		8162915	May 23, 2024	DP		
<u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u>						
N 020533	007	7828787	Oct 18, 2025	DP		
		7857802	Nov 28, 2026	DP		
		8162915	May 23, 2024	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	001	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	002	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	003	10413543	Feb 12, 2036	DP		
<u>ROSUVASTATIN CALCIUM - EZALLOR SPRINKLE</u>						
N 208647	004	10413543	Feb 12, 2036	DP		
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	001	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	DP		
		8246979	Sep 01, 2027	DP U-1272		
		8246979	Sep 01, 2027	DP U-1273		
		8246980	Nov 27, 2025	DP		
		9925150	Mar 01, 2032	DP		
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	002	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	DP		
		8246979	Sep 01, 2027	DP U-1272		
		8246979	Sep 01, 2027	DP U-1273		
		8246980	Nov 27, 2025	DP		
		9925150	Mar 01, 2032	DP		
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	003	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	DP		
		8246979	Sep 01, 2027	DP U-1272		
		8246979	Sep 01, 2027	DP U-1273		
		8246980	Nov 27, 2025	DP		
		9925150	Mar 01, 2032	DP		
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	004	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	DP		
		8246979	Sep 01, 2027	DP U-1272		
		8246979	Sep 01, 2027	DP U-1273		
		8246980	Nov 27, 2025	DP		
		9925150	Mar 01, 2032	DP		
<u>ROTIGOTINE - NEUPRO</u>						
N 021829	005	10130589	Dec 22, 2030	DP		
		10350174	Dec 22, 2030	DP		
		8246979	Sep 01, 2027	DP U-1272		

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<u>ROTIGOTINE - NEUPRO</u>						
N 021829 005	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>ROTIGOTINE - NEUPRO</u>						
N 021829 006	10130589	Dec 22, 2030	DP			
	10350174	Dec 22, 2030	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	9925150	Mar 01, 2032	DP			
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 001	10130636	Aug 17, 2035	U-2012		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2101			
	10130636	Aug 17, 2035	U-2273			
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031	DP			
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2273			
	7351701	Jul 23, 2024	U-2830			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2830			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			
	9861638	Feb 10, 2031	U-2273			
	9987285	Aug 17, 2035	DP			
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 002	10130636	Aug 17, 2035	U-2012		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2101			
	10130636	Aug 17, 2035	U-2273			
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031	DP			
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2273			
	7351701	Jul 23, 2024	U-2830			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2830			

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<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 002	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			
	9861638	Feb 10, 2031	U-2273			
	9987285	Aug 17, 2035	DP			
<u>RUCAPARIB CAMSYLATE - RUBRACA</u>						
N 209115 003	10130636	Aug 17, 2035	U-2012		ODE-168	Apr 06, 2025
	10130636	Aug 17, 2035	U-2101			
	10130636	Aug 17, 2035	U-2273			
	10130636	Aug 17, 2035	U-2830			
	10278974	Feb 10, 2031	DP			
	7351701	Jul 23, 2024	U-2012			
	7351701	Jul 23, 2024	U-2273			
	7351701	Jul 23, 2024	U-2830			
	7531530	Jul 23, 2024	U-2012			
	7531530	Jul 23, 2024	U-2273			
	7531530	Jul 23, 2024	U-2830			
	8071579	Aug 12, 2027	U-2012			
	8071579	Aug 12, 2027	U-2273			
	8071579	Aug 12, 2027	U-2830			
	8143241	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2273			
	8143241	Aug 12, 2027	U-2830			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	8859562	Aug 04, 2031	U-2273			
	8859562	Aug 04, 2031	U-2830			
	9045487	Feb 10, 2031	DS DP			
	9861638	Feb 10, 2031	U-2012			
	9861638	Feb 10, 2031	U-2273			
	9987285	Aug 17, 2035	DP			
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 001	10016429	Jun 12, 2028	U-3226		I-872	Sep 22, 2024
	10016429	Jun 12, 2028	U-3230		M-285	Dec 19, 2025
	10016429*PED	Dec 12, 2028			ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP U-3227		ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3228		PED	Mar 22, 2025
	7598257*PED	Jun 24, 2028			PED	Jun 19, 2026
	8415362	Dec 24, 2027	DS DP		PED	Nov 24, 2026
	8415362*PED	Jun 24, 2028			PED	Mar 22, 2029

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<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 001	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028		U-1573		
	8822481	Jun 12, 2028		U-3226		
	8822481	Jun 12, 2028		U-3227		
	8822481	Jun 12, 2028		U-3228		
	8822481	Jun 12, 2028		U-3230		
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	8829013	Jun 12, 2028		U-3227		
	8829013	Jun 12, 2028		U-3228		
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026		U-3226		
	9079912	Dec 12, 2026		U-3227		
	9079912	Dec 12, 2026		U-3228		
	9079912	Dec 12, 2026		U-3230		
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026		U-3226		
	9814722	Dec 12, 2026		U-3230		
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 002	10016429	Jun 12, 2028		U-3226	I-872	Sep 22, 2024
	10016429	Jun 12, 2028		U-3230	M-285	Dec 19, 2025
	10016429*PED	Dec 12, 2028			ODE-238	May 24, 2026
	7598257	Dec 24, 2027	DS DP	U-3227	ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP	U-3228	PED	Mar 22, 2025
	7598257*PED	Jun 24, 2028			PED	Jun 19, 2026
	8415362	Dec 24, 2027	DS DP		PED	Nov 24, 2026
	8415362*PED	Jun 24, 2028			PED	Mar 22, 2029
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028		U-1573		
	8822481	Jun 12, 2028		U-3226		
	8822481	Jun 12, 2028		U-3227		
	8822481	Jun 12, 2028		U-3228		
	8822481	Jun 12, 2028		U-3230		
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	8829013	Jun 12, 2028		U-3227		
	8829013	Jun 12, 2028		U-3228		
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026		U-3226		
	9079912	Dec 12, 2026		U-3227		
	9079912	Dec 12, 2026		U-3228		
	9079912	Dec 12, 2026		U-3230		
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026		U-3226		

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<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 002	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 003	10016429	Jun 12, 2028	U-3226		M-285	Dec 19, 2025
	10016429	Jun 12, 2028	U-3230		ODE-238	May 24, 2026
	10016429*PED	Dec 12, 2028			ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3227		PED	Jun 19, 2026
	7598257	Dec 24, 2027	DS DP U-3228		PED	Nov 24, 2026
	7598257*PED	Jun 24, 2028			PED	Mar 22, 2029
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-1573			
	8822481	Jun 12, 2028	U-3226			
	8822481	Jun 12, 2028	U-3227			
	8822481	Jun 12, 2028	U-3228			
	8822481	Jun 12, 2028	U-3230			
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	8829013	Jun 12, 2028	U-3227			
	8829013	Jun 12, 2028	U-3228			
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3226			
	9079912	Dec 12, 2026	U-3227			
	9079912	Dec 12, 2026	U-3228			
	9079912	Dec 12, 2026	U-3230			
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026	U-3226			
	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 004	10016429	Jun 12, 2028	U-3226		M-285	Dec 19, 2025
	10016429	Jun 12, 2028	U-3230		ODE-238	May 24, 2026
	10016429*PED	Dec 12, 2028			ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3227		PED	Jun 19, 2026
	7598257	Dec 24, 2027	DS DP U-3228		PED	Nov 24, 2026
	7598257*PED	Jun 24, 2028			PED	Mar 22, 2029
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-1573			
	8822481	Jun 12, 2028	U-3226			
	8822481	Jun 12, 2028	U-3227			
	8822481	Jun 12, 2028	U-3228			
	8822481	Jun 12, 2028	U-3230			
	8822481*PED	Dec 12, 2028				

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<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 004	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	8829013	Jun 12, 2028	U-3227			
	8829013	Jun 12, 2028	U-3228			
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3226			
	9079912	Dec 12, 2026	U-3227			
	9079912	Dec 12, 2026	U-3228			
	9079912	Dec 12, 2026	U-3230			
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026	U-3226			
	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>						
N 202192 005	10016429	Jun 12, 2028	U-3226		M-285	Dec 19, 2025
	10016429	Jun 12, 2028	U-3230		ODE-238	May 24, 2026
	10016429*PED	Dec 12, 2028			ODE-373	Sep 22, 2028
	7598257	Dec 24, 2027	DS DP U-3227		PED	Jun 19, 2026
	7598257	Dec 24, 2027	DS DP U-3228		PED	Nov 24, 2026
	7598257*PED	Jun 24, 2028			PED	Mar 22, 2029
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-1573			
	8822481	Jun 12, 2028	U-3226			
	8822481	Jun 12, 2028	U-3227			
	8822481	Jun 12, 2028	U-3228			
	8822481	Jun 12, 2028	U-3230			
	8822481*PED	Dec 12, 2028				
	8829013	Jun 12, 2028	U-1201			
	8829013	Jun 12, 2028	U-1622			
	8829013	Jun 12, 2028	U-3227			
	8829013	Jun 12, 2028	U-3228			
	8829013*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3226			
	9079912	Dec 12, 2026	U-3227			
	9079912	Dec 12, 2026	U-3228			
	9079912	Dec 12, 2026	U-3230			
	9079912*PED	Jun 12, 2027				
	9814722	Dec 12, 2026	U-3226			
	9814722	Dec 12, 2026	U-3230			
	9814722*PED	Jun 12, 2027				
<u>RUXOLITINIB PHOSPHATE - OPZELURA</u>						
N 215309 001	10610530	Jun 12, 2028	U-3229		I-896	Jul 18, 2025
	10610530	Jun 12, 2028	U-3404		NP	Sep 21, 2024
	10610530*PED	Dec 12, 2028			PED	Mar 21, 2025
	10639310	Dec 12, 2026	U-3229		PED	Jan 18, 2026
	10639310*PED	Jun 12, 2027				

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<u>RUXOLITINIB PHOSPHATE - OPZELURA</u>						
N 215309 001	10758543	May 20, 2031	DP			
	10758543*PED	Nov 20, 2031				
	10869870	May 20, 2031	U-3229			
	10869870	May 20, 2031	U-3404			
	10869870*PED	Nov 20, 2031				
	11219624	May 20, 2031	DP U-3229			
	11219624*PED	Nov 20, 2031				
	11510923	Sep 04, 2040	U-3505			
	11571425	May 20, 2031	DP			
	11590136	May 20, 2031	U-3229			
	11590136	May 20, 2031	U-3404			
	11590137	Sep 04, 2040	U-3505			
	11590138	Jun 10, 2040	U-3551			
	11602536	May 05, 2041	U-3550			
	7598257	Dec 24, 2027	DS DP			
	7598257*PED	Jun 24, 2028				
	8415362	Dec 24, 2027	DS DP			
	8415362*PED	Jun 24, 2028				
	8722693	Jun 12, 2028	DS DP			
	8722693*PED	Dec 12, 2028				
	8822481	Jun 12, 2028	U-3229			
	8822481	Jun 12, 2028	U-3404			
	8822481*PED	Dec 12, 2028				
	9079912	Dec 12, 2026	U-3229			
	9079912	Dec 12, 2026	U-3404			
	9079912*PED	Jun 12, 2027				
	9974790	Dec 12, 2026	U-3229			
	9974790	Dec 12, 2026	U-3404			
	9974790*PED	Jun 12, 2027				
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620 001	11058667	May 09, 2036	U-3170		M-82	Feb 16, 2024
	11135192	Aug 22, 2033	U-3084			
	7468390*PED	May 27, 2024				
	8101659	Jan 15, 2025	DP			
	8101659*PED	Jul 15, 2025				
	8877938	May 27, 2027	DS DP			
	8877938*PED	Nov 27, 2027				
	9388134	Nov 08, 2026	U-1723			
	9388134*PED	May 08, 2027				
	9517226	Aug 22, 2033	U-3084			
	9937143	Aug 22, 2033	U-3084			
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620 002	11058667	May 09, 2036	U-3170		M-82	Feb 16, 2024
	11135192	Aug 22, 2033	U-3084			
	7468390*PED	May 27, 2024				
	8101659	Jan 15, 2025	DP			
	8101659*PED	Jul 15, 2025				
	8877938	May 27, 2027	DS DP			
	8877938*PED	Nov 27, 2027				

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<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	002 9388134	Nov 08, 2026	U-1723			
	9388134*PED	May 08, 2027				
	9517226	Aug 22, 2033	U-3084			
	9937143	Aug 22, 2033	U-3084			
<u>SACUBITRIL; VALSARTAN - ENTRESTO</u>						
N 207620	003 11058667	May 09, 2036	U-3170		M-82	Feb 16, 2024
	11135192	Aug 22, 2033	U-3084			
	7468390*PED	May 27, 2024				
	8101659	Jan 15, 2025	DP			
	8101659*PED	Jul 15, 2025				
	8877938	May 27, 2027	DS DP			
	8877938*PED	Nov 27, 2027				
	9388134	Nov 08, 2026	U-1723			
	9388134*PED	May 08, 2027				
	9517226	Aug 22, 2033	U-3084			
	9937143	Aug 22, 2033	U-3084			
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145	001 8076515	Dec 10, 2028	DS DP U-1993			
	8278485	Jun 08, 2027	DS U-1993			
	8283380	Mar 21, 2031	U-1993			
<u>SAFINAMIDE MESYLATE - XADAGO</u>						
N 207145	002 8076515	Dec 10, 2028	DS DP U-1993			
	8278485	Jun 08, 2027	DS U-1993			
	8283380	Mar 21, 2031	U-1993			
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 022181	001 7566462	Nov 16, 2025	DP			
	7566462*PED	May 16, 2026				
	7566714	Nov 17, 2024	U-989			
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024	U-1010			
	7612073*PED	May 17, 2025				
	7727987	Nov 17, 2024	DP			
	7727987*PED	May 17, 2025				
	8003126	Nov 16, 2025				
	8003126*PED	May 16, 2026				
	8067416	Nov 17, 2024	U-989			
	8067416*PED	May 17, 2025				
	8318745	Nov 17, 2024	DP			
	8318745*PED	May 17, 2025				
	9433624	Nov 17, 2024	U-1589			
	9433624*PED	May 17, 2025				
	RE43797	Nov 17, 2024	U-1156			
	RE43797*PED	May 17, 2025				
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065	001 7566714	Nov 17, 2024	U-1589			
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024	U-1010			
	7612073*PED	May 17, 2025				

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<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065	001	8067416	Nov 17, 2024		U-1589	
		8067416*PED	May 17, 2025			
		9216178	Nov 01, 2032	DP		
		9216178*PED	May 01, 2033			
		9433624	Nov 17, 2024		U-1589	
		9433624*PED	May 17, 2025			
		RE43797	Nov 17, 2024		U-1590	
		RE43797*PED	May 17, 2025			
<u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u>						
N 205065	002	7566714	Nov 17, 2024		U-1589	
		7566714*PED	May 17, 2025			
		7612073	Nov 17, 2024		U-1010	
		7612073*PED	May 17, 2025			
		8067416	Nov 17, 2024		U-1589	
		8067416*PED	May 17, 2025			
		9216178	Nov 01, 2032	DP		
		9216178*PED	May 01, 2033			
		9433624	Nov 17, 2024		U-1589	
		9433624*PED	May 17, 2025			
		RE43797	Nov 17, 2024		U-1590	
		RE43797*PED	May 17, 2025			
<u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u>						
N 209521	001	8318706	May 01, 2031	DS DP	U-2405	
		8513223	Dec 07, 2029		U-2406	
		9255068	Feb 09, 2033	DS DP	U-2407	
		9255068	Feb 09, 2033	DS DP	U-2408	
		9481639	Aug 10, 2028		U-2409	
<u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u>						
N 209521	002	8318706	May 01, 2031	DS DP	U-2405	
		8513223	Dec 07, 2029		U-2406	
		9255068	Feb 09, 2033	DS DP	U-2407	
		9255068	Feb 09, 2033	DS DP	U-2408	
		9481639	Aug 10, 2028		U-2409	
<u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u>						
N 209521	003	8318706	May 01, 2031	DS DP	U-2405	
		8513223	Dec 07, 2029		U-2406	
		9255068	Feb 09, 2033	DS DP	U-2407	
		9255068	Feb 09, 2033	DS DP	U-2408	
		9481639	Aug 10, 2028		U-2409	
<u>SAXAGLIPTIN HYDROCHLORIDE - SAXAGLIPTIN</u>						
A 205941	001				PC	Jan 27, 2024
<u>SAXAGLIPTIN HYDROCHLORIDE - SAXAGLIPTIN</u>						
A 205941	002				PC	Jan 27, 2024
<u>SAXAGLIPTIN HYDROCHLORIDE - SAXAGLIPTIN</u>						
A 205972	001				PC	Jan 27, 2024

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<u>SAXAGLIPTIN HYDROCHLORIDE - SAXAGLIPTIN</u>						
A 205972	002				PC	Jan 27, 2024
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
N 022350	001 7951400	Nov 30, 2028	DP			
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
N 022350	002 7951400	Nov 30, 2028	DP			
<u>SECNIDAZOLE - SOLOSEC</u>						
N 209363	001 10335390	Sep 04, 2035	DP U-2583		I-866	Jun 30, 2024
	10335390	Sep 04, 2035	DP U-3270		NCE	Sep 15, 2022
	10682338	Sep 04, 2035	DP U-2583		NPP	Jan 26, 2025
	10682338	Sep 04, 2035	DP U-3270		GAIN	Sep 15, 2027
	10849884	Sep 04, 2035	DP U-2583			
	10849884	Sep 04, 2035	DP U-3169			
	10849884	Sep 04, 2035	DP U-3270			
	10849884	Sep 04, 2035	DP U-3302			
	10857133	Sep 04, 2035	DP U-2583			
	10857133	Sep 04, 2035	DP U-3270			
	11000507	Sep 04, 2035	DP U-2583			
	11000507	Sep 04, 2035	DP U-3169			
	11000507	Sep 04, 2035	DP U-3270			
	11000507	Sep 04, 2035	DP U-3302			
	11000508	Sep 04, 2035	DP U-2583			
	11000508	Sep 04, 2035	DP U-3169			
	11000508	Sep 04, 2035	DP U-3270			
	11000508	Sep 04, 2035	DP U-3302			
	11020377	Sep 04, 2035	DP U-2583			
	11020377	Sep 04, 2035	DP U-3169			
	11020377	Sep 04, 2035	DP U-3270			
	11020377	Sep 04, 2035	DP U-3302			
	11324721	Sep 04, 2035	DP U-2583			
	11324721	Sep 04, 2035	DP U-3169			
	11324721	Sep 04, 2035	DP U-3270			
	11324721	Sep 04, 2035	DP U-3302			
	11602522	Sep 04, 2035	DP U-3169			
	11602522	Sep 04, 2035	DP U-3302			
	11684607	Sep 16, 2035	DP U-2583			
	11684607	Sep 16, 2035	DP U-3169			
	11684607	Sep 16, 2035	DP U-3270			
	11684607	Sep 16, 2035	DP U-3302			
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379	001				NCE	Apr 30, 2024
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379	002				NCE	Apr 30, 2024
<u>SELENIOS ACID - SELENIOS ACID</u>						
N 209379	003				NCE	Apr 30, 2024
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	001 10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			

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<u>SELEXIPAG - UPTRAVI</u>						
N 207947	001	7205302	Oct 31, 2026	DS DP	U-1797	
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029		U-1798	
		9284280	Jun 25, 2030		U-1831	
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	002	10821108	Dec 01, 2036	DP	U-2992	
		10828298	Dec 01, 2036	DP	U-2991	
		7205302	Oct 31, 2026	DS DP	U-1797	
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029		U-1798	
		9284280	Jun 25, 2030		U-1831	
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	003	10821108	Dec 01, 2036	DP	U-2992	
		10828298	Dec 01, 2036	DP	U-2991	
		7205302	Oct 31, 2026	DS DP	U-1797	
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029		U-1798	
		9284280	Jun 25, 2030		U-1831	
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	004	10821108	Dec 01, 2036	DP	U-2992	
		10828298	Dec 01, 2036	DP	U-2991	
		7205302	Oct 31, 2026	DS DP	U-1797	
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029		U-1798	
		9284280	Jun 25, 2030		U-1831	
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	005	10821108	Dec 01, 2036	DP	U-2992	
		10828298	Dec 01, 2036	DP	U-2991	
		7205302	Oct 31, 2026	DS DP	U-1797	
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029		U-1798	
		9284280	Jun 25, 2030		U-1831	
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	006	10821108	Dec 01, 2036	DP	U-2992	
		10828298	Dec 01, 2036	DP	U-2991	
		7205302	Oct 31, 2026	DS DP	U-1797	
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029		U-1798	
		9284280	Jun 25, 2030		U-1831	
<u>SELEXIPAG - UPTRAVI</u>						
N 207947	007	10821108	Dec 01, 2036	DP	U-2992	
		10828298	Dec 01, 2036	DP	U-2991	
		7205302	Oct 31, 2026	DS DP	U-1797	
		8791122	Aug 01, 2030	DS DP		
		9173881	Aug 12, 2029		U-1798	
		9284280	Jun 25, 2030		U-1831	

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<u>SELEXIPAG - UPTRAVI</u>						
N 207947 008	10821108	Dec 01, 2036	DP U-2992			
	10828298	Dec 01, 2036	DP U-2991			
	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELEXIPAG - UPTRAVI</u>						
N 214275 001	7205302	Oct 31, 2026	DS DP U-1797			
	8791122	Aug 01, 2030	DS DP			
	9173881	Aug 12, 2029	U-1798			
	9284280	Jun 25, 2030	U-1831			
<u>SELINEXOR - XPOVIO</u>						
N 212306 001	10519139	Aug 14, 2035	DS DP U-2584		NCE	Jul 03, 2024
	10519139	Aug 14, 2035	DS DP U-2855		ODE-257	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-3018		ODE-310	Jun 22, 2027
	10544108	Jul 26, 2032	U-2584		ODE-346	Dec 18, 2027
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	11746102	Aug 14, 2035	U-2584			
	11746102	Aug 14, 2035	U-2855			
	11746102	Aug 14, 2035	U-3018			
	11753401	Aug 14, 2035	DP U-2584			
	11753401	Aug 14, 2035	DP U-2855			
	11753401	Aug 14, 2035	DP U-3018			
	11787771	Jul 26, 2032	U-2584			
	11787771	Jul 26, 2032	U-2855			
	11787771	Jul 26, 2032	U-3018			
	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 002	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	11746102	Aug 14, 2035	U-2584			
	11746102	Aug 14, 2035	U-2855			
	11746102	Aug 14, 2035	U-3018			
	11753401	Aug 14, 2035	DP U-2584			
	11753401	Aug 14, 2035	DP U-2855			
	11753401	Aug 14, 2035	DP U-3018			

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<u>SELINEXOR - XPOVIO</u>						
N 212306 002	11787771	Jul 26, 2032	U-2584			
	11787771	Jul 26, 2032	U-2855			
	11787771	Jul 26, 2032	U-3018			
	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 003	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	11746102	Aug 14, 2035	U-2584			
	11746102	Aug 14, 2035	U-2855			
	11746102	Aug 14, 2035	U-3018			
	11753401	Aug 14, 2035	DP U-2584			
	11753401	Aug 14, 2035	DP U-2855			
	11753401	Aug 14, 2035	DP U-3018			
	11787771	Jul 26, 2032	U-2584			
	11787771	Jul 26, 2032	U-2855			
	11787771	Jul 26, 2032	U-3018			
	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032	U-2584			
	9079865	Jul 26, 2032	U-2855			
	9079865	Jul 26, 2032	U-3018			
	9714226	Jul 26, 2032	DS DP			
<u>SELINEXOR - XPOVIO</u>						
N 212306 004	10519139	Aug 14, 2035	DS DP U-2584		ODE*	Jul 03, 2026
	10519139	Aug 14, 2035	DS DP U-2855		ODE*	Jun 22, 2027
	10519139	Aug 14, 2035	DS DP U-3018		ODE*	Dec 18, 2027
	10544108	Jul 26, 2032	U-2584			
	10544108	Jul 26, 2032	U-3018			
	11034660	Jul 26, 2032	U-2584			
	11034660	Jul 26, 2032	U-3018			
	11746102	Aug 14, 2035	U-2584			
	11746102	Aug 14, 2035	U-2855			
	11746102	Aug 14, 2035	U-3018			
	11753401	Aug 14, 2035	DP U-2584			
	11753401	Aug 14, 2035	DP U-2855			
	11753401	Aug 14, 2035	DP U-3018			
	11787771	Jul 26, 2032	U-2584			
	11787771	Jul 26, 2032	U-2855			
	11787771	Jul 26, 2032	U-3018			

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<u>SELINEXOR - XPOVIO</u>						
N 212306 004	11807629	Aug 14, 2035	DS DP			
	8999996	Jul 03, 2033	DS DP			
	9079865	Jul 26, 2032		U-2584		
	9079865	Jul 26, 2032		U-2855		
	9079865	Jul 26, 2032		U-3018		
	9714226	Jul 26, 2032	DS DP			
<u>SELPERCATINIB - RETEVMO</u>						
N 213246 001	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037		U-2826	ODE-301	May 08, 2027
	10137124	Oct 10, 2037		U-2827	ODE-302	May 08, 2027
	10137124	Oct 10, 2037		U-2828	ODE-303	May 08, 2027
	10137124	Oct 10, 2037		U-3450	ODE-409	Sep 21, 2029
	10137124	Oct 10, 2037		U-3451	ODE-412	Sep 21, 2029
	10137124	Oct 10, 2037		U-3452		
	10137124	Oct 10, 2037		U-3453		
	10172851	Oct 10, 2037		U-2826		
	10172851	Oct 10, 2037		U-2827		
	10172851	Oct 10, 2037		U-2828		
	10172851	Oct 10, 2037		U-3450		
	10172851	Oct 10, 2037		U-3451		
	10172851	Oct 10, 2037		U-3452		
	10172851	Oct 10, 2037		U-3453		
	10584124	Oct 10, 2038	DS	U-2826		
	10584124	Oct 10, 2038	DS	U-2827		
	10584124	Oct 10, 2038	DS	U-2828		
	10584124	Oct 10, 2038	DS	U-3450		
	10584124	Oct 10, 2038	DS	U-3451		
	10584124	Oct 10, 2038	DS	U-3452		
	10584124	Oct 10, 2038	DS	U-3453		
	10786489	Oct 10, 2038	DP	U-2971		
	10786489	Oct 10, 2038	DP	U-2972		
	10786489	Oct 10, 2038	DP	U-2973		
	10786489	Oct 10, 2038	DP	U-2974		
	10786489	Oct 10, 2038	DP	U-2975		
	10786489	Oct 10, 2038	DP	U-2976		
	10786489	Oct 10, 2038	DP	U-2977		
	10786489	Oct 10, 2038	DP	U-3450		
	10786489	Oct 10, 2038	DP	U-3451		
	10786489	Oct 10, 2038	DP	U-3452		
	10786489	Oct 10, 2038	DP	U-3453		
<u>SELPERCATINIB - RETEVMO</u>						
N 213246 002	10112942	Oct 10, 2037	DS DP		NCE	May 08, 2025
	10137124	Oct 10, 2037		U-2826	ODE-301	May 08, 2027
	10137124	Oct 10, 2037		U-2827	ODE-302	May 08, 2027
	10137124	Oct 10, 2037		U-2828	ODE-303	May 08, 2027
	10137124	Oct 10, 2037		U-3450	ODE-409	Sep 21, 2029
	10137124	Oct 10, 2037		U-3451	ODE-412	Sep 21, 2029
	10137124	Oct 10, 2037		U-3452		
	10137124	Oct 10, 2037		U-3453		

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<u>SEPERCATINIB - RETEVMO</u>						
N 213246	002 10172851	Oct 10, 2037		U-2826		
	10172851	Oct 10, 2037		U-2827		
	10172851	Oct 10, 2037		U-2828		
	10584124	Oct 10, 2038	DS	U-2826		
	10584124	Oct 10, 2038	DS	U-2827		
	10584124	Oct 10, 2038	DS	U-2828		
	10584124	Oct 10, 2038	DS	U-3450		
	10584124	Oct 10, 2038	DS	U-3451		
	10584124	Oct 10, 2038	DS	U-3452		
	10584124	Oct 10, 2038	DS	U-3453		
	10786489	Oct 10, 2038	DP	U-2971		
	10786489	Oct 10, 2038	DP	U-2972		
	10786489	Oct 10, 2038	DP	U-2973		
	10786489	Oct 10, 2038	DP	U-2974		
	10786489	Oct 10, 2038	DP	U-2975		
	10786489	Oct 10, 2038	DP	U-2976		
	10786489	Oct 10, 2038	DP	U-2977		
	10786489	Oct 10, 2038	DP	U-3450		
	10786489	Oct 10, 2038	DP	U-3451		
	10786489	Oct 10, 2038	DP	U-3452		
	10786489	Oct 10, 2038	DP	U-3453		
<u>SELUMETINIB SULFATE - KOSELUGO</u>						
N 213756	001 11813246	Mar 26, 2029		DP	NCE	Apr 10, 2025
	7425637	Mar 13, 2024	DS		ODE-288	Apr 10, 2027
	9156795	Dec 12, 2026	DS	DP		
	9562017	Dec 12, 2026	DS	U-2800		
<u>SELUMETINIB SULFATE - KOSELUGO</u>						
N 213756	002 11813246	Mar 26, 2029		DP	NCE	Apr 10, 2025
	7425637	Mar 13, 2024	DS		ODE-288	Apr 10, 2027
	9156795	Dec 12, 2026	DS	DP		
	9562017	Dec 12, 2026	DS	U-2800		
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637	001 10220155	Jul 17, 2026		DP		
	10335462	Jun 21, 2033		U-2580		
	10357616	Jan 20, 2026		DP		
	10376652	Jan 20, 2026		DP		
	11097063	Jul 17, 2026		DP		
	11311679	Jan 20, 2026		DP		
	11446443	Oct 20, 2025		DP		
	7762994	May 23, 2024		DP		
	8114833	Aug 13, 2025		DP		
	8129343	Dec 05, 2031	DS	DP U-2202		
	8536122	Mar 20, 2026	DS	DP U-2202		
	8684969	Oct 20, 2025		DP		
	8920383	Jul 17, 2026		DP		
	9108002	Jan 20, 2026		DP		
	9132239	Feb 01, 2032		DP		
	9457154	Sep 27, 2027		DP		

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<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637	001	9616180	Jan 20, 2026	DP		
		9687611	Feb 27, 2027	DP		
		9775953	Jul 17, 2026	DP		
		9861757	Jan 20, 2026	DP		
		RE46363	Aug 03, 2026	DP		
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637	002	10220155	Jul 17, 2026	DP		
		10335462	Jun 21, 2033	U-2580		
		10357616	Jan 20, 2026	DP		
		10376652	Jan 20, 2026	DP		
		11097063	Jul 17, 2026	DP		
		11311679	Jan 20, 2026	DP		
		11446443	Oct 20, 2025	DP		
		7762994	May 23, 2024	DP		
		8114833	Aug 13, 2025	DP		
		8129343	Dec 05, 2031	DS DP U-2202		
		8536122	Mar 20, 2026	DS DP U-2202		
		8684969	Oct 20, 2025	DP		
		8920383	Jul 17, 2026	DP		
		9108002	Jan 20, 2026	DP		
		9132239	Feb 01, 2032	DP		
		9457154	Sep 29, 2027	DP		
		9616180	Jan 20, 2026	DP		
		9687611	Feb 27, 2027	DP		
		9775953	Jul 17, 2026	DP		
		9861757	Jan 20, 2026	DP		
		RE46363	Aug 03, 2026	DP		
<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637	003	10220155	Jul 17, 2026	DP	D-185	Mar 28, 2025
		10357616	Jan 20, 2026	DP		
		10376652	Jan 20, 2026	DP		
		11097063	Jul 17, 2026	DP		
		11311679	Jan 20, 2026	DP		
		11446443	Oct 20, 2025	DP		
		7762994	May 23, 2024	DP		
		8114833	Aug 13, 2025	DP		
		8129343	Dec 05, 2031	DS DP U-3355		
		8536122	Mar 20, 2026	DS DP U-3355		
		8684969	Oct 20, 2025	DP		
		8920383	Jul 17, 2026	DP		
		9108002	Jan 20, 2026	DP		
		9132239	Feb 01, 2032	DP		
		9457154	Sep 29, 2027	DP		
		9616180	Jan 20, 2026	DP		
		9687611	Feb 27, 2027	DP		
		9775953	Jul 17, 2026	DP		
		9861757	Jan 20, 2026	DP		
		RE46363	Aug 03, 2026	DP		

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<u>SEMAGLUTIDE - OZEMPIC</u>						
N 209637	004 10220155	Jul 17, 2026	DP			
	10335462	Jun 21, 2033	DP	U-2580		
	10357616	Jan 20, 2026	DP			
	10376652	Jan 20, 2026	DP			
	11097063	Jul 17, 2026	DP			
	11311679	Jan 20, 2026	DP			
	11446443	Oct 20, 2025	DP			
	7762994	May 23, 2024	DP			
	8114833	Aug 13, 2025	DP			
	8129343	Dec 05, 2031	DS DP	U-3469		
	8536122	Mar 20, 2026	DS DP	U-3469		
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 29, 2027	DP			
	9616180	Jan 20, 2026	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	9861757	Jan 20, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	001 10086047	Dec 16, 2031	DP			
	10278923	May 02, 2034		U-2628		
	10933120	Mar 15, 2033	DP			
	10960052	Dec 16, 2031	DP			
	11382957	Dec 16, 2031	DP			
	11759501	Mar 15, 2033	DP			
	11759502	Mar 15, 2033	DP			
	11759503	Mar 15, 2033	DP			
	8129343	Dec 05, 2031	DS DP	U-2628		
	8536122	Mar 20, 2026	DS DP	U-2628		
	9278123	Dec 16, 2031	DP	U-2628		
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	002 10086047	Dec 16, 2031	DP			
	10278923	May 02, 2034		U-2628		
	10933120	Mar 15, 2033	DP			
	10960052	Dec 16, 2031	DP			
	11382957	Dec 16, 2031	DP			
	11759501	Mar 15, 2033	DP			
	11759502	Mar 15, 2033	DP			
	11759503	Mar 15, 2033	DP			
	8129343	Dec 05, 2031	DS DP	U-2628		
	8536122	Mar 20, 2026	DS DP	U-2628		
	9278123	Dec 16, 2031	DP	U-2628		
<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	003 10086047	Dec 16, 2031	DP			
	10278923	May 02, 2034		U-2628		

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<u>SEMAGLUTIDE - RYBELSUS</u>						
N 213051	003 10933120	Mar 15, 2033	DP			
	10960052	Dec 16, 2031	DP			
	11382957	Dec 16, 2031	DP			
	11759501	Mar 15, 2033	DP			
	11759502	Mar 15, 2033	DP			
	11759503	Mar 15, 2033	DP			
	8129343	Dec 05, 2031	DS DP U-2628			
	8536122	Mar 20, 2026	DS DP U-2628			
	9278123	Dec 16, 2031	DP U-2628			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	001 10888605	Aug 24, 2038	DP U-3162		NP	Jun 04, 2024
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	002 10888605	Aug 24, 2038	DP U-3162		NP	Jun 04, 2024
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	003 10888605	Aug 24, 2038	DP U-3162		NP	Jun 04, 2024
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	004 10888605	Aug 24, 2038	DP U-3162		D-190	Jul 21, 2026
	11318191	Feb 17, 2041	DP U-3162		NP	Jun 04, 2024
	11752198	Aug 24, 2038	DP U-3162		NPP	Dec 23, 2025
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SEMAGLUTIDE - WEGOVY</u>						
N 215256	005 10888605	Aug 24, 2038	DP U-3162		NP	Jun 04, 2024
	11318191	Feb 17, 2041	DP U-3162		NPP	Dec 23, 2025
	11752198	Aug 24, 2038	DP U-3162			
	8129343	Dec 05, 2031	DS DP			
	8536122	Mar 20, 2026	DS DP			
	9764003	Jun 21, 2033	U-3161			
<u>SETMELANOTIDE ACETATE - IMCIVREE</u>						
N 213793	001 11129869	Jul 04, 2034	DP		I-892	Jun 16, 2025
	8039435	Oct 13, 2027	DS DP		NCE	Nov 25, 2025

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<u>SETMELANOTIDE ACETATE - IMCIVREE</u>						
N 213793	001 9458195	Oct 13, 2027	DS DP		ODE-336	Nov 25, 2027
					ODE-402	Jun 16, 2029
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022127	001 7985418	Oct 27, 2025	DP			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022318	001 9095509	Dec 06, 2030	DP			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N 022318	002 9095509	Dec 06, 2030	DP			
<u>SILDENAFIL CITRATE - REVATIO</u>						
N 021845	001				M-287	Jan 31, 2026
<u>SILDENAFIL CITRATE - REVATIO</u>						
N 022473	001				M-287	Jan 31, 2026
<u>SILDENAFIL CITRATE - REVATIO</u>						
N 203109	001				M-287	Jan 31, 2026
<u>SILDENAFIL CITRATE - LIOREV</u>						
N 214952	001 11337979	Dec 24, 2038	DP U-3582			
	11464778	Dec 24, 2038	DP U-3582			
	11759468	Dec 24, 2038	DP U-3582			
<u>SIMEPREVIR SODIUM - OLYSIO</u>						
N 205123	001 7671032	May 19, 2025	DS DP			
	8148399	Sep 05, 2029	DS DP U-1467			
	8349869	Jul 28, 2026	DS DP U-1467			
	8741926	Jul 28, 2026	DS U-1467			
	8754106	Jul 28, 2026	DS U-1467			
	9040562	Jul 28, 2026	DS DP U-1467			
	9353103	Jul 28, 2026	U-1467			
	9623022	Jul 28, 2026	U-1467			
	9856265	Jul 28, 2026	DS DP U-1467			
<u>SIMVASTATIN - FLOLIPID</u>						
N 206679	001 10300041	Apr 26, 2027	DP			
	9597289	Feb 23, 2030	DP			
<u>SIMVASTATIN - FLOLIPID</u>						
N 206679	002 10300041	Apr 26, 2027	DP			
	9597289	Feb 23, 2030	DP			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	001 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	002 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	003 7326708	Apr 11, 2026	DS DP U-1188			
	7326708*PED	Oct 11, 2026				

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<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	004	7326708	Apr 11, 2026	DS DP U-1188		
		7326708*PED	Oct 11, 2026			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	005	7326708	Apr 11, 2026	DS DP U-1188		
		7326708*PED	Oct 11, 2026			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>						
N 202343	006	7326708	Apr 11, 2026	DS DP U-1188		
		7326708*PED	Oct 11, 2026			
<u>SINCALIDE - SINCALIDE</u>						
N 210850	001	11110063	Apr 20, 2038	DP U-3481		
		11110063	Apr 20, 2038	DP U-3482		
		11110063	Apr 20, 2038	DP U-3483		
		11318100	Apr 20, 2038	DP U-3477		
		11318100	Apr 20, 2038	DP U-3478		
		11318100	Apr 20, 2038	DP U-3479		
		11318100	Apr 20, 2038	DP U-3480		
		11737983	Apr 20, 2038	DP U-3482		
		11737983	Apr 20, 2038	DP U-3689		
<u>SINECATECHINS - VEREGEN</u>						
N 021902	001	7858662	Oct 02, 2026	DP U-172		
		9770406	Jul 12, 2025	DP U-172		
<u>SIPONIMOD - MAYZENT</u>						
N 209884	001	7939519	Aug 27, 2028	DS DP	M-274	Mar 01, 2025
		8492441	Nov 30, 2030	U-2511	NCE	Mar 26, 2024
<u>SIPONIMOD - MAYZENT</u>						
N 209884	002	7939519	Aug 27, 2028	DS DP	M-274	Mar 01, 2025
		8492441	Nov 30, 2030	U-2511	NCE	Mar 26, 2024
<u>SIPONIMOD - MAYZENT</u>						
N 209884	003	7939519	Aug 27, 2028	DS DP	M-274	Mar 01, 2025
		8492441	Nov 30, 2030	U-2511	NCE	Mar 26, 2024
<u>SIROLIMUS - FYARRO</u>						
N 213312	001	10206887	Apr 15, 2030	DP	NP	Nov 22, 2024
		10705070	Mar 05, 2036	DP	ODE-386	Nov 22, 2028
		10973806	Jun 29, 2036	U-3258		
		11497737	Oct 28, 2040	DP		
		8911786	Feb 14, 2029	DP U-3259		
<u>SIROLIMUS - HYFTOR</u>						
N 213478	001				NP	Mar 22, 2025
					ODE-391	Mar 22, 2029
<u>SITAGLIPTIN - ZITUVIO</u>						
N 211566	001	10925871	Feb 25, 2035	DP		
<u>SITAGLIPTIN - ZITUVIO</u>						
N 211566	002	10925871	Feb 25, 2035	DP		

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<u>SITAGLIPTIN - ZITUVIO</u>						
N 211566 003	10925871	Feb 25, 2035	DP			
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 001	7326708	Nov 24, 2026	DS DP U-802		M-187	Dec 04, 2023
	7326708*PED	May 24, 2027			PED	Jun 04, 2024
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 002	7326708	Nov 24, 2026	DS DP U-802		M-187	Dec 04, 2023
	7326708*PED	May 24, 2027			PED	Jun 04, 2024
<u>SITAGLIPTIN PHOSPHATE - JANUVIA</u>						
N 021995 003	7326708	Nov 24, 2026	DS DP U-802		M-187	Dec 04, 2023
	7326708*PED	May 24, 2027			PED	Jun 04, 2024
<u>SODIUM BENZOATE; SODIUM PHENYLACETATE - SODIUM PHENYLACETATE AND SODIUM BENZOATE</u>						
A 217526 001					CGT	Mar 17, 2024
<u>SODIUM CHLORIDE - SODIUM CHLORIDE 23.4%</u>						
A 217796 001					CGT	Jan 23, 2024
<u>SODIUM NITRITE - SODIUM NITRITE</u>						
N 203922 001	8568793	Dec 24, 2031	DS DP			
	9687506	Feb 10, 2030	DP U-3394			
	9687506	Feb 10, 2030	DP U-3395			
<u>SODIUM NITRITE; SODIUM THIOSULFATE - NITHIODOTE</u>						
N 201444 001	10479686	Jul 07, 2030	DP U-3390			
	11753301	Feb 10, 2030	DS DP U-3681			
	8496973	Mar 29, 2031	DS DP U-1419			
	8568793	Dec 24, 2031	DS DP			
	9345724	Jul 07, 2030	DS DP U-2015			
	9585912	Jul 07, 2030	DS DP			
	9687506	Feb 10, 2030	DP U-3394			
	9687506	Feb 10, 2030	DP U-3395			
<u>SODIUM OXYBATE - XYREM</u>						
N 021196 001	10213400	Mar 15, 2033	U-2499		ODE-231	Oct 26, 2025
	10213400*PED	Sep 15, 2033			PED	Apr 26, 2026
	10864181	Mar 15, 2033	U-1532			
	10864181*PED	Sep 15, 2033				
	11253494	Mar 15, 2033	U-3323			
	11253494	Mar 15, 2033	U-3324			
	11253494*PED	Sep 15, 2033				
	7668730	Jun 16, 2024	U-1110	Y		
	7668730*PED	Dec 16, 2024				
	8772306	Mar 15, 2033	U-1532			
	8772306*PED	Sep 15, 2033				
	9050302	Mar 15, 2033	U-1532			
	9050302*PED	Sep 15, 2033				
	9486426	Mar 15, 2033	U-1532			
	9486426*PED	Sep 15, 2033				
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 001	10272062	Jul 21, 2037	DP		NP	May 01, 2026
	10736866	Jul 21, 2037	DP		ODE-431	May 01, 2030

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<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 001	10925844	Feb 28, 2040	DP			
	10952986	Jul 21, 2037	U-3601			
	10973795	Jul 21, 2037	DP			
	11000498	Jul 21, 2037	DP U-3580			
	11052061	Jul 21, 2037	DP			
	11065224	Jul 21, 2037	DP			
	11400065	Jul 21, 2037	U-3579			
	11504347	Jul 21, 2037	DP			
	11583510	Feb 07, 2042	U-3578			
	11602512	Jul 21, 2037	U-3577			
	11602513	Jul 21, 2037	U-3576			
	11766418	Jul 21, 2037	DP			
	11779557	Mar 16, 2042	U-3705			
	11839597	Jul 21, 2037	DP			
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 002	10272062	Jul 21, 2037	DP		NP	May 01, 2026
	10736866	Jul 21, 2037	DP		ODE-431	May 01, 2030
	10925844	Feb 28, 2040	DP			
	10952986	Jul 21, 2037	U-3601			
	10973795	Jul 21, 2037	DP			
	11000498	Jul 21, 2037	DP U-3580			
	11052061	Jul 21, 2037	DP			
	11065224	Jul 21, 2037	DP			
	11400065	Jul 21, 2037	U-3579			
	11504347	Jul 21, 2037	DP			
	11583510	Feb 07, 2042	U-3578			
	11602512	Jul 21, 2037	U-3577			
	11602513	Jul 21, 2037	U-3576			
	11766418	Jul 21, 2037	DP			
	11779557	Mar 16, 2042	U-3705			
	11826335	Jul 21, 2037	U-3751			
	11839597	Jul 21, 2037	DP			
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755 003	10272062	Jul 21, 2037	DP		NP	May 01, 2026
	10736866	Jul 21, 2037	DP		ODE-431	May 01, 2030
	10925844	Feb 28, 2040	DP			
	10952986	Jul 21, 2037	U-3601			
	10973795	Jul 21, 2037	DP			
	11000498	Jul 21, 2037	DP U-3580			
	11052061	Jul 21, 2037	DP			
	11065224	Jul 21, 2037	DP			
	11400065	Jul 21, 2037	U-3579			
	11504347	Jul 21, 2037	DP			
	11583510	Feb 07, 2042	U-3578			
	11602512	Jul 21, 2037	U-3577			
	11602513	Jul 21, 2037	U-3576			
	11766418	Jul 21, 2037	DP			
	11779557	Mar 16, 2042	U-3705			
	11839597	Jul 21, 2037	DP			

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<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755	003	10272062	Jul 21, 2037	DP	NP	May 01, 2026
		10736866	Jul 21, 2037	DP	ODE-431	May 01, 2030
		10925844	Feb 28, 2040	DP		
		10952986	Jul 21, 2037	U-3601		
		10973795	Jul 21, 2037	DP		
		11000498	Jul 21, 2037	DP U-3580		
		11052061	Jul 21, 2037	DP		
		11065224	Jul 21, 2037	DP		
		11400065	Jul 21, 2037	U-3579		
		11504347	Jul 21, 2037	DP		
		11583510	Feb 07, 2042	U-3578		
		11602512	Jul 21, 2037	U-3577		
		11602513	Jul 21, 2037	U-3576		
		11766418	Jul 21, 2037	DP		
		11779557	Mar 16, 2042	U-3705		
		11839597	Jul 21, 2037	DP		
<u>SODIUM OXYBATE - LUMRYZ</u>						
N 214755	004	10272062	Jul 21, 2037	DP	NP	May 01, 2026
		10736866	Jul 21, 2037	DP	ODE-431	May 01, 2030
		10925844	Feb 28, 2040	DP		
		10952986	Jul 21, 2037	U-3601		
		10973795	Jul 21, 2037	DP		
		11000498	Jul 21, 2037	DP U-3580		
		11052061	Jul 21, 2037	DP		
		11065224	Jul 21, 2037	DP		
		11400065	Jul 21, 2037	U-3579		
		11504347	Jul 21, 2037	DP		
		11583510	Feb 07, 2042	U-3578		
		11602512	Jul 21, 2037	U-3577		
		11602513	Jul 21, 2037	U-3576		
		11766418	Jul 21, 2037	DP		
		11779557	Mar 16, 2042	U-3705		
		11839597	Jul 21, 2037	DP		
<u>SODIUM PHENYL BUTYRATE - OLPRUVA</u>						
N 214860	001	11154521	Oct 17, 2036	DP		
		11202767	Oct 17, 2036	U-3502		
		11433041	Oct 17, 2036	DP		
<u>SODIUM PHENYL BUTYRATE - OLPRUVA</u>						
N 214860	002	11154521	Oct 17, 2036	DP		
		11202767	Oct 17, 2036	U-3502		
		11433041	Oct 17, 2036	DP		
<u>SODIUM PHENYL BUTYRATE - OLPRUVA</u>						
N 214860	003	11154521	Oct 17, 2036	DP		
		11202767	Oct 17, 2036	U-3502		
		11433041	Oct 17, 2036	DP		
<u>SODIUM PHENYL BUTYRATE - OLPRUVA</u>						
N 214860	004	11154521	Oct 17, 2036	DP		
		11202767	Oct 17, 2036	U-3502		

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<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860	004 11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860	005 11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036	U-3502			
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE - OLPRUVA</u>						
N 214860	006 11154521	Oct 17, 2036	DP			
	11202767	Oct 17, 2036	U-3502			
	11433041	Oct 17, 2036	DP			
<u>SODIUM PHENYLBUTYRATE; TAURURSODIOL - RELYVRIO</u>						
N 216660	001 10251896	Dec 24, 2033	U-3460		NCE	Sep 29, 2027
	10857162	Dec 24, 2033	U-3460		ODE-411	Sep 29, 2029
	11071742	Dec 24, 2033	DP			
	11583542	Jul 27, 2040	DP			
	9872865	Dec 24, 2033	U-3460			
<u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</u>						
N 021892	001 7687075	Jun 22, 2028	DS DP			
<u>SODIUM THIOSULFATE - SODIUM THIOSULFATE</u>						
N 203923	001 10479686	Jul 07, 2030	DP U-3390			
	11753301	Feb 10, 2030	DS DP U-3682			
	8496973	Mar 29, 2031	DS DP U-1419			
	9345724	Jul 07, 2030	DS DP U-2015			
	9585912	Jul 07, 2030	DS DP			
<u>SODIUM THIOSULFATE - PEDMARK</u>						
N 212937	001 10596190	Jan 05, 2038	U-3443	Y	NP	Sep 20, 2025
	11291728	Jul 01, 2039	DP		ODE-384	Sep 20, 2029
	11510984	Jul 01, 2039	DP			
	11617793	Jul 01, 2039	DP			
<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078	001 10300087	Oct 14, 2035	DS U-2312			
	10335432	Feb 10, 2032	U-2312			
	10398730	Feb 10, 2032	U-2312			
	10413569	Feb 10, 2032	DS			
	10695365	Oct 22, 2033	DS			
	11406662	Feb 10, 2032	DS			
	11738044	Oct 14, 2035	U-2312			
	8802152	Apr 19, 2032	DS			
	8808750	Feb 10, 2032	U-2312			
	8877255	Oct 22, 2033	DS			
	9592253	Oct 14, 2035	DS U-2312			
	9844567	Feb 10, 2032	U-2312			
	9861658	Feb 10, 2032	U-2312			
	9913860	Oct 22, 2033	DS U-2312			
<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078	002 10300087	Oct 14, 2035	DS U-2312			
	10398730	Feb 10, 2032	U-2312			

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<u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u>						
N 207078 002	10413569	Feb 10, 2032	DS			
	10695365	Oct 22, 2033	DS			
	11406662	Feb 10, 2032	DS			
	11738044	Oct 14, 2035		U-2312		
	8802152	Apr 19, 2032	DS			
	8808750	Feb 10, 2032		U-2312		
	8877255	Oct 22, 2033	DS			
	9592253	Oct 14, 2035	DS	U-2312		
	9844567	Feb 10, 2032		U-2312		
	9861658	Feb 10, 2032		U-2312		
	9913860	Oct 22, 2033	DS	U-2312		
<u>SOFOSEBUVIR - SOVALDI</u>						
N 204671 001	7964580	Mar 26, 2029	DS DP U-1470		ODE*	Aug 28, 2026
	7964580*PED	Sep 26, 2029			ODE-135	Apr 07, 2024
	8334270	Mar 21, 2028	DS DP U-1470		PED	Oct 07, 2024
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9549941	Mar 26, 2029		U-1958		
	9549941*PED	Sep 26, 2029				
<u>SOFOSEBUVIR - SOVALDI</u>						
N 204671 002	7964580	Mar 26, 2029	DS DP U-1470		ODE*	Apr 07, 2024
	7964580*PED	Sep 26, 2029			ODE*	Aug 28, 2026
	8334270	Mar 21, 2028	DS DP U-1470		PED	Oct 07, 2024
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				

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<u>SOFOSBUVIR - SOVALDI</u>						
N 212480 001	7964580	Mar 26, 2029	DS DP U-1470		ODE*	Apr 07, 2024
	7964580*PED	Sep 26, 2029			ODE-258	Aug 28, 2026
	8334270	Mar 21, 2028	DS DP U-1470		PED	Oct 07, 2024
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOSBUVIR - SOVALDI</u>						
N 212480 002	7964580	Mar 26, 2029	DS DP U-1470		ODE*	Apr 07, 2024
	7964580*PED	Sep 26, 2029			ODE-258	Aug 28, 2026
	8334270	Mar 21, 2028	DS DP U-1470		PED	Oct 07, 2024
	8334270*PED	Sep 21, 2028				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341 001	10086011	Jan 30, 2034	U-1470		M-264	Jul 14, 2023
	10086011*PED	Jul 30, 2034			M-277	Apr 27, 2025
	11116783	Jan 30, 2034	DP U-1470		ODE-293	Mar 19, 2027
	11116783*PED	Jul 30, 2034			ODE-376	Jun 10, 2028
	7964580	Mar 26, 2029	DS DP U-1470		PED	Jan 14, 2024
	7964580*PED	Sep 26, 2029			PED	Sep 19, 2027
	8334270	Mar 21, 2028	DS DP U-1470		PED	Dec 10, 2028
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			

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<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341 001	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9757406	Jan 30, 2034	DP			
	9757406*PED	Jul 30, 2034				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 208341 002	10086011	Jan 30, 2034	U-1470		M-264	Jul 14, 2023
	10086011*PED	Jul 30, 2034			M-277	Apr 27, 2025
	11116783	Jan 30, 2034	DP U-1470		ODE-293	Mar 19, 2027
	11116783*PED	Jul 30, 2034			ODE-376	Jun 10, 2028
	7964580	Mar 26, 2029	DS DP U-1470		PED	Jan 14, 2024
	7964580*PED	Sep 26, 2029			PED	Sep 19, 2027
	8334270	Mar 21, 2028	DS DP U-1470		PED	Dec 10, 2028
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
	9757406	Jan 30, 2034	DP			
	9757406*PED	Jul 30, 2034				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 001	11116783	Jan 30, 2034	DP U-1470		M-277	Apr 27, 2025
	11116783*PED	Jul 30, 2034			ODE-376	Jun 10, 2028

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<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 001	11707479	Jan 30, 2034	DP U-1470		PED	Dec 10, 2028
	11707479*PED	Jul 30, 2034				
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 002	11116783	Jan 30, 2034	DP U-1470		M-277	Apr 27, 2025
	11116783*PED	Jul 30, 2034			ODE-376	Jun 10, 2028
	11707479	Jan 30, 2034	DP U-1470		PED	Dec 10, 2028
	11707479*PED	Jul 30, 2034				
	7964580	Mar 26, 2029	DS DP U-1470			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-1470			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-1470			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-1470			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-1470			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-1470			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	U-1470			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DP U-1470			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-1470			
	8921341*PED	May 16, 2033				

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<u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u>						
N 214187 002	8940718	Nov 16, 2032	DS DP U-1470			
	8940718*PED	May 16, 2033				
	9085573	Mar 21, 2028	DS DP U-1470			
	9085573*PED	Sep 21, 2028				
	9284342	Sep 13, 2030	DS DP U-1470			
	9284342*PED	Mar 13, 2031				
<u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195 001	10912814	Jun 01, 2037	DP			
	11116783	Jan 30, 2034	DP U-2039			
	11116783	Jan 30, 2034	DP U-2040			
	11116783*PED	Jul 30, 2034				
	11338007	Jun 01, 2037	DP U-2039			
	11338007	Jun 01, 2037	DP U-2040			
	11338007*PED	Dec 01, 2037				
	7964580	Mar 26, 2029	DS DP U-2039			
	7964580	Mar 26, 2029	DS DP U-2040			
	7964580*PED	Sep 26, 2029				
	8334270	Mar 21, 2028	DS DP U-2039			
	8334270	Mar 21, 2028	DS DP U-2040			
	8334270*PED	Sep 21, 2028				
	8575135	Nov 16, 2032	DS DP U-2039			
	8575135	Nov 16, 2032	DS DP U-2040			
	8575135*PED	May 16, 2033				
	8580765	Mar 21, 2028	DS DP U-2039			
	8580765	Mar 21, 2028	DS DP U-2040			
	8580765*PED	Sep 21, 2028				
	8618076	Dec 11, 2030	DS DP U-2039			
	8618076	Dec 11, 2030	DS DP U-2040			
	8618076*PED	Jun 11, 2031				
	8633309	Mar 26, 2029	DS DP U-2039			
	8633309	Mar 26, 2029	DS DP U-2040			
	8633309*PED	Sep 26, 2029				
	8735372	Mar 21, 2028	DS DP U-2039			
	8735372	Mar 21, 2028	DS DP U-2040			
	8735372*PED	Sep 21, 2028				
	8889159	Mar 26, 2029	DS DP U-2039			
	8889159	Mar 26, 2029	DS DP U-2040			
	8889159*PED	Sep 26, 2029				
	8921341	Nov 16, 2032	DS DP U-2039			
	8921341	Nov 16, 2032	DS DP U-2040			
	8921341*PED	May 16, 2033				
	8940718	Nov 16, 2032	DS DP U-2039			
	8940718	Nov 16, 2032	DS DP U-2040			
	8940718*PED	May 16, 2033				
	8957046	Mar 21, 2028	U-2039			
	8957046	Mar 21, 2028	U-2040			
	9085573	Mar 21, 2028	DS DP U-2039			
	9085573	Mar 21, 2028	DS DP U-2040			
	9085573*PED	Sep 21, 2028				

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<u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u>						
N 209195 001	9284342	Sep 13, 2030	DS DP U-2039			
	9284342	Sep 13, 2030	DS DP U-2040			
	9284342*PED	Mar 13, 2031				
	9296782	Jul 17, 2034	DS DP			
	9585906	Mar 21, 2028	DS DP U-2039			
	9585906	Mar 21, 2028	DS DP U-2040			
	9868745	Nov 16, 2032	DS DP			
<u>SOLIFENACIN SUCCINATE - VESICARE LS</u>						
N 209529 001	9918970	May 18, 2031	DP			
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230 001	10195151	Sep 05, 2037	DP		NCE	Jun 17, 2024
	10351517	Jun 07, 2026	U-2548		ODE-254	Jun 17, 2026
	10512609	Sep 05, 2037	U-2548			
	10912754	Jun 01, 2038	U-3082			
	10940133	Mar 19, 2040	U-3099			
	10959976	Jun 01, 2038	U-3151			
	11160779	Mar 19, 2040	U-3521			
	11439597	Sep 05, 2037	DP			
	11560354	Mar 06, 2039	DP U-3520			
	11648232	Jun 01, 2038	U-3602			
	11753368	Jun 07, 2026	U-2548			
	11771666	Dec 30, 2042	U-3693			
	11771667	Dec 30, 2042	U-3693			
	11779554	Dec 30, 2042	U-3693			
	11793776	Dec 30, 2042	U-3693			
	11839598	Mar 19, 2040	U-3765			
	11839599	Mar 19, 2040	U-3764			
	11850226	Mar 19, 2040	U-3775			
	11850227	Mar 19, 2040	U-3775			
	11850228	Mar 19, 2040	U-3775			
	8440715	Jun 11, 2031	U-2548			
	8877806	Jun 07, 2026	U-2548			
	9604917	Jun 07, 2026	U-2548			
<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230 002	10195151	Sep 05, 2037	DP		NCE	Jun 17, 2024
	10351517	Jun 07, 2026	U-2548		ODE-254	Jun 17, 2026
	10512609	Sep 05, 2037	U-2548			
	10912754	Jun 01, 2038	U-3082			
	10959976	Jun 01, 2038	U-3151			
	11160779	Mar 19, 2040	U-3521			
	11439597	Sep 05, 2037	DP			
	11560354	Mar 06, 2039	DP U-3520			
	11648232	Jun 01, 2038	U-3602			
	11753368	Jun 07, 2026	U-2548			
	11771666	Dec 30, 2042	U-3693			
	11771667	Dec 30, 2042	U-3693			
	11779554	Dec 30, 2042	U-3693			
	11793776	Dec 30, 2042	U-3693			

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<u>SOLRIAMFETOL HYDROCHLORIDE - SUNOSI</u>						
N 211230	002	11839598	Mar 19, 2040	U-3765		
		11839599	Mar 19, 2040	U-3764		
		11850226	Mar 19, 2040	U-3775		
		11850227	Mar 19, 2040	U-3775		
		11850228	Mar 19, 2040	U-3775		
		8440715	Jun 11, 2031	U-2548		
		8877806	Jun 07, 2026	U-2548		
		9604917	Jun 07, 2026	U-2548		
<u>SONIDEGIB PHOSPHATE - ODOMZO</u>						
N 205266	001	10266523	Mar 30, 2036	DS DP		
		8063043	Sep 15, 2029	DS DP		
		8178563	Jul 24, 2029	DS	U-1722	
<u>SORAFENIB TOSYLATE - NEXAVAR</u>						
N 021923	001	8877933	Dec 24, 2027	DS DP	U-1624	
		9737488	Sep 10, 2028	DP	U-1480	
		9737488	Sep 10, 2028	DP	U-1696	
		9737488	Sep 10, 2028	DP	U-2107	
<u>SOTAGLIFLOZIN - INPEFA</u>						
N 216203	001	7781577	May 04, 2028	DS DP	U-3628	NCE May 26, 2028
		8217156	Oct 07, 2030	DS DP		
		8476413	May 29, 2028	DS DP	U-3628	
<u>SOTAGLIFLOZIN - INPEFA</u>						
N 216203	002	7781577	May 04, 2028	DS DP	U-3628	NCE May 26, 2028
		8217156	Oct 07, 2030	DS DP		
		8476413	May 29, 2028	DS DP	U-3628	
<u>SOTALOL HYDROCHLORIDE - SOTALOL HYDROCHLORIDE</u>						
N 022306	001	10512620	Aug 14, 2038		U-2769	
		10512620	Aug 14, 2038		U-3547	
		10799138	Apr 05, 2039		U-3125	
		10799138	Apr 05, 2039		U-3549	
		11583216	Aug 21, 2039		U-3549	
		11696902	Aug 14, 2038		U-2769	
<u>SOTALOL HYDROCHLORIDE - SOTYLIZE</u>						
N 205108	001	10206895	Apr 01, 2034	DP	U-2096	
		10206895	Apr 01, 2034	DP	U-2494	
		11013703	Apr 01, 2034	DP		
		11850222	Nov 19, 2034		U-2096	
		11850222	Nov 19, 2034		U-2494	
		9724297	Aug 31, 2035	DP	U-2096	
<u>SOTORASIB - LUMAKRAS</u>						
N 214665	001	10519146	May 21, 2038	DS DP		NCE May 28, 2026
		11236091	May 20, 2040	DS DP	U-3306	ODE-352 May 28, 2028
		11426404	Sep 15, 2040		U-3306	
		11827635	May 20, 2040	DS DP	U-3306	
<u>SOTORASIB - LUMAKRAS</u>						
N 214665	002	10519146	May 21, 2038	DS DP		NCE May 28, 2026

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<u>SOTORASIB - LUMAKRAS</u>						
N 214665	002 11236091	May 20, 2040	DS DP U-3306		ODE-352	May 28, 2028
	11426404	Sep 15, 2040	U-3306			
	11827635	May 20, 2040	DS DP U-3306			
<u>SPARSENTAN - FILSPARI</u>						
N 216403	001 9993461	Mar 29, 2030	U-3269		NCE ODE-389	Feb 17, 2028 Feb 17, 2030
<u>SPARSENTAN - FILSPARI</u>						
N 216403	002 9993461	Mar 29, 2030	U-3269		NCE ODE-389	Feb 17, 2028 Feb 17, 2030
<u>SPINOSAD - NATROBA</u>						
N 022408	001 9895388	Nov 25, 2033	U-3365		I-858	Apr 28, 2024
<u>SPIRONOLACTONE - CAROSPIR</u>						
N 209478	001 10493083	Oct 28, 2036	DP			
	10624906	Oct 28, 2036	DP			
	10660907	Oct 28, 2036	DP			
	10888570	Oct 28, 2036	DP			
	11389461	Oct 28, 2036	DP			
	11395828	Oct 28, 2036	U-2109			
	11395828	Oct 28, 2036	U-3401			
	11395828	Oct 28, 2036	U-3402			
	11491166	Oct 28, 2036	DP			
	9757394	Oct 28, 2036	DP U-2109			
<u>STIRIPENTOL - DIACOMIT</u>						
N 206709	001				M-281 ODE-198 ODE-403	Jul 14, 2025 Aug 20, 2025 Jul 14, 2029
<u>STIRIPENTOL - DIACOMIT</u>						
N 206709	002				M-281 ODE-198 ODE-403	Jul 14, 2025 Aug 20, 2025 Jul 14, 2029
<u>STIRIPENTOL - DIACOMIT</u>						
N 207223	001				NPP ODE-198 ODE-403	Jul 14, 2025 Aug 20, 2025 Jul 14, 2029
<u>STIRIPENTOL - DIACOMIT</u>						
N 207223	002				NPP ODE-198 ODE-403	Jul 14, 2025 Aug 20, 2025 Jul 14, 2029
<u>SUFENTANIL CITRATE - DSUVIA</u>						
N 209128	001 10245228	Jan 05, 2027	DP U-1351			
	10342762	Jan 05, 2027	DP			
	10507180	Jan 05, 2027	DP U-1351			
	10896751	Mar 16, 2030	DP			
	11672738	Feb 02, 2038	DP			
	11676691	Mar 16, 2030	DP			
	8202535	Oct 22, 2030	U-1351			

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>SUFENTANIL CITRATE - DSUVIA</u>						
N 209128	001 8226978	Jan 05, 2027	DP U-1351			
	8231900	Jan 05, 2027	DP			
	8252328	Jan 05, 2027	DP			
	8252329	Jan 05, 2027	DP			
	8535714	Jan 05, 2027	DP U-1351			
	8574189	Mar 16, 2030	DP			
	8778393	Jan 05, 2027	U-1351			
	8778394	Jan 05, 2027	U-1351			
	8865211	Jan 05, 2027	U-1351			
	8865743	Oct 22, 2030	U-1351			
	8945592	Jul 29, 2031	DP			
	9320710	Jan 05, 2027	U-1351			
	9744129	Jan 05, 2027	DP			
<u>SUGAMMADEX SODIUM - BRIDION</u>						
N 022225	001 RE44733	Jan 27, 2026	DS DP U-1794		M-291	Jan 22, 2024
					NPP	Jun 25, 2024
<u>SUGAMMADEX SODIUM - BRIDION</u>						
N 022225	002 RE44733	Jan 27, 2026	DS DP U-1794		M-291	Jan 22, 2024
					NPP	Jun 25, 2024
<u>SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES - LUMASON</u>						
N 203684	001 10232061	Jul 06, 2038	DP			
	10335502	Jul 06, 2038	DP			
	11723869	May 15, 2039	DP U-3666			
	11723869	May 15, 2039	DP U-3667			
	11723869	May 15, 2039	DP U-3668			
<u>SUMATRIPTAN - TOSYMRA</u>						
N 210884	001 10603305	Jun 16, 2030	DP U-1719			
	11337962	Jun 16, 2030	DP U-1719			
	8268791	May 09, 2026	DP			
	8440631	May 09, 2026	DP U-1719			
	9211282	Jul 19, 2031	DP U-1719			
	9283280	May 09, 2026	DP			
	9610280	Jun 16, 2030	DP U-1719			
	9974770	Jun 16, 2030	DP U-1719			
<u>SUMATRIPTAN SUCCINATE - SUMAVEL DOSEPRO</u>						
N 022239	001 7776007	Nov 22, 2026	DP			
	7901385	Jul 31, 2026	DP			
	8287489	Dec 06, 2024	DP			
<u>SUMATRIPTAN SUCCINATE - ALSUMA</u>						
N 022377	001 7811254	Aug 26, 2027	DP U-1083			
<u>SUMATRIPTAN SUCCINATE - ZECUITY</u>						
N 202278	001 7973058	Apr 12, 2027	U-1328			
	8155737	Apr 12, 2027	U-1328			
	8366600	Apr 21, 2029	U-1327			
	8470853	Apr 12, 2027	U-1328			
	8597272	Apr 12, 2027	DP			
	8983594	Nov 19, 2030	DP U-1328			

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<u>SUMATRIPTAN SUCCINATE - ZECUITY</u>						
N 202278	001 9272137	Sep 07, 2027	DP			
	9327114	Oct 08, 2032	DP U-1328			
	9427578	Apr 12, 2027	DP U-1328			
<u>SUMATRIPTAN SUCCINATE - ONZETRA XSAIL</u>						
N 206099	001 10076614	Oct 20, 2034	DP			
	10076615	Jul 30, 2029	U-2010			
	10076615	Jul 30, 2029	U-2011			
	10076615	Jul 30, 2029	U-2404			
	10124132	Mar 06, 2027	DP U-1719			
	10124132	Mar 06, 2027	DP U-2010			
	10124132	Mar 06, 2027	DP U-2011			
	10398859	Dec 19, 2027	DP			
	10478574	Nov 04, 2033	U-2404			
	10722667	Dec 30, 2028	DP			
	11571531	Feb 23, 2026	U-1809			
	7975690	Aug 18, 2025	DP U-1809			
	8550073	Oct 22, 2029	DP			
	8590530	Sep 15, 2025	DP U-1809			
	8875704	Apr 07, 2028	DP U-1809			
	8899229	Aug 18, 2030	DP			
	8978647	Dec 06, 2030	DP			
	9108015	Sep 15, 2025	DP			
	9119932	Apr 23, 2024	DP			
	9649456	Oct 21, 2030	DP U-1719			
	9649456	Oct 21, 2030	DP U-2010			
	9649456	Oct 21, 2030	DP U-2011			
<u>SUMATRIPTAN SUCCINATE - ZEMBRACE SYMTOUCH</u>						
N 208223	001 10537554	Jan 29, 2036	U-72			
	11364224	Jan 29, 2036	U-72			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	001 10098892	May 29, 2033	DP			
	7951797	Nov 20, 2029	DS DP U-620			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	002 10098892	May 29, 2033	DP			
	7951797	Nov 20, 2029	DS DP U-620			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	003 10098892	May 29, 2033	DP			
	7951797	Nov 20, 2029	DS DP U-620			
<u>SUVOREXANT - BELSOMRA</u>						
N 204569	004 10098892	May 29, 2033	DP			
	7951797	Nov 20, 2029	DS DP U-620			
<u>TACROLIMUS - PROGRAF</u>						
N 050708	001				ODE-294	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050708	002				ODE-294	May 24, 2025

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<u>TACROLIMUS - PROGRAF</u>						
N 050708	002				ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050708	003				ODE-294 ODE-360	May 24, 2025 Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 050709	001				ODE-294 ODE-360	May 24, 2025 Jul 16, 2028
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	001				ODE*	May 24, 2025
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	002				ODE*	May 24, 2025
<u>TACROLIMUS - ASTAGRAF XL</u>						
N 204096	003				ODE*	May 24, 2025
<u>TACROLIMUS - ENVARUSUS XR</u>						
N 206406	001	10166190	May 30, 2028	DP		
		10548880	Aug 30, 2024	U-2677		
		10548880	Aug 30, 2024	U-2678		
		10864199	May 30, 2028	U-2677		
		10864199	May 30, 2028	U-2678		
		11077096	Aug 30, 2024	DP		
		11110081	May 30, 2028	U-2678		
		11123331	May 30, 2028	U-2677		
		11419823	May 30, 2028	DP		
		7994214	Aug 30, 2024	DP		
		8486993	Aug 30, 2024	DP U-1752		
		8586084	Aug 30, 2024	U-1752		
		8591946	Aug 30, 2024	DP		
		8617599	Aug 30, 2024	DP		
		8623410	Aug 30, 2024	DP		
		8623411	Aug 30, 2024	U-1752		
		8664239	Aug 30, 2028	U-1752		
		8664239	Aug 30, 2028	U-2677		
		8664239	Aug 30, 2028	U-2678		
		8685998	Aug 30, 2028	DP U-1752		
		8685998	Aug 30, 2028	DP U-2677		
		8685998	Aug 30, 2028	DP U-2678		
		8889185	Aug 30, 2024	U-1752		
		8889186	Aug 30, 2024	U-1752		
		9161907	Aug 30, 2024	DP U-1752		
		9549918	May 30, 2028	DP		
		9757362	Aug 30, 2024	DP		
		9763920	Aug 30, 2024	DP		
<u>TACROLIMUS - ENVARUSUS XR</u>						
N 206406	002	10166190	May 30, 2028	DP		
		10548880	Aug 30, 2024	U-2677		
		10548880	Aug 30, 2024	U-2678		
		10864199	May 30, 2028	U-2677		

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<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406 002	10864199	May 30, 2028	U-2678			
	11077096	Aug 30, 2024	DP			
	11110081	May 30, 2028	U-2678			
	11123331	May 30, 2028	U-2677			
	11419823	May 30, 2028	DP			
	7994214	Aug 30, 2024	DP			
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	8664239	Aug 30, 2028	U-1752			
	8664239	Aug 30, 2028	U-2677			
	8664239	Aug 30, 2028	U-2678			
	8685998	Aug 30, 2028	DP U-1752			
	8685998	Aug 30, 2028	DP U-2677			
	8685998	Aug 30, 2028	DP U-2678			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			
	9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406 003	10166190	May 30, 2028	DP			
	10548880	Aug 30, 2024	U-2677			
	10548880	Aug 30, 2024	U-2678			
	10864199	May 30, 2028	U-2677			
	10864199	May 30, 2028	U-2678			
	11077096	Aug 30, 2024	DP			
	11110081	May 30, 2028	U-2678			
	11123331	May 30, 2028	U-2677			
	11419823	May 30, 2028	DP			
	7994214	Aug 30, 2024	DP			
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	8664239	Aug 30, 2028	U-1752			
	8664239	Aug 30, 2028	U-2677			
	8664239	Aug 30, 2028	U-2678			
	8685998	Aug 30, 2028	DP U-1752			
	8685998	Aug 30, 2028	DP U-2677			
	8685998	Aug 30, 2028	DP U-2678			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			

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<u>TACROLIMUS - ENVARSUS XR</u>						
N 206406	003 9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<u>TACROLIMUS - PROGRAF</u>						
N 210115	001				ODE-269	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TACROLIMUS - PROGRAF</u>						
N 210115	002				ODE-269	May 24, 2025
					ODE-360	Jul 16, 2028
<u>TADALAFIL - TADLIQ</u>						
N 214522	001 11382917	Dec 24, 2038	DP U-3397			
	11666576	Dec 24, 2038	DP U-3397			
<u>TAFAMIDIS - VYNDAMAX</u>						
N 212161	001 7214695	Dec 19, 2024	DS DP		NCE	May 03, 2024
	7214696	Dec 19, 2024	U-2524		ODE-237	May 03, 2026
	9770441	Aug 31, 2035	DS DP U-2524			
<u>TAFAMIDIS MEGLUMINE - VYNDAOEL</u>						
N 211996	001 7214695	Dec 19, 2024	DS DP		NCE	May 03, 2024
	7214696	Dec 19, 2024	U-2524		ODE-237	May 03, 2026
<u>TAFENOQUINE SUCCINATE - ARAKODA</u>						
N 210607	001 10342791	Dec 02, 2035	U-2582			
	10888558	Dec 02, 2035	U-2582			
	11744828	Dec 02, 2035	U-2582			
<u>TAFENOQUINE SUCCINATE - KRINTAFEL</u>						
N 210795	001				ODE-201	Jul 20, 2025
<u>TAFLUPROST - ZIOPTAN</u>						
N 202514	001 10864159	May 28, 2029	DP U-778			
	9999593	May 28, 2029	DP			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	001 10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029	U-3651			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	002 10189837	Oct 20, 2031	DS DP			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029	U-2437			
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651	003 10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029	U-3651			

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<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651 003	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029		U-2437		
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651 004	10189837	Oct 20, 2031	DS DP			
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029		U-2437		
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651 005	10189837	Oct 20, 2031	DS DP		I-920	Jun 20, 2026
	10780088	Jul 27, 2029		U-3651		
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029		U-2437		
<u>TALAZOPARIB TOSYLATE - TALZENNA</u>						
N 211651 006	10189837	Oct 20, 2031	DS DP			
	10780088	Jul 27, 2029		U-3651		
	8012976	Oct 19, 2029	DS DP			
	8420650	Jul 27, 2029	DS DP			
	8735392	Oct 20, 2031	DS DP			
	9820985	Jul 27, 2029		U-2437		
<u>TALC - STERITALC</u>						
N 205555 001					ODE-143	May 01, 2024
					ODE-191	May 01, 2024
<u>TALC - STERITALC</u>						
N 205555 002					ODE-143	May 01, 2024
					ODE-191	May 01, 2024
<u>TALC - STERITALC</u>						
N 205555 003					ODE-143	May 01, 2024
					ODE-191	May 01, 2024
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА</u>						
N 022304 001	7994364	Jun 27, 2025	DS DP U-931		NPP	Jul 03, 2026
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА</u>						
N 022304 002	7994364	Jun 27, 2025	DS DP U-931		NPP	Jul 03, 2026
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА</u>						
N 022304 003	7994364	Jun 27, 2025	DS DP U-931		NPP	Jul 03, 2026
<u>TAPENTADOL HYDROCHLORIDE - NUCYNТА ER</u>						
N 200533 001	11344512	Apr 21, 2028		U-3391		
	11344512	Apr 21, 2028		U-3392		
	7994364	Jun 27, 2025	DS DP U-1178			
	7994364	Jun 27, 2025	DS DP U-1276			
	8114383	Oct 10, 2024	DP		Y	
	8536130	Sep 22, 2028		U-1276		

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<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	001	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8114383	Oct 10, 2024	DP	Y	
		8536130	Sep 22, 2028	U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	002	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8114383	Oct 10, 2024	DP	Y	
		8536130	Sep 22, 2028	U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	003	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8114383	Oct 10, 2024	DP	Y	
		8536130	Sep 22, 2028	U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	004	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8114383	Oct 10, 2024	DP	Y	
		8536130	Sep 22, 2028	U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>						
N 200533	005	11344512	Apr 21, 2028	U-3391		
		11344512	Apr 21, 2028	U-3392		
		7994364	Jun 27, 2025	DS DP U-1178		
		7994364	Jun 27, 2025	DS DP U-1276		
		8114383	Oct 10, 2024	DP	Y	
		8536130	Sep 22, 2028	U-1276		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N 203794	001	7994364	Jun 27, 2025	DS DP U-1289	NPP	Jul 03, 2026
<u>TAPINAROF - VTAMA</u>						
N 215272	001	10195160	May 19, 2036	DP	NCE	May 23, 2027
		10426743	May 19, 2036	U-2625		
		10647649	Nov 13, 2038	DS		
		11458108	May 19, 2036	DP		
		11590088	Nov 13, 2039	U-2625		
		11597692	Nov 13, 2038	DS DP		
		11612573	May 19, 2036	U-2625		
		11617724	May 19, 2036	DP		
		11622945	May 19, 2036	DP		

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<u>TASIMELTEON - HETLIOZ</u>						
N 205677 001	10071977	Feb 12, 2035	DS DP		ODE-330	Dec 01, 2027
	10149829	Jan 25, 2033		U-2477		
	10149829	Jan 25, 2033		U-3006		
	10179119	Aug 29, 2035		U-3003		
	10376487	Jul 27, 2035		U-2615		
	10376487	Jul 27, 2035		U-3007		
	10449176	Jan 25, 2033		U-2149		
	10610510	Jan 25, 2033		U-2805		
	10610510	Jan 25, 2033		U-3009		
	10610511	Oct 10, 2034		U-2615		
	10610511	Oct 10, 2034		U-3007		
	10829465	Feb 12, 2035	DS DP			
	10945988	Jan 25, 2033		U-2149		
	10980770	Jan 25, 2033		U-3106		
	10980770	Jan 25, 2033		U-3107		
	11141400	Oct 10, 2034		U-2615		
	11141400	Oct 10, 2034		U-3007		
	11266622	Aug 29, 2035		U-3003		
	11285129	Jan 25, 2033		U-3342		
	11285129	Jan 25, 2033		U-3343		
	11566011	Feb 12, 2035	DS DP			
	11633377	Jan 25, 2033		U-2149		
	11633377	Jan 25, 2033		U-3003		
	11759446	Feb 21, 2041		U-3003		
	11760740	Feb 12, 2035	DS DP			
	11786502	Oct 10, 2034		U-3007		
	11786502	Oct 10, 2034		U-3739		
	11833130	Jan 25, 2033		U-2149		
	11833130	Jan 25, 2033		U-3003		
	11850229	Jan 25, 2033		U-3342		
	11850229	Jan 25, 2033		U-3343		
	9060995	Jan 25, 2033		U-1710		
	9539234	Jan 25, 2033		U-1934		
	9539234	Jan 25, 2033		U-3004		
	9549913	Jan 25, 2033		U-1486		
	9730910	May 17, 2034		U-2085		
	9730910	May 17, 2034		U-3005		
	9855241	Jan 25, 2033		U-2149		
	RE46604	Jan 25, 2033		U-2147		
<u>TASIMELTEON - HETLIOZ LQ</u>						
N 214517 001	10071977	Feb 12, 2035	DS DP		ODE-329	Dec 01, 2027
	10149829	Jan 25, 2033		U-3006		
	10179119	Aug 29, 2035		U-3003		
	10376487	Jul 27, 2035		U-3007		
	10610510	Jan 25, 2033		U-3009		
	10610511	Oct 10, 2034		U-3007		
	10829465	Feb 12, 2035	DS DP			
	10980770	Jan 25, 2033		U-3106		
	11141400	Oct 10, 2034		U-3007		

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<u>TASIMELTEON - HETLIOZ LQ</u>						
N 214517 001	11202770	Dec 11, 2040	DP			
	11266622	Aug 29, 2035	U-3003			
	11285129	Jan 25, 2033	U-3342			
	11566011	Feb 12, 2035	DS DP			
	11633377	Jan 25, 2033	U-3003			
	11759446	Feb 21, 2041	U-3003			
	11760740	Feb 12, 2035	DS DP			
	11786502	Oct 10, 2034	U-3007			
	11833130	Jan 25, 2033	U-3003			
	11850229	Jan 25, 2033	U-3342			
	9539234	Jan 25, 2033	U-3004			
	9730910	May 17, 2034	U-3005			
<u>TAZAROTENE - FABIOR</u>						
N 202428 001	10568859	Feb 24, 2030	DP U-2760			
	10688071	Feb 24, 2030	DP U-2760			
	8808716	Feb 24, 2030	DP			
<u>TAZAROTENE - ARAZLO</u>						
N 211882 001	11311482	May 11, 2038	U-2368			
	11679116	Jun 06, 2036	DP			
<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723 001	10155002	Apr 13, 2032	U-2736		NCE	Jan 23, 2025
	10245269	Apr 11, 2033	U-2737		ODE-299	Jan 23, 2027
	10245269	Apr 11, 2033	U-2851		ODE-314	Jun 18, 2027
	10245269	Apr 11, 2033	U-2854			
	10369155	Oct 16, 2035	U-2736			
	10369155	Oct 16, 2035	U-2852			
	10369155	Oct 16, 2035	U-2853			
	10420775	Apr 13, 2032	U-2736			
	10420775	Apr 13, 2032	U-2852			
	10420775	Apr 13, 2032	U-2853			
	10786511	Dec 19, 2035	DP			
	10821113	Apr 11, 2033	DS DP			
	11052093	Apr 13, 2032	DS DP U-2736			
	11052093	Apr 13, 2032	DS DP U-3179			
	11491163	Apr 11, 2033	U-3491			
	11491163	Apr 11, 2033	U-3492			
	8410088	Jan 23, 2034	DS DP			
	8691507	Sep 12, 2031	U-2852			
	8765732	Apr 13, 2032	U-2852			
	8765732	Apr 13, 2032	U-2853			
	8895245	Sep 12, 2031	U-2852			
	9090562	Apr 13, 2032	DS DP			
	9175331	Sep 12, 2031	U-2852			
	9333217	Sep 12, 2031	U-2852			
	9334527	Sep 12, 2031	U-2852			
	9394283	Apr 11, 2033	DS DP U-2852			
	9394283	Apr 11, 2033	DS DP U-2853			
	9522152	Apr 13, 2032	U-2738			

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<u>TAZEMETOSTAT HYDROBROMIDE - TAZVERIK</u>						
N 211723	001 9549931	Apr 13, 2032	U-2736			
	9549931	Apr 13, 2032	U-2852			
	9549931	Apr 13, 2032	U-2853			
	9688665	Aug 22, 2034	U-2736			
	9855275	Apr 13, 2032	U-2736			
	9872862	Apr 11, 2033	U-2738			
	9889138	Oct 16, 2035	U-2736			
	9889138	Oct 16, 2035	U-2852			
	9889138	Oct 16, 2035	U-2853			
	9949999	Sep 12, 2031	U-2851			
<u>TECHNETIUM TC-99M LABELED CARBON - TECHNEGAS KIT</u>						
N 022335	001				NP	Sep 29, 2026
<u>TECHNETIUM TC-99M SULFUR COLLOID KIT - TECHNETIUM TC-99M SULFUR COLLOID KIT</u>						
A 213516	001				CGT	Jun 01, 2024
<u>TECHNETIUM TC-99M TETROFOSMIN KIT - MYOVUE 30ML</u>						
N 020372	002 9549999	Mar 10, 2030	DP			
<u>TECHNETIUM TC-99M TILMANOCEPT - LYMPHOSEEK KIT</u>						
N 202207	001 6409990	May 12, 2025	DS		NPP	May 19, 2024
	9439985	Jan 30, 2029	DP			
<u>TECOVIRIMAT - TPOXX</u>						
N 208627	001 7737168	Sep 04, 2031	U-2346		ODE-200	Jul 13, 2025
	8039504	Jul 23, 2027	DP			
	8124643	Jun 18, 2024	DS DP			
	9339466	Mar 23, 2031	DS DP			
<u>TECOVIRIMAT - TPOXX</u>						
N 214518	001 10576165	Aug 02, 2031	DP			
	7737168	Sep 04, 2031	U-3377			
	8039504	Jul 23, 2027	DP			
	8124643	Jun 18, 2024	DS DP			
	8530509	Jun 18, 2024	DP			
	8802714	Jun 18, 2024	U-3377			
	9233097	Aug 02, 2031	DP			
	9907859	Aug 02, 2031	U-3377			
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205435	001 10065947	Feb 03, 2030	DP		NCE	Jun 20, 2019
	10442829	Feb 03, 2030	DS		GAIN	Jun 20, 2024
	7816379	Jun 20, 2028	DS DP U-2507			
	8420676	Feb 23, 2028	DS DP U-282			
	8426389	Dec 31, 2030	DS DP U-282			
	9624250	Feb 03, 2030	DS DP U-2507			
	9988406	Feb 03, 2030	DP			
<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205436	001 10065947	Feb 03, 2030	DP		NCE	Jun 20, 2019
	10442829	Feb 03, 2030	DS		GAIN	Jun 20, 2024
	7816379	Jun 20, 2028	DS DP U-2507			
	8420676	Feb 23, 2028	DS DP U-282			

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<u>TEDIZOLID PHOSPHATE - SIVEXTRO</u>						
N 205436	001	8426389	Dec 31, 2030	DS DP U-282		
		9624250	Feb 03, 2030	DS DP U-2507		
		9988406	Feb 03, 2030	DP		
<u>TEDUGLUTIDE RECOMBINANT - GATTEX KIT</u>						
N 203441	001	7847061	Nov 01, 2025	U-1320	Y	ODE-240 May 16, 2026
		7847061*PED	May 01, 2026			
		9060992	Nov 01, 2025	U-1320	Y	
		9060992*PED	May 01, 2026			
<u>TELAPREVIR - INCIVEK</u>						
N 201917	001	7820671	Feb 25, 2025	DS DP		
		8431615	May 30, 2028	U-1398		
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110	001	7531623	Jan 01, 2027	DS		
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>						
N 022110	002	7531623	Jan 01, 2027	DS		
<u>TELOTTRISTAT ETIPRATE - XERMELO</u>						
N 208794	001	7553840	Dec 11, 2027	DS		ODE-132 Feb 28, 2024
		7709493	Feb 28, 2031	DS U-1979		
		7968559	Dec 11, 2027	U-1979		
		8193204	Feb 27, 2031	DS		
		8653094	Dec 19, 2028	U-1979		
<u>TEMSIROLIMUS - TORISEL</u>						
N 022088	001	8026276	Jan 20, 2026	DP		
		8455539*PED	Jan 25, 2024			
		8722700*PED	Jan 25, 2024			
		8791097	May 10, 2032	U-1550		
		8791097	May 10, 2032	U-1551		
		8791097*PED	Nov 10, 2032			
<u>TENAPANOR HYDROCHLORIDE - IBSRELA</u>						
N 211801	001	8541448	Aug 01, 2033	DS DP		NCE Sep 12, 2024
		8969377	Dec 30, 2029	DS DP		
		9006281	May 02, 2030	U-2626		
		9408840	Dec 30, 2029	U-2626		
<u>TENAPANOR HYDROCHLORIDE - XPHOZAH</u>						
N 213931	001	10272079	Apr 10, 2034	U-3736		NCE Sep 12, 2024
		10272079	Apr 10, 2034	U-381		NP Oct 17, 2026
		10940146	Apr 10, 2034	U-3736		
		10940146	Apr 10, 2034	U-381		
		8541448	Aug 01, 2033	DS DP		
		8969377	Dec 30, 2029	DS DP		
<u>TENAPANOR HYDROCHLORIDE - XPHOZAH</u>						
N 213931	002	10272079	Apr 10, 2034	U-3736		NCE Sep 12, 2024
		10272079	Apr 10, 2034	U-381		NP Oct 17, 2026
		10940146	Apr 10, 2034	U-3736		
		10940146	Apr 10, 2034	U-381		
		8541448	Aug 01, 2033	DS DP		

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<u>TENAPANOR HYDROCHLORIDE - XPHOZAH</u>						
N 213931	002 8969377	Dec 30, 2029	DS DP			
<u>TENAPANOR HYDROCHLORIDE - XPHOZAH</u>						
N 213931	003 10272079	Apr 10, 2034		U-3736	NCE	Sep 12, 2024
	10272079	Apr 10, 2034		U-381	NP	Oct 17, 2026
	10940146	Apr 10, 2034		U-3736		
	10940146	Apr 10, 2034		U-381		
	8541448	Aug 01, 2033	DS DP			
	8969377	Dec 30, 2029	DS DP			
<u>TENOFOVIR ALAFENAMIDE FUMARATE - VEMLIDY</u>						
N 208464	001 7390791	Apr 17, 2025	DS DP		NPP	Oct 17, 2025
	7390791*PED	Oct 17, 2025				
	8754065	Aug 15, 2032	DS DP	U-1275		
	8754065	Aug 15, 2032	DS DP	U-999		
	8754065*PED	Feb 15, 2033				
	9296769	Aug 15, 2032	DS DP	U-1275		
	9296769	Aug 15, 2032	DS DP	U-999		
	9296769*PED	Feb 15, 2033				
<u>TEPOTINIB HYDROCHLORIDE - TEPMETKO</u>						
N 214096	001 8329692	Oct 30, 2029	DS DP		NCE	Feb 03, 2026
	8580781	Mar 19, 2030	DS DP		ODE-325	Feb 03, 2028
	8658643	Jul 04, 2028		U-3077		
	8921357	May 30, 2028	DS DP			
	8927540	Jul 21, 2028		U-3078		
	9062029	Jul 04, 2028	DP			
	9284300	Apr 29, 2028	DP			
	9403799	Jul 04, 2028		U-3077		
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992	001 6794410	Sep 12, 2026		U-1285	M-61	Apr 30, 2024
	6794410*PED	Mar 12, 2027			PED	Oct 30, 2024
	8802735	Sep 14, 2030	DP			
	8802735*PED	Mar 14, 2031				
	9186346	Feb 04, 2034		U-1786		
	9186346*PED	Aug 04, 2034				
<u>TERIFLUNOMIDE - AUBAGIO</u>						
N 202992	002 6794410	Sep 12, 2026		U-1285	M-61	Apr 30, 2024
	6794410*PED	Mar 12, 2027			PED	Oct 30, 2024
	8802735	Sep 14, 2030	DP			
	8802735*PED	Mar 14, 2031				
	9186346	Feb 04, 2034		U-1786		
	9186346*PED	Aug 04, 2034				
<u>TERIPARATIDE - FORTEO</u>						
N 021318	001 7517334	Mar 25, 2025	DP			
<u>TERIPARATIDE - FORTEO</u>						
N 021318	002 7517334	Mar 25, 2025	DP			
<u>TERLIPRESSIN ACETATE - TERLIVAZ</u>						
N 022231	001 10335452	Apr 05, 2037		U-3711	NCE	Sep 14, 2027

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<u>TERLIPRESSIN ACETATE - TERLIVAZ</u>						
N 022231	001				ODE-406	Sep 14, 2029
<u>TESTOSTERONE - TESTIM</u>						
N 021454	001	7320968	Jan 18, 2025	U-843		
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	001	8466136	Oct 12, 2026	DP		
		8466137	Oct 12, 2026	U-1103		
		8466138	Oct 12, 2026	U-1103		
		8486925	Oct 12, 2026	DP		
		8729057	Oct 12, 2026	DP		
		8741881	Oct 12, 2026	U-1103		
		8754070	Oct 12, 2026	DP		
		8759329	Oct 12, 2026	DP		
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	002	8466136	Oct 12, 2026	DP		
		8466137	Oct 12, 2026	U-1103		
		8466138	Oct 12, 2026	U-1103		
		8486925	Oct 12, 2026	DP		
		8729057	Oct 12, 2026	DP		
		8741881	Oct 12, 2026	U-1103		
		8754070	Oct 12, 2026	DP		
		8759329	Oct 12, 2026	DP		
<u>TESTOSTERONE - ANDROGEL</u>						
N 022309	003	8466136	Oct 12, 2026	DP		
		8466137	Oct 12, 2026	U-1103		
		8466138	Oct 12, 2026	U-1103		
		8486925	Oct 12, 2026	DP		
		8729057	Oct 12, 2026	DP		
		8741881	Oct 12, 2026	U-1103		
		8754070	Oct 12, 2026	DP		
		8759329	Oct 12, 2026	DP		
<u>TESTOSTERONE - AXIRON</u>						
N 022504	001	8419307	Feb 26, 2027	U-1386		
		8435944	Sep 27, 2027	U-1390		
		8807861	Feb 26, 2027	DP U-1563		
		8993520	Jun 02, 2026	U-1390		
		9180194	Jun 02, 2026	U-1390		
		9289586	Feb 26, 2027	U-1390		
<u>TESTOSTERONE - VOGELXO</u>						
N 204399	002	8785426	Feb 11, 2034	DP U-1531		
		9295675	Feb 11, 2034	DP U-1531		
		9662340	Feb 11, 2034	DP U-1531		
<u>TESTOSTERONE - VOGELXO</u>						
N 204399	003	8785426	Feb 11, 2034	DP U-1531		
		9295675	Feb 11, 2034	DP U-1531		
		9662340	Feb 11, 2034	DP U-1531		

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<u>TESTOSTERONE - NATESTO</u>						
N 205488	001	11090312	Mar 17, 2034		U-1616	
		8574622	Feb 04, 2024	DP		
		8784869	Feb 04, 2024	DP		
		8784882	Feb 04, 2024	DP	U-1557	
		8877230	Feb 04, 2024		U-1616	
<u>TESTOSTERONE CYPIONATE - TESTOSTERONE CYPIONATE</u>						
N 216318	001	11311554	Mar 25, 2039	DP		
		11642355	Mar 25, 2039	DP		
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	001	10238662	Feb 19, 2035	DP	U-2418	
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP	U-2418	
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		
		10912782	Feb 19, 2035	DP	U-2418	
		11160751	Oct 07, 2034	DP	U-2418	
		11191908	Oct 18, 2035	DP		
		11446440	Aug 21, 2031	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11771646	Apr 10, 2034	DP		
		11813435	Feb 25, 2035	DP		
		8021335	Oct 04, 2026	DP		
		8562564	Jan 24, 2026	DP		
		9180259	Jan 24, 2026	DP		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9744302	Nov 19, 2035	DP		
		9950125	Sep 04, 2036	DP	U-2418	
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	002	10238662	Feb 19, 2035	DP	U-2418	
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP	U-2418	
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		
		10912782	Feb 19, 2035	DP	U-2418	
		11160751	Oct 07, 2034	DP	U-2418	
		11191908	Oct 18, 2035	DP		
		11446440	Aug 21, 2031	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11771646	Apr 10, 2034	DP		

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<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	002	11813435	Feb 25, 2035	DP		
		8021335	Oct 04, 2026	DP		
		8562564	Jan 24, 2026	DP		
		9180259	Jan 24, 2026	DP		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9744302	Nov 19, 2035	DP		
		9950125	Sep 04, 2036	DP	U-2418	
<u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u>						
N 209863	003	10238662	Feb 19, 2035	DP	U-2418	
		10279131	Jul 31, 2031	DP		
		10357609	Aug 21, 2031	DP		
		10478560	Jan 24, 2026	DP		
		10646495	Aug 30, 2038	DP		
		10821072	Jun 04, 2033	DP	U-2418	
		10881798	Feb 11, 2034	DP		
		10905827	Aug 21, 2031	DP		
		10912782	Feb 19, 2035	DP	U-2418	
		11160751	Oct 07, 2034	DP	U-2418	
		11191908	Oct 18, 2035	DP		
		11446440	Aug 21, 2031	DP		
		11446441	Jan 24, 2026	DP		
		11497753	Mar 19, 2030	DP		
		11771646	Apr 10, 2034	DP		
		11813435	Feb 25, 2035	DP		
		8021335	Oct 04, 2026	DP		
		8562564	Jan 24, 2026	DP		
		9180259	Jan 24, 2026	DP		
		9533102	Jan 24, 2026	DP		
		9629959	Jan 24, 2026	DP		
		9744302	Nov 19, 2035	DP		
		9950125	Sep 04, 2036	DP	U-2418	
<u>TESTOSTERONE UNDECANOATE - AVEED</u>						
N 022219	001	7718640	Mar 14, 2027	DP		
		8338395	May 08, 2027		U-1500	
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089	001	10543219	Apr 12, 2030		U-2506	
		10617696	Apr 12, 2030	DS DP		
		11179402	Apr 14, 2026	DS DP		
		11179403	Apr 12, 2030		U-2506	
		11331325	Jan 06, 2027		U-2506	
		11426416	Apr 12, 2030		U-3420	
		11564933	Apr 12, 2039		U-1103	
		8241664	Mar 29, 2029	DP	U-2506	
		8492369	Dec 20, 2030	DP	U-2506	
		8778916	Apr 12, 2030	DP		

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<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 002	10543219	Apr 12, 2030	U-2506			
	10617696	Apr 12, 2030	DS DP			
	11179402	Apr 14, 2026	DS DP			
	11179403	Apr 12, 2030	U-2506			
	11331325	Jan 06, 2027	U-2506			
	11426416	Apr 12, 2030	U-3420			
	11564933	Apr 12, 2039	U-1103			
	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>TESTOSTERONE UNDECANOATE - JATENZO</u>						
N 206089 003	10543219	Apr 12, 2030	U-2506			
	10617696	Apr 12, 2030	DS DP			
	11179402	Apr 14, 2026	DS DP			
	11179403	Apr 12, 2030	U-2506			
	11331325	Jan 06, 2027	U-2506			
	11426416	Apr 12, 2030	U-3420			
	11564933	Apr 12, 2039	U-1103			
	8241664	Mar 29, 2029	DP U-2506			
	8492369	Dec 20, 2030	DP U-2506			
	8778916	Apr 12, 2030	DP			
<u>TESTOSTERONE UNDECANOATE - TLANDO</u>						
N 208088 001	10226473	Nov 30, 2030	U-1500		NP	Mar 28, 2025
	10716794	Nov 30, 2030	U-1500			
	11304960	Jan 08, 2029	DP U-1500			
	11311555	Nov 30, 2030	U-1500			
	11364249	Nov 30, 2030	U-1500			
	11433083	Nov 30, 2030	U-1500			
	11464735	Apr 28, 2041	DP U-1500			
	11559530	Nov 28, 2037	U-1500			
	8778922	Jan 08, 2029	DP U-1500			
	8865695	Jan 08, 2029	U-1500			
	9943527	Nov 30, 2030	U-1500			
	9949985	Nov 30, 2030	U-1500			
<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953 001	10576089	Dec 31, 2030	DP U-2506		NP	Jul 27, 2025
	10576090	Dec 31, 2030	DP			
	11590146	Dec 31, 2030	DP			
	11617758	Mar 15, 2033	DP U-2506			
<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953 002	10576089	Dec 31, 2030	DP U-2506		NP	Jul 27, 2025
	10576090	Dec 31, 2030	DP			
	11590146	Dec 31, 2030	DP			
	11617758	Mar 15, 2033	DP U-2506			
<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953 003	10576089	Dec 31, 2030	DP U-2506		NP	Jul 27, 2025
	10576090	Dec 31, 2030	DP			
	11590146	Dec 31, 2030	DP			

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<u>TESTOSTERONE UNDECANOATE - KYZATREX</u>						
N 213953	003 11617758	Mar 15, 2033	DP U-2506			
<u>THIOTEPA - TEPADINA</u>						
N 208264	001				ODE-129	Jan 26, 2024
<u>THIOTEPA - TEPADINA</u>						
N 208264	002				ODE-129	Jan 26, 2024
<u>TICAGRELOR - BRILINTA</u>						
N 022433	001 10300065	Jan 27, 2036	U-2541		I-848	Nov 05, 2023
	10300065	Jan 27, 2036	U-2542		M-283	May 09, 2025
	10300065*PED	Jul 27, 2036			PED	May 05, 2024
	8425934	Apr 17, 2030	DP		PED	Nov 09, 2025
	8425934*PED	Oct 17, 2030				
	RE46276	Oct 30, 2024	DS DP U-1935			
	RE46276	Oct 30, 2024	DS DP U-1936			
	RE46276	Oct 30, 2024	DS DP U-1937			
	RE46276	Oct 30, 2024	DS DP U-1938			
	RE46276	Oct 30, 2024	DS DP U-2838			
	RE46276	Oct 30, 2024	DS DP U-2839			
	RE46276	Oct 30, 2024	DS DP U-2988			
	RE46276*PED	Apr 30, 2025				
<u>TICAGRELOR - BRILINTA</u>						
N 022433	002 10300065	Jan 27, 2036	U-2541		I-848	Nov 05, 2023
	10300065	Jan 27, 2036	U-2542		M-283	May 09, 2025
	10300065*PED	Jul 27, 2036			PED	May 05, 2024
	8425934	Apr 17, 2030	DP		PED	Nov 09, 2025
	8425934*PED	Oct 17, 2030				
	RE46276	Oct 30, 2024	DS DP U-1935			
	RE46276	Oct 30, 2024	DS DP U-1936			
	RE46276	Oct 30, 2024	DS DP U-1937			
	RE46276	Oct 30, 2024	DS DP U-1938			
	RE46276	Oct 30, 2024	DS DP U-2838			
	RE46276	Oct 30, 2024	DS DP U-2839			
	RE46276	Oct 30, 2024	DS DP U-2988			
	RE46276*PED	Apr 30, 2025				
<u>TIGECYCLINE - TYGACIL</u>						
N 021821	001 10588975	Mar 13, 2026	DP			
	7879828	Feb 05, 2029	DP			
	8372995	Oct 08, 2030	DP			
	8975242	Oct 24, 2028	DP			
	9254328	Mar 13, 2026	DP			
	9694078	Mar 13, 2026	DP			
<u>TIGECYCLINE - TIGECYCLINE</u>						
N 211158	001 9855335	Apr 07, 2033	DP			
<u>TIOPRONIN - THIOLA</u>						
N 019569	001				ODE-267	Jun 28, 2026
<u>TIOPRONIN - THIOLA EC</u>						
N 211843	001 11458104	Nov 14, 2038	U-3441		ODE*	Jun 28, 2026

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<u>TIOPRONIN - THIOLA EC</u>						
N 211843	002 11458104	Nov 14, 2038	U-3441		ODE*	Jun 28, 2026
<u>TIOTROPIUM BROMIDE - SPIRIVA</u>						
N 021395	001 7694676	Mar 12, 2027	DP			
	7694676*PED	Sep 12, 2027				
	8022082	Jan 19, 2026	DP U-1186			
	8022082*PED	Jul 19, 2026				
	9010323	Apr 19, 2030	DP			
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	001 7284474	Aug 26, 2024	DP			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7896264*PED	Nov 26, 2025				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Oct 01, 2027				
<u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u>						
N 021936	002 7284474	Aug 26, 2024	DP			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7896264*PED	Nov 26, 2025				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Oct 01, 2027				
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981	001 10456399	Feb 03, 2037	U-3657		ODE-229	Feb 22, 2026
	10456399	Feb 03, 2037	U-3658			
	10457666	Jun 17, 2034	DS DP			
	10960004	Feb 03, 2037	U-3657			
	10960004	Feb 03, 2037	U-3658			
	9527833	Jun 17, 2034	DS DP			
	9943537	Sep 05, 2034	U-3659			
	RE46284	Sep 22, 2029	U-1751			
	RE46284	Sep 22, 2029	U-2503			
	RE46284	Sep 22, 2029	U-3656			
<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981	002 10456399	Feb 03, 2037	U-3657		ODE-229	Feb 22, 2026
	10456399	Feb 03, 2037	U-3658			
	10457666	Jun 17, 2034	DS DP			

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<u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u>						
N 207981	002	10960004	Feb 03, 2037	U-3657		
		10960004	Feb 03, 2037	U-3658		
		9527833	Jun 17, 2034	DS DP		
		9943537	Sep 05, 2034	U-3659		
		RE46284	Sep 22, 2029	U-1751		
		RE46284	Sep 22, 2029	U-2503		
		RE46284	Sep 22, 2029	U-3656		
<u>TIRBANIBULIN - KLISYRI</u>						
N 213189	001	10323001	Dec 28, 2027	DP	NCE	Dec 14, 2025
		10617693	Mar 12, 2038	U-3015		
		10669236	Sep 07, 2038	DS DP		
		11497750	Mar 12, 2038	U-3015		
		7300931	Feb 06, 2026	DS DP		
		7851470	Feb 02, 2029	DS DP U-3015		
		8236799	Dec 28, 2025	DS DP		
		8980890	Dec 28, 2025	DS DP		
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866	001	11357820	Jun 14, 2039	DP	NCE	May 13, 2027
		8734394	Feb 24, 2031	DP		
		9402957	Jun 29, 2031	DP		
		9474780	Jan 05, 2036	DS DP U-3378		
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866	002	11357820	Jun 14, 2039	DP	NCE	May 13, 2027
		8734394	Feb 24, 2031	DP		
		9402957	Jun 29, 2031	DP		
		9474780	Jan 05, 2036	DS DP U-3378		
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866	003	11357820	Jun 14, 2039	DP	NCE	May 13, 2027
		8734394	Feb 24, 2031	DP		
		9402957	Jun 29, 2031	DP		
		9474780	Jan 05, 2036	DS DP U-3378		
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866	004	11357820	Jun 14, 2039	DP	NCE	May 13, 2027
		8734394	Feb 24, 2031	DP		
		9402957	Jun 29, 2031	DP		
		9474780	Jan 05, 2036	DS DP U-3378		
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866	005	11357820	Jun 14, 2039	DP	NCE	May 13, 2027
		8734394	Feb 24, 2031	DP		
		9402957	Jun 29, 2031	DP		
		9474780	Jan 05, 2036	DS DP U-3378		
<u>TIRZEPATIDE - MOUNJARO (AUTOINJECTOR)</u>						
N 215866	006	11357820	Jun 14, 2039	DP	NCE	May 13, 2027
		8734394	Feb 24, 2031	DP		
		9402957	Jun 29, 2031	DP		
		9474780	Jan 05, 2036	DS DP U-3378		

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<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 001	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	8734394	Feb 24, 2031	DP		NP	Nov 08, 2026
	9402957	Jun 29, 2031	DP			
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 002	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	8734394	Feb 24, 2031	DP		NP	Nov 08, 2026
	9402957	Jun 29, 2031	DP			
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 003	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	8734394	Feb 24, 2031	DP		NP	Nov 08, 2026
	9402957	Jun 29, 2031	DP			
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 004	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	8734394	Feb 24, 2031	DP		NP	Nov 08, 2026
	9402957	Jun 29, 2031	DP			
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 005	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	8734394	Feb 24, 2031	DP		NP	Nov 08, 2026
	9402957	Jun 29, 2031	DP			
	9474780	Jan 05, 2036	DS DP			
<u>TIRZEPATIDE - ZEPBOUND</u>						
N 217806 006	11357820	Jun 14, 2039	DP		NCE	May 13, 2027
	8734394	Feb 24, 2031	DP		NP	Nov 08, 2026
	9402957	Jun 29, 2031	DP			
	9474780	Jan 05, 2036	DS DP			
<u>TIVOZANIB HYDROCHLORIDE - FOTIVDA</u>						
N 212904 001	11504365	Nov 05, 2039		U-3476	NCE	Mar 10, 2026
	6821987	Apr 26, 2024	DS DP	U-3100		
<u>TIVOZANIB HYDROCHLORIDE - FOTIVDA</u>						
N 212904 002	11504365	Nov 05, 2039		U-3476	NCE	Mar 10, 2026
	6821987	Apr 26, 2024	DS DP	U-3100		
<u>TOBRAMYCIN - TOBI PODHALER</u>						
N 201688 001	10207066	Nov 04, 2030	DP	U-909		
	11484671	Nov 07, 2024	DP	U-909		
	7516741	Jan 11, 2024	DP			
	7559325	Oct 27, 2025	DP			
	8069851	Sep 24, 2024	DP			
	8664187	Jun 20, 2025		U-909		
	8869794	Sep 12, 2028	DP	U-909		
	RE47526	Apr 09, 2024	DP			

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<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 203214	001 RE41783	Dec 08, 2025	DS		I-879	Dec 14, 2024
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 203214	002 RE41783	Dec 08, 2025	DS		I-879	Dec 14, 2024
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246	001 11253523	Mar 14, 2034		U-3326	I-879	Dec 14, 2024
	11253523	Mar 14, 2034		U-3327		
	11253523	Mar 14, 2034		U-3328		
	11253523	Mar 14, 2034		U-3329		
	9937181	Mar 14, 2034		DP		
	RE41783	Dec 08, 2025	DS			
<u>TOFACITINIB CITRATE - XELJANZ XR</u>						
N 208246	002 10639309	Mar 14, 2034		DP		
	11253523	Mar 14, 2034		U-3326		
	11253523	Mar 14, 2034		U-3327		
	11253523	Mar 14, 2034		U-3328		
	11253523	Mar 14, 2034		U-3329		
	RE41783	Dec 08, 2025	DS			
<u>TOFACITINIB CITRATE - XELJANZ</u>						
N 213082	001 RE41783	Dec 08, 2025	DS			
<u>TOFERSEN - QALSODY</u>						
N 215887	001 10385341	Apr 01, 2035	DS DP		NCE	Apr 25, 2028
	10669546	Apr 01, 2035		U-3575	ODE-432	Apr 25, 2030
	10968453	Apr 01, 2035		U-3575		
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	001 10905694	Apr 07, 2030		DP		
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	002 10905694	Apr 07, 2030		DP		
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - SAMSCA</u>						
N 022275	003 10905694	Apr 07, 2030		DP		
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	001 10905694	Apr 07, 2030		DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	002 10905694	Apr 07, 2030		DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	003 10905694	Apr 07, 2030		DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOLVAPTAN - JYNARQUE</u>						
N 204441	004 10905694	Apr 07, 2030		DP	ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			

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<u>TOLVAPTAN - JYNARQUE</u>						
N 204441 005	10905694	Apr 07, 2030	DP		ODE-178	Apr 23, 2025
	8501730	Sep 01, 2026	DS			
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635 001	10314790	Nov 16, 2027	DP U-1675			
	10314790	Nov 16, 2027	DP U-1992			
	8298576	Apr 04, 2028	DP U-106			
	8298576	Apr 04, 2028	DP U-1992			
	8298580	Nov 16, 2027	DP U-106			
	8298580	Nov 16, 2027	DP U-1992			
	8663683	Nov 16, 2027	DP U-106			
	8663683	Nov 16, 2027	DP U-1992			
	8877248	Nov 16, 2027	DP U-106			
	8877248	Nov 16, 2027	DP U-1992			
	8889191	Nov 16, 2027	U-106			
	8889191	Nov 16, 2027	U-1992			
	8992989	Nov 16, 2027	DP U-1675			
	8992989	Nov 16, 2027	DP U-1992			
	9549940	Nov 16, 2027	DP U-1675			
	9549940	Nov 16, 2027	DP U-1992			
	9555004	Nov 16, 2027	DP U-1675			
	9555004	Nov 16, 2027	DP U-1992			
	9622983	Nov 16, 2027	DP U-1675			
	9622983	Nov 16, 2027	DP U-1992			
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635 002	10314790	Nov 16, 2027	DP U-1675			
	10314790	Nov 16, 2027	DP U-1992			
	8298576	Apr 04, 2028	DP U-106			
	8298576	Apr 04, 2028	DP U-1992			
	8298580	Nov 16, 2027	DP U-106			
	8298580	Nov 16, 2027	DP U-1992			
	8663683	Nov 16, 2027	DP U-106			
	8663683	Nov 16, 2027	DP U-1992			
	8877248	Nov 16, 2027	DP U-106			
	8877248	Nov 16, 2027	DP U-1992			
	8889191	Nov 16, 2027	U-106			
	8889191	Nov 16, 2027	U-1992			
	8992989	Nov 16, 2027	DP U-1675			
	8992989	Nov 16, 2027	DP U-1992			
	9549940	Nov 16, 2027	DP U-1675			
	9549940	Nov 16, 2027	DP U-1992			
	9555004	Nov 16, 2027	DP U-1675			
	9555004	Nov 16, 2027	DP U-1992			
	9622983	Nov 16, 2027	DP U-1675			
	9622983	Nov 16, 2027	DP U-1992			
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635 003	10314790	Nov 16, 2027	DP U-1675			
	10314790	Nov 16, 2027	DP U-1992			
	8298576	Apr 04, 2028	DP U-106			

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<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	003	8298576	Apr 04, 2028	DP U-1992		
		8298580	Nov 16, 2027	DP U-106		
		8298580	Nov 16, 2027	DP U-1992		
		8663683	Nov 16, 2027	DP U-106		
		8663683	Nov 16, 2027	DP U-1992		
		8877248	Nov 16, 2027	DP U-106		
		8877248	Nov 16, 2027	DP U-1992		
		8889191	Nov 16, 2027	U-106		
		8889191	Nov 16, 2027	U-1992		
		8992989	Nov 16, 2027	DP U-1675		
		8992989	Nov 16, 2027	DP U-1992		
		9549940	Nov 16, 2027	DP U-1675		
		9549940	Nov 16, 2027	DP U-1992		
		9555004	Nov 16, 2027	DP U-1675		
		9555004	Nov 16, 2027	DP U-1992		
		9622983	Nov 16, 2027	DP U-1675		
		9622983	Nov 16, 2027	DP U-1992		
<u>TOPIRAMATE - TROKENDI XR</u>						
N 201635	004	10314790	Nov 16, 2027	DP U-1675		
		10314790	Nov 16, 2027	DP U-1992		
		8298576	Apr 04, 2028	DP U-106		
		8298576	Apr 04, 2028	DP U-1992		
		8298580	Nov 16, 2027	DP U-106		
		8298580	Nov 16, 2027	DP U-1992		
		8663683	Nov 16, 2027	DP U-106		
		8663683	Nov 16, 2027	DP U-1992		
		8877248	Nov 16, 2027	DP U-106		
		8877248	Nov 16, 2027	DP U-1992		
		8889191	Nov 16, 2027	U-106		
		8889191	Nov 16, 2027	U-1992		
		8992989	Nov 16, 2027	DP U-1675		
		8992989	Nov 16, 2027	DP U-1992		
		9549940	Nov 16, 2027	DP U-1675		
		9549940	Nov 16, 2027	DP U-1992		
		9555004	Nov 16, 2027	DP U-1675		
		9555004	Nov 16, 2027	DP U-1992		
		9622983	Nov 16, 2027	DP U-1675		
		9622983	Nov 16, 2027	DP U-1992		
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	001	10363224	Mar 19, 2033	U-766		
		8652527	Mar 19, 2033	DP		
		8889190	Mar 19, 2033	DP		
		9101545	Mar 19, 2033	DP		
		9555005	Mar 19, 2033	DP		
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	002	10363224	Mar 19, 2033	U-766		
		8652527	Mar 19, 2033	DP		
		8889190	Mar 19, 2033	DP		

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<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	002	9101545	Mar 19, 2033	DP		
		9555005	Mar 19, 2033	DP		
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	003	10363224	Mar 19, 2033	U-766		
		8652527	Mar 19, 2033	DP		
		8889190	Mar 19, 2033	DP		
		9101545	Mar 19, 2033	DP		
		9555005	Mar 19, 2033	DP		
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	004	10363224	Mar 19, 2033	U-766		
		8652527	Mar 19, 2033	DP		
		8889190	Mar 19, 2033	DP		
		9101545	Mar 19, 2033	DP		
		9555005	Mar 19, 2033	DP		
<u>TOPIRAMATE - OUDEXY XR</u>						
N 205122	005	10363224	Mar 19, 2033	U-766		
		8652527	Mar 19, 2033	DP		
		8889190	Mar 19, 2033	DP		
		9101545	Mar 19, 2033	DP		
		9555005	Mar 19, 2033	DP		
<u>TOPIRAMATE - EPRONTIA</u>						
N 214679	001	11433046	Aug 21, 2040	U-3413		
		11433046	Aug 21, 2040	U-3414		
		11433046	Aug 21, 2040	U-3415		
		11633374	Aug 21, 2040	DP		
		11826343	Aug 21, 2040	DP		
<u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u>						
N 020981	001	8158645	Dec 10, 2024	DP		
<u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u>						
N 020981	002	8158645	Dec 10, 2024	DP		
<u>TORSEMIDE - SOAANZ</u>						
N 213218	001	10154963	Oct 06, 2033	DP		
<u>TORSEMIDE - SOAANZ</u>						
N 213218	002	10154963	Oct 06, 2033	DP		
<u>TRABECTEDIN - YONDELIS</u>						
N 207953	001	8895557	Jan 07, 2028	DP		
		8895557*PED	Jul 07, 2028			
<u>TRAMADOL HYDROCHLORIDE - ODOLO</u>						
N 214044	001	11103452	Sep 01, 2040	DP	U-3197	
		11752103	Sep 01, 2040	DP	U-3197	
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	001	10869869	Aug 30, 2033	U-3184	I-895	Jun 22, 2025
		10869869*PED	Mar 02, 2034		I-908	Mar 16, 2026
		7378423	May 29, 2027	DS DP	ODE-148	Jun 22, 2024
		7378423*PED	Nov 29, 2027		ODE-182	Apr 30, 2025

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<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114 001	8580304	Jan 28, 2032	DP		ODE-183	May 04, 2025
	8580304*PED	Jul 28, 2032			ODE-428	Mar 16, 2030
	8703781	Oct 15, 2030	DS DP U-1712		PED	Dec 22, 2024
	8703781	Oct 15, 2030	DS DP U-2020		PED	Oct 30, 2025
	8703781	Oct 15, 2030	DS DP U-2037		PED	Nov 04, 2025
	8703781	Oct 15, 2030	DS DP U-2302		PED	Dec 22, 2025
	8703781	Oct 15, 2030	DS DP U-2305			
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-1581			
	8835443	Jun 10, 2025	U-1582			
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	8835443	Jun 10, 2025	U-2302			
	8835443	Jun 10, 2025	U-2305			
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030	U-2020			
	8952018*PED	Apr 15, 2031				
	9155706	Jan 28, 2032	DP			
	9155706*PED	Jul 28, 2032				
	9271941	Jan 28, 2032	DP			
	9271941*PED	Jul 28, 2032				
	9399021	Jan 28, 2032	DP			
	9399021*PED	Jul 28, 2032				
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114 002	10869869	Aug 30, 2033	U-3184		I-895	Jun 22, 2025
	10869869*PED	Mar 02, 2034			I-908	Mar 16, 2026
	7378423	May 29, 2027	DS DP		ODE-148	Jun 22, 2024
	7378423*PED	Nov 29, 2027			ODE-182	Apr 30, 2025
	8580304	Jan 28, 2032	DP		ODE-183	May 04, 2025
	8580304*PED	Jul 28, 2032			ODE-428	Mar 16, 2030
	8703781	Oct 15, 2030	DS DP U-1712		PED	Dec 22, 2024
	8703781	Oct 15, 2030	DS DP U-2020		PED	Oct 30, 2025
	8703781	Oct 15, 2030	DS DP U-2037		PED	Nov 04, 2025
	8703781	Oct 15, 2030	DS DP U-2302		PED	Dec 22, 2025
	8703781	Oct 15, 2030	DS DP U-2305			
	8703781*PED	Apr 15, 2031				
	8835443	Jun 10, 2025	U-1581			
	8835443	Jun 10, 2025	U-1582			
	8835443	Jun 10, 2025	U-2020			
	8835443	Jun 10, 2025	U-2037			
	8835443	Jun 10, 2025	U-2302			
	8835443	Jun 10, 2025	U-2305			
	8835443*PED	Dec 10, 2025				
	8952018	Oct 15, 2030	U-2020			
	8952018*PED	Apr 15, 2031				
	9155706	Jan 28, 2032	DP			
	9155706*PED	Jul 28, 2032				
	9271941	Jan 28, 2032	DP			
	9271941*PED	Jul 28, 2032				

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<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	002	9399021	Jan 28, 2032	DP		
		9399021*PED	Jul 28, 2032			
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 204114	003	10869869	Aug 30, 2033	U-3184	I-895	Jun 22, 2025
		10869869*PED	Mar 02, 2034		I-908	Mar 16, 2026
		7378423	May 29, 2027	DS DP	ODE-148	Jun 22, 2024
		7378423*PED	Nov 29, 2027		ODE-182	Apr 30, 2025
		8580304	Jan 28, 2032	DP	ODE-183	May 04, 2025
		8580304*PED	Jul 28, 2032		ODE-428	Mar 16, 2030
		8703781	Oct 15, 2030	DS DP U-1712	PED	Dec 22, 2024
		8703781	Oct 15, 2030	DS DP U-2020	PED	Oct 30, 2025
		8703781	Oct 15, 2030	DS DP U-2037	PED	Nov 04, 2025
		8703781	Oct 15, 2030	DS DP U-2302	PED	Dec 22, 2025
		8703781	Oct 15, 2030	DS DP U-2305		
		8703781*PED	Apr 15, 2031			
		8835443	Jun 10, 2025	U-1581		
		8835443	Jun 10, 2025	U-1582		
		8835443	Jun 10, 2025	U-2020		
		8835443	Jun 10, 2025	U-2037		
		8835443	Jun 10, 2025	U-2302		
		8835443	Jun 10, 2025	U-2305		
		8835443*PED	Dec 10, 2025			
		8952018	Oct 15, 2030	U-2020		
		8952018*PED	Apr 15, 2031			
		9155706	Jan 28, 2032	DP		
		9155706*PED	Jul 28, 2032			
		9271941	Jan 28, 2032	DP		
		9271941*PED	Jul 28, 2032			
		9399021	Jan 28, 2032	DP		
		9399021*PED	Jul 28, 2032			
<u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u>						
N 217513	001	7378423	May 29, 2027	DS DP	NP	Mar 16, 2026
		8703781	Oct 15, 2030	DS DP U-3564	ODE-428	Mar 16, 2030
		8835443	Jun 10, 2025	U-3564		
<u>TRANEXAMIC ACID - TRANEXAMIC ACID</u>						
A 217155	001				CGT	Apr 16, 2024
<u>TRANEXAMIC ACID - LYSTEDA</u>						
N 022430	001	7947739	Mar 04, 2025	DP		
		8022106	Mar 04, 2025	U-1182		
		8273795	Mar 04, 2025	U-1182		
		8487005	Mar 04, 2025	DP U-1182		
		8791160	Mar 04, 2025	DP U-1182		
		8809394	Mar 04, 2025	DP U-1182		
		8957113	Mar 04, 2025	DP U-1182		
		9060939	Mar 04, 2025	DP		
<u>TRAVOPROST - TRAVATAN Z</u>						
N 021994	001	8268299	Oct 13, 2029	DP		
		8323630	Sep 20, 2027	DP		

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<u>TRAVOPROST - TRAVATAN Z</u>						
N 021994	001 8388941	Sep 20, 2027	DP			
<u>TRAVOPROST - IZBA</u>						
N 204822	001 8178582	Oct 10, 2029	DP			
	8722735	Oct 10, 2029	DP			
	8754123	May 19, 2029	DP			
	9144561	Mar 13, 2029	DP			
<u>TRAVOPROST - IDOSE TR</u>						
N 218010	001 10206813	Oct 17, 2030	DP			
	11426306	Oct 17, 2030	DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	001 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	002 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	003 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - DESYREL</u>						
N 018207	004 8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N 022411	001 7829120	Mar 27, 2027	DP U-796			
	8133893	Mar 13, 2029	DS DP			
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N 022411	002 7829120	Mar 27, 2027	DP U-796			
	8133893	Mar 13, 2029	DS DP			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	001 10076505	Dec 16, 2024	DP			
	10695308	Dec 16, 2024	U-2845			
	11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9199908	May 24, 2024	U-1771			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024	U-2036			
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	002 10076505	Dec 16, 2024	DP			
	10695308	Dec 16, 2024	U-2845			
	11723887	Dec 15, 2028	DS			
	7999007	Mar 29, 2029	DP U-1437			
	8653137	Sep 05, 2028	U-1437			
	8658694	Sep 05, 2028	U-1437			
	9199908	May 24, 2024	U-1771			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024	U-2036			

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<u>TREPROSTINIL - REMODULIN</u>						
N 021272	003	10076505	Dec 16, 2024	DP		
		10695308	Dec 16, 2024		U-2845	
		11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9199908	May 24, 2024		U-1771	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	004	10076505	Dec 16, 2024	DP		
		10695308	Dec 16, 2024		U-2845	
		11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9199908	May 24, 2024		U-1771	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	005	10076505	Dec 16, 2024	DP		
		10695308	Dec 16, 2024		U-2845	
		11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9199908	May 24, 2024		U-1771	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	006	11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9199908	May 24, 2024		U-1771	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	007	11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9199908	May 24, 2024		U-1771	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		

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<u>TREPROSTINIL - REMODULIN</u>						
N 021272	007	11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9199908	May 24, 2024		U-1771	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
<u>TREPROSTINIL - REMODULIN</u>						
N 021272	008	11723887	Dec 15, 2028	DS		
		7999007	Mar 29, 2029	DP	U-1437	
		8653137	Sep 05, 2028		U-1437	
		8658694	Sep 05, 2028		U-1437	
		9199908	May 24, 2024		U-1771	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
<u>TREPROSTINIL - TYVASO</u>						
N 022387	001	10376525	May 14, 2027		U-1849	I-856 Mar 31, 2024
		10716793	May 14, 2027		U-1849	
		11723887	Dec 15, 2028	DS		
		11826327	Jan 04, 2042		U-3749	
		9339507	Mar 10, 2028	DP		
		9358240	May 05, 2028		U-1849	
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	001	10076505	Dec 16, 2024	DP		
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	002	10076505	Dec 16, 2024	DP		
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	003	10076505	Dec 16, 2024	DP		
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - REMODULIN</u>						
N 208276	004	10076505	Dec 16, 2024	DP		
		9593066	Dec 15, 2028	DS		
		9604901	Dec 15, 2028	DS		
		9713599	Dec 16, 2024		U-2036	
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324	001	10130685	Aug 23, 2025	DP		
		10421729	Apr 01, 2035	DP		

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<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324 001	10716793	May 14, 2027	U-1849			
	10772883	Jun 11, 2030	DP			
	11723887	Dec 15, 2028	DS			
	11826327	Jan 04, 2042	U-3749			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324 002	10130685	Aug 23, 2025	DP			
	10421729	Apr 01, 2035	DP			
	10716793	May 14, 2027	U-1849			
	10772883	Jun 11, 2030	DP			
	11723887	Dec 15, 2028	DS			
	11826327	Jan 04, 2042	U-3749			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324 003	10130685	Aug 23, 2025	DP			
	10421729	Apr 01, 2035	DP			
	10716793	May 14, 2027	U-1849			
	10772883	Jun 11, 2030	DP			
	11723887	Dec 15, 2028	DS			
	11826327	Jan 04, 2042	U-3749			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL - TYVASO DPI</u>						
N 214324 004	10130685	Aug 23, 2025	DP			
	10421729	Apr 01, 2035	DP			
	10716793	May 14, 2027	U-1849			
	10772883	Jun 11, 2030	DP			
	11723887	Dec 15, 2028	DS			
	11826327	Jan 04, 2042	U-3749			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 001	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	7544713	Jul 14, 2024	U-1475			
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP U-2724			
	8747897	Aug 11, 2031	DP U-2725			
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024	U-1475			
	9393203	Apr 27, 2026	DP U-1877			
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			

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<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 001	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP	U-1877		
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 002	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS	Y		
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP	U-1877		
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 003	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS	Y		
	8747897	Aug 11, 2031	DP	U-2724		
	8747897	Aug 11, 2031	DP	U-2725		
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP	U-1877		
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 004	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026

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<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 004	7417070	Jul 30, 2026	DS			
	7544713	Jul 14, 2024			U-1475	
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS	Y		
	8747897	Aug 11, 2031	DP		U-2724	
	8747897	Aug 11, 2031	DP		U-2725	
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024			U-1475	
	9393203	Apr 27, 2026	DP		U-1877	
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TREPROSTINIL DIOLAMINE - ORENITRAM</u>						
N 203496 005	11723887	Dec 15, 2028	DS		ODE-272	Oct 18, 2026
	7417070	Jul 30, 2026	DS			
	7544713	Jul 14, 2024			U-1475	
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8747897	Aug 11, 2031	DP		U-2724	
	8747897	Aug 11, 2031	DP		U-2725	
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024			U-1475	
	9393203	Apr 27, 2026	DP		U-1877	
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<u>TRETINOIN - TRETINOIN MICROSPHERE</u>						
A 215609 001					CGT	Mar 06, 2024
<u>TRETINOIN - ALTRENO</u>						
N 209353 001	10653656	Aug 22, 2038	DP		U-2368	
	11324710	Aug 22, 2038			U-2368	
<u>TRIAMCINOLONE ACETONIDE - TRIESENC</u>						
N 022048 001	8128960	Dec 17, 2029	DP			
	8211880	Mar 10, 2029			U-1257	
	8211880	Mar 10, 2029			U-1258	
<u>TRIAMCINOLONE ACETONIDE - ZILRETTA</u>						
N 208845 001	8828440	Aug 04, 2031	DP			
	9555048	Aug 04, 2031			U-2151	
<u>TRIAMCINOLONE ACETONIDE - XIPERE</u>						
N 211950 001	8636713	May 02, 2027			U-3234	Oct 22, 2024
	9636332	Nov 08, 2033			U-3234	
	9937075	May 02, 2034	DP			

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<u>TRICLABENDAZOLE - EGATEN</u>						
N 208711	001				NCE	Feb 13, 2024
					ODE-228	Feb 13, 2026
<u>TRIENTINE TETRAHYDROCHLORIDE - CUVRIOR</u>						
N 215760	001	10988436	May 03, 2039	DS	NP	Apr 28, 2025
		11072577	May 03, 2039	U-3370	ODE-401	Apr 28, 2029
<u>TRIFAROTENE - AKLIEF</u>						
N 211527	001	7807708	Jul 19, 2031	DS DP	NCE	Oct 04, 2024
		8227507	Dec 21, 2025	U-818		
		8470871	Dec 21, 2025	U-2639		
		9084778	May 30, 2033	DP U-134		
		9498465	May 30, 2033	DP U-1033		
<u>TRIEPTANOIN - DOJOLVI</u>						
N 213687	001	8697748	Apr 28, 2029	DP	NCE	Jun 30, 2025
		9186344	Jul 01, 2025	DP	ODE-311	Jun 30, 2027
<u>TRILACICLIB DIHYDROCHLORIDE - COSELA</u>						
N 214200	001	10085992	Mar 14, 2034	U-3081	NCE	Feb 12, 2026
		10189849	Oct 25, 2031	DS		
		10189850	Oct 25, 2031	DP		
		10927120	Oct 25, 2031	DP		
		10966984	Mar 14, 2034	U-3079		
		10966984	Mar 14, 2034	U-3080		
		11040042	Mar 14, 2034	DP		
		11529352	Jul 23, 2039	U-3504		
		11717523	Mar 14, 2034	U-3678		
		11717523	Mar 14, 2034	U-3679		
		8598186	Oct 25, 2031	DS DP		
		8598197	Oct 25, 2031	DS DP		
		9487530	Mar 14, 2034	U-3079		
		9487530	Mar 14, 2034	U-3080		
		9957276	Oct 25, 2031	DS		
<u>TRIPTORELIN PAMOATE - TRELSTAR</u>						
N 022437	001	10166181	Jun 30, 2029	DP		
<u>TRIPTORELIN PAMOATE - TRIPTODUR KIT</u>						
N 208956	001	10166181	Jun 30, 2029	DP	ODE-149	Jun 29, 2024
<u>TROFINETIDE - DAYBUE</u>						
N 217026	001	11370755	Aug 03, 2040	DS DP	NCE	Mar 10, 2028
		9212204	Jan 27, 2032	U-3556	ODE-425	Mar 10, 2030
<u>TUCATINIB - TUKYSA</u>						
N 213411	001	11207324	Apr 27, 2038	U-3510	I-906	Jan 19, 2026
		11504370	Mar 25, 2033	U-2788	NCE	Apr 17, 2025
		7452895	Nov 16, 2024	DS DP U-2788	ODE-309	Apr 17, 2027
		8648087	Apr 12, 2031	DS DP	ODE-422	Jan 19, 2030
		9457093	Oct 12, 2032	DP U-2788		
		9693989	May 09, 2027	DP U-2788		

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<u>TUCATINIB - TUKYSA</u>						
N 213411 002	11207324	Apr 27, 2038	U-3510		I-906	Jan 19, 2026
	11504370	Mar 25, 2033	U-2788		NCE	Apr 17, 2025
	7452895	Nov 16, 2024	DS DP U-2788		ODE-309	Apr 17, 2027
	8648087	Apr 12, 2031	DS DP		ODE-422	Jan 19, 2030
	9457093	Oct 12, 2032	DP U-2788			
	9693989	May 09, 2027	DP U-2788			
<u>UBROGEPANT - UBRELVY</u>						
N 211765 001	10117836	Jan 30, 2035	DP		NCE	Dec 23, 2024
	11717515	Dec 22, 2041	U-3677			
	8754096	Jul 19, 2032	DS DP U-2717			
	8912210	Dec 23, 2033	DS DP			
	9499545	Nov 10, 2031	DS DP U-2718			
	9833448	Nov 10, 2031	U-2718			
<u>UBROGEPANT - UBRELVY</u>						
N 211765 002	10117836	Jan 30, 2035	DP		NCE	Dec 23, 2024
	8754096	Jul 19, 2032	DS DP U-2717			
	8912210	Dec 23, 2033	DS DP			
	9499545	Nov 10, 2031	DS DP U-2718			
	9833448	Nov 10, 2031	U-2718			
<u>ULIPRISTAL ACETATE - ELLA</u>						
N 022474 001	10159681	Apr 13, 2030	U-2510		M-271	Jun 24, 2024
	10772897	Apr 13, 2030	U-2958			
	8426392	Jun 12, 2030	U-1389			
	8512745	Jun 02, 2030	DP			
	8735380	Feb 20, 2029	DP			
	8962603	Jun 12, 2030	U-1657			
	9283233	Apr 13, 2030	U-1821			
	9844510	Dec 08, 2028	DP			
<u>UMBRALISIB TOSYLATE - UKONIO</u>						
N 213176 001	10072013	Jul 02, 2033	U-3063		NCE	Feb 05, 2026
	10072013	Jul 02, 2033	U-3064			
	10414773	May 26, 2035	DS DP U-3063			
	10414773	May 26, 2035	DS DP U-3064			
	10570142	Jul 02, 2033	DS DP U-3063			
	10570142	Jul 02, 2033	DS DP U-3064			
	10947244	May 26, 2035	U-3063			
	10947244	May 26, 2035	U-3064			
	10981919	Jul 02, 2033	U-3063			
	10981919	Jul 02, 2033	U-3064			
	9150579	Jul 02, 2033	DS DP			
	9669033	Jul 02, 2033	U-3063			
	9669033	Jul 02, 2033	U-3064			
	9969740	May 26, 2035	DS DP U-3063			
	9969740	May 26, 2035	DS DP U-3064			
<u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u>						
N 205382 001	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	8113199	Oct 23, 2027	DP			

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<u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u>						
N 205382 001	8161968	Feb 05, 2028	DP			
	8183257	Jul 27, 2025	U-1476			
	8201556	Feb 05, 2029	DP			
	8309572	Apr 27, 2025	U-1476			
	8534281	Mar 08, 2030	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
<u>UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - ANORO ELLIPTA</u>						
N 203975 001	11090294	Nov 29, 2030	U-3203			
	7439393	May 21, 2025	DS DP U-1476			
	7439393*PED	Nov 21, 2025				
	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	8113199	Oct 23, 2027	DP			
	8113199*PED	Apr 23, 2028				
	8161968	Feb 05, 2028	DP			
	8161968*PED	Aug 05, 2028				
	8183257	Jul 27, 2025	U-1476			
	8309572	Apr 27, 2025	U-1476			
	8511304	Jun 14, 2027	DP U-1476			
	8511304*PED	Dec 14, 2027				
	8534281	Mar 08, 2030	DP			
	8534281*PED	Sep 08, 2030				
	8746242	Oct 11, 2030	DP			
	8746242*PED	Apr 11, 2031				
	9333310	Oct 02, 2027	DP			
	9333310*PED	Apr 02, 2028				
	9750726	Nov 29, 2030	DP			
<u>UPADACITINIB - RINVOQ</u>						
N 211675 001	10519164	Oct 17, 2036	DP		I-880	Dec 14, 2024
	10597400	Oct 17, 2036	U-3255		I-883	Jan 14, 2025
	10981923	Oct 17, 2036	DS		I-886	Mar 16, 2025
	10981924	Oct 17, 2036	DP		I-888	Apr 29, 2025
	10995095	Oct 17, 2036	U-3298		I-919	May 18, 2026
	11186584	Oct 17, 2036	DS		NCE	Aug 16, 2024
	11198697	Oct 17, 2036	DP			
	11365198	Oct 17, 2036	U-3275			
	11512092	Oct 17, 2036	U-3275			
	11512092	Oct 17, 2036	U-3371			
	11512092	Oct 17, 2036	U-3487			
	11524964	Oct 17, 2036	U-3371			
	11535624	Oct 17, 2036	U-3255			
	11535625	Oct 17, 2036	U-3298			
	11564922	Mar 09, 2038	U-3624			
	11607411	Mar 09, 2038	U-3341			
	11661425	Oct 17, 2036	DS DP			
	11680069	Oct 17, 2036	DS DP			
	11718627	Oct 17, 2036	DS DP			
	11767326	Oct 17, 2036	U-3298			

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<u>UPADACITINIB - RINVOQ</u>						
N 211675 001	11773105	Oct 17, 2036	DS DP			
	11773106	Oct 17, 2036		U-3275		
	11773106	Oct 17, 2036		U-3371		
	11773106	Oct 17, 2036		U-3487		
	11780847	Oct 17, 2036	DP			
	11780848	Oct 17, 2036		U-3371		
	11787815	Oct 17, 2036	DP			
	11795175	Oct 17, 2036		U-3255		
	8962629	Jan 15, 2031	DS	U-3255		
	8962629	Jan 15, 2031	DS	U-3275		
	8962629	Jan 15, 2031	DS	U-3341		
	8962629	Jan 15, 2031	DS	U-3371		
	8962629	Jan 15, 2031	DS	U-3624		
	9951080	Oct 17, 2036	DS DP			
	9963459	Oct 17, 2036	DP			
	RE47221	Dec 01, 2030	DS			
<u>UPADACITINIB - RINVOQ</u>						
N 211675 002	10344036	Oct 17, 2036	DP		I-883	Jan 14, 2025
	10550126	Oct 17, 2036		U-3298	I-886	Mar 16, 2025
	10730883	Oct 17, 2036	DP		I-919	May 18, 2026
	10981923	Oct 17, 2036	DS		NCE	Aug 16, 2024
	10981924	Oct 17, 2036	DP			
	11186584	Oct 17, 2036	DS			
	11198697	Oct 17, 2036	DP			
	11535626	Oct 17, 2036		U-3298		
	11564922	Mar 09, 2038		U-3624		
	11607411	Mar 09, 2038		U-3341		
	11661425	Oct 17, 2036	DS DP			
	11680069	Oct 17, 2036	DS DP			
	11718627	Oct 17, 2036	DS DP			
	11767326	Oct 17, 2036		U-3298		
	11773105	Oct 17, 2036	DS DP			
	8962629	Jan 15, 2031	DS	U-3341		
	8962629	Jan 15, 2031	DS	U-3624		
	9951080	Oct 17, 2036	DS DP			
	RE47221	Dec 01, 2030	DS			
<u>UPADACITINIB - RINVOQ</u>						
N 211675 003	10202393	Oct 17, 2036	DP		I-886	Mar 16, 2025
	10981923	Oct 17, 2036	DS		I-919	May 18, 2026
	11186584	Oct 17, 2036	DS		NCE	Aug 16, 2024
	11198697	Oct 17, 2036	DP			
	11564922	Mar 09, 2038		U-3624		
	11607411	Mar 09, 2038		U-3341		
	11661425	Oct 17, 2036	DS DP			
	11680069	Oct 17, 2036	DS DP			
	11718627	Oct 17, 2036	DS DP			
	11773105	Oct 17, 2036	DS DP			
	8962629	Jan 15, 2031	DS	U-3341		
	8962629	Jan 15, 2031	DS	U-3624		

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<u>UPADACITINIB - RINVOQ</u>						
N 211675	003 9951080	Oct 17, 2036	DS DP			
	RE47221	Dec 01, 2030	DS			
<u>URIDINE TRIACETATE - VISTOGARD</u>						
N 208159	001 7776838	Aug 17, 2027		U-1791		
<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241	001 10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055		ODE-440	Aug 18, 2030
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037		U-1995		
	10857148	Oct 10, 2037		U-1995		
	10874648	Oct 10, 2037		U-1995		
	10874648	Oct 10, 2037		U-3046		
	10874648	Oct 10, 2037		U-3055		
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037		U-1995		
	10912771	Oct 10, 2037		U-3055		
	10912771	Oct 10, 2037		U-3076		
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040		U-1995		
	10952997	Oct 10, 2037		U-1995		
	10993941	Oct 10, 2037		U-1995		
	11026931	Aug 14, 2039		U-1995		
	11026939	Sep 18, 2038	DP U-1995			
	11026939	Sep 18, 2038	DP U-3055			
	11040029	Oct 10, 2037		U-1995		
	11311532	Sep 18, 2038	DP U-1995			
	11311532	Sep 18, 2038	DP U-3055			
	11439629	Oct 10, 2037		U-3055		
	11654142	Nov 14, 2038		U-3055		
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027		U-1995		
	8357697	Nov 08, 2027		U-3055		
<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241	002 10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055		ODE-440	Aug 18, 2030
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037		U-1995		

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<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241 002	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040	U-1995			
	10952997	Oct 10, 2037	U-1995			
	10993941	Oct 10, 2037	U-1995			
	11026931	Aug 14, 2039	U-1995			
	11026939	Sep 18, 2038	DP U-1995			
	11026939	Sep 18, 2038	DP U-3055			
	11040029	Oct 10, 2037	U-1995			
	11311532	Sep 18, 2038	DP U-1995			
	11311532	Sep 18, 2038	DP U-3055			
	11439629	Oct 10, 2037	U-3055			
	11654142	Nov 14, 2038	U-3055			
	8039627	Apr 11, 2031	DS DP			
	8357697	Nov 08, 2027	U-1995			
	8357697	Nov 08, 2027	U-3055			
<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241 003	10065952	Oct 28, 2036	DS DP U-1995		I-925	Aug 18, 2026
	10065952	Oct 28, 2036	DS DP U-3055		ODE-440	Aug 18, 2030
	10844058	Oct 28, 2036	DS DP U-1995			
	10844058	Oct 28, 2036	DS DP U-3055			
	10851103	Oct 28, 2036	DS DP U-1995			
	10851103	Oct 28, 2036	DS DP U-3055			
	10851104	Oct 28, 2036	DS U-1995			
	10851104	Oct 28, 2036	DS U-3055			
	10857137	Oct 10, 2037	U-1995			
	10857148	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-1995			
	10874648	Oct 10, 2037	U-3046			
	10874648	Oct 10, 2037	U-3055			
	10906902	Dec 22, 2036	DS DP			
	10906903	Dec 22, 2036	DS DP			
	10912771	Oct 10, 2037	U-1995			
	10912771	Oct 10, 2037	U-3055			
	10912771	Oct 10, 2037	U-3076			
	10919892	Dec 22, 2036	DS DP			
	10940141	Aug 10, 2040	U-1995			
	10952997	Oct 10, 2037	U-1995			
	10993941	Oct 10, 2037	U-1995			
	11026931	Aug 14, 2039	U-1995			
	11026939	Sep 18, 2038	DP U-1995			

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<u>VALBENZAZINE TOSYLATE - INGREZZA</u>						
N 209241	003	11026939	Sep 18, 2038	DP U-3055		
		11040029	Oct 10, 2037	U-1995		
		11311532	Sep 18, 2038	DP U-1995		
		11311532	Sep 18, 2038	DP U-3055		
		11439629	Oct 10, 2037	U-3055		
		11654142	Nov 14, 2038	U-3055		
		8039627	Apr 11, 2031	DS DP		
		8357697	Nov 08, 2027	U-1995		
		8357697	Nov 08, 2027	U-3055		
<u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u>						
N 022257	001	8889109	Dec 11, 2027	DP		
		9642911	Dec 11, 2027	DP		
<u>VALSARTAN - DIOVAN</u>						
N 021283	001				NPP	Apr 19, 2024
<u>VALSARTAN - DIOVAN</u>						
N 021283	002				NPP	Apr 19, 2024
<u>VALSARTAN - DIOVAN</u>						
N 021283	004				NPP	Apr 19, 2024
<u>VAMOROLONE - AGAMREE</u>						
N 215239	001	10857161	May 28, 2029	DP U-3747	NCE	Oct 26, 2028
		11382922	Jul 16, 2040	DS DP	ODE-450	Oct 26, 2030
		11471471	Mar 17, 2040	U-3747		
		11690853	Jun 29, 2036	U-3747		
		11833159	May 28, 2029	DP		
		8334279	May 28, 2029	U-3747		
<u>VANCOMYCIN - VANCOMYCIN</u>						
N 213895	001	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
		11000567	Nov 06, 2035	DP		
		11517609	Nov 06, 2035	U-282		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
A 214913	001				PC	Feb 25, 2024
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
A 214913	002				PC	Feb 25, 2024
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
A 217489	001				CGT	Feb 12, 2024
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
A 217489	002				CGT	Feb 12, 2024
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910	001	10493028	Mar 13, 2035	DP		
		10688046	Mar 13, 2035	DP		
		10959946	Mar 13, 2035	DP		
		10959947	Mar 13, 2035	DP U-3104		
		10959947	Mar 13, 2035	DP U-3105		

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<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910	001	10959948	Mar 13, 2035	DP	U-3104	
		10959948	Mar 13, 2035	DP	U-3105	
		10959949	Mar 13, 2035	DP	U-3104	
		10959949	Mar 13, 2035	DP	U-3105	
		11638692	Mar 13, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - FIRVANO KIT</u>						
N 208910	002	10493028	Mar 13, 2035	DP		
		10688046	Mar 13, 2035	DP		
		10959946	Mar 13, 2035	DP		
		10959947	Mar 13, 2035	DP	U-3104	
		10959947	Mar 13, 2035	DP	U-3105	
		10959948	Mar 13, 2035	DP	U-3104	
		10959948	Mar 13, 2035	DP	U-3105	
		10959949	Mar 13, 2035	DP	U-3104	
		10959949	Mar 13, 2035	DP	U-3105	
		11638692	Mar 13, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	001	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	002	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	003	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	004	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	005	10039804	Nov 06, 2035	DP	U-282	
		10188697	Nov 06, 2035	DP	U-282	
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035		U-282	
		11628200	Nov 06, 2035	DP		

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<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	006	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035	U-282		
		11628200	Nov 06, 2035	DP		
<u>VANCOMYCIN HYDROCHLORIDE - VANCOMYCIN HYDROCHLORIDE</u>						
N 211962	007	10039804	Nov 06, 2035	DP U-282		
		10188697	Nov 06, 2035	DP U-282		
		10849956	Nov 06, 2035	DP		
		11517609	Nov 06, 2035	U-282		
		11628200	Nov 06, 2035	DP		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	001	8067427	Aug 08, 2028	DP		
<u>VANDETANIB - CAPRELSA</u>						
N 022405	002	8067427	Aug 08, 2028	DP		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	001	8273876	Jul 23, 2027	U-1288		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	002	8273876	Jul 23, 2027	U-1288		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	003	8273876	Jul 23, 2027	U-1288		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>						
N 021400	004	8273876	Jul 23, 2027	U-1288		
<u>VARDENAFIL HYDROCHLORIDE - STAXYN</u>						
N 200179	001	8613950	Dec 23, 2028	DP		
<u>VARENICLINE TARTRATE - TYRVAYA</u>						
N 213978	001	10456396	Oct 19, 2035	DP U-1900	NP	Oct 15, 2024
		11224598	Oct 19, 2035	U-1900		
		9504644	Oct 19, 2035	U-1900		
		9504645	Oct 19, 2035	DP		
		9532944	Oct 19, 2035	U-1900		
		9597284	Oct 19, 2035	U-1900		
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	001	9375478	Jan 30, 2035	U-1857		
		9687526	Jan 30, 2035	U-1857		
		9744209	Jan 30, 2035	U-1857		
		9744239	Jan 30, 2035	U-1857		
		9750785	Jan 30, 2035	DP		
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485	002	9375478	Jan 30, 2035	U-1857		
		9687526	Jan 30, 2035	U-1857		
		9744209	Jan 30, 2035	U-1857		
		9744239	Jan 30, 2035	U-1857		
		9750785	Jan 30, 2035	DP		
		9937223	Jan 30, 2035	U-1857		

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<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485 003	10010575	Jan 30, 2035	U-1857			
	9919026	Jan 30, 2035	DP			
	9925233	Jan 30, 2035	U-1857			
	9925234	Jan 30, 2035	U-1857			
	9962422	Jan 30, 2035	U-1857			
	9968649	Jan 30, 2035	U-1857			
	9974827	Jan 30, 2035	U-1857			
	9981006	Jan 30, 2035	U-1857			
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485 004	10010575	Jan 30, 2035	U-1857			
	9919026	Jan 30, 2035	DP			
	9925233	Jan 30, 2035	U-1857			
	9925234	Jan 30, 2035	U-1857			
	9962422	Jan 30, 2035	U-1857			
	9968649	Jan 30, 2035	U-1857			
	9974827	Jan 30, 2035	U-1857			
	9981006	Jan 30, 2035	U-1857			
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485 005	10010575	Jan 30, 2035	U-1857			
	9919026	Jan 30, 2035	DP			
	9925233	Jan 30, 2035	U-1857			
	9925234	Jan 30, 2035	U-1857			
	9962422	Jan 30, 2035	U-1857			
	9968649	Jan 30, 2035	U-1857			
	9974827	Jan 30, 2035	U-1857			
	9981006	Jan 30, 2035	U-1857			
<u>VASOPRESSIN - VASOSTRICT</u>						
N 204485 006	10010575	Jan 30, 2035	U-1857			
	9919026	Jan 30, 2035	DP			
	9925233	Jan 30, 2035	U-1857			
	9925234	Jan 30, 2035	U-1857			
	9962422	Jan 30, 2035	U-1857			
	9968649	Jan 30, 2035	U-1857			
	9974827	Jan 30, 2035	U-1857			
	9981006	Jan 30, 2035	U-1857			
<u>VEMURAFENIB - ZELBORAF</u>						
N 202429 001	7504509	Oct 22, 2026	DS DP		ODE-158	Nov 06, 2024
	7863288	Jun 20, 2029	DS DP			
	8143271	Jun 21, 2026	DS DP			
	8470818	Aug 02, 2026			U-1418	
	8470818	Aug 02, 2026			U-2164	
	8741920	Jul 27, 2030	DS DP			
	9447089	Jun 06, 2032	DP			
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573 001	10730873	Nov 21, 2031	DS		ODE-185	Jun 08, 2025
	10993942	Sep 06, 2033	U-3114		ODE-211	Nov 21, 2025
	11110087	Sep 06, 2033	U-3222		ODE-239	May 15, 2026
	11110087	Sep 06, 2033	U-3223			

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<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	001	11369599	May 23, 2032	DP		
		11413282	Sep 06, 2033		U-3412	
		11590128	Sep 06, 2033		U-3548	
		8546399	Jun 27, 2031	DS DP		
		8722657	Jan 29, 2032	DS		
		9174982	May 26, 2030		U-2323	
		9174982	May 26, 2030		U-2445	
		9174982	May 26, 2030		U-2446	
		9174982	May 26, 2030		U-2537	
		9539251	Sep 06, 2033		U-2538	
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	002	10730873	Nov 21, 2031	DS	ODE-185	Jun 08, 2025
		10993942	Sep 06, 2033		ODE-211	Nov 21, 2025
		11110087	Sep 06, 2033		ODE-239	May 15, 2026
		11110087	Sep 06, 2033		U-3222	
		11110087	Sep 06, 2033		U-3223	
		11369599	May 23, 2032	DP		
		11413282	Sep 06, 2033		U-3412	
		11590128	Sep 06, 2033		U-3548	
		8546399	Jun 27, 2031	DS DP		
		8722657	Jan 29, 2032	DS		
		9174982	May 26, 2030		U-2323	
		9174982	May 26, 2030		U-2445	
		9174982	May 26, 2030		U-2446	
		9174982	May 26, 2030		U-2537	
		9539251	Sep 06, 2033		U-2538	
<u>VENETOCLAX - VENCLEXTA</u>						
N 208573	003	10730873	Nov 21, 2031	DS	ODE-185	Jun 08, 2025
		10993942	Sep 06, 2033		ODE-211	Nov 21, 2025
		11110087	Sep 06, 2033		ODE-239	May 15, 2026
		11110087	Sep 06, 2033		U-3222	
		11110087	Sep 06, 2033		U-3223	
		11369599	May 23, 2032	DP		
		11413282	Sep 06, 2033		U-3412	
		11590128	Sep 06, 2033		U-3548	
		8546399	Jun 27, 2031	DS DP		
		8722657	Jan 29, 2032	DS		
		9174982	May 26, 2030		U-2323	
		9174982	May 26, 2030		U-2445	
		9174982	May 26, 2030		U-2446	
		9174982	May 26, 2030		U-2537	
		9539251	Sep 06, 2033		U-2538	
<u>VENLAFAXINE BESYLATE - VENLAFAXINE BESYLATE</u>						
N 215429	001	7776358	May 16, 2028	DP		
<u>VERICIGUAT - VEROUVO</u>						
N 214377	001	10736896	May 19, 2031	DS DP	NCE	Jan 19, 2026
		11439642	May 19, 2031		U-3062	
		8420656	May 19, 2031	DS DP		
		8921377	May 19, 2031		U-3062	
		9604948	Nov 26, 2032	DS DP	U-3062	

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<u>VERICIGUAT - VEROUVO</u>						
N 214377	001 9993476	May 19, 2031	U-3062			
<u>VERICIGUAT - VEROUVO</u>						
N 214377	002 10736896	May 19, 2031	DS DP		NCE	Jan 19, 2026
	11439642	May 19, 2031	U-3062			
	8420656	May 19, 2031	DS DP			
	8921377	May 19, 2031	U-3062			
	9604948	Nov 26, 2032	DS DP U-3062			
	9993476	May 19, 2031	U-3062			
<u>VERICIGUAT - VEROUVO</u>						
N 214377	003 10736896	May 19, 2031	DS DP		NCE	Jan 19, 2026
	11439642	May 19, 2031	U-3062			
	8420656	May 19, 2031	DS DP			
	8921377	May 19, 2031	U-3062			
	9604948	Nov 26, 2032	DS DP U-3062			
	9993476	May 19, 2031	U-3062			
<u>VIBEGRON - GEMTESA</u>						
N 213006	001 8247415	Dec 01, 2030	DS DP U-3045		NCE	Dec 23, 2025
	8653260	Apr 02, 2029	DS			
<u>VILOXAZINE HYDROCHLORIDE - OELBREE</u>						
N 211964	001 11324753	Sep 04, 2029	U-727		NCE	Apr 02, 2026
	11458143	Sep 04, 2029	U-727		NPP	Apr 29, 2025
	9358204	Feb 07, 2033	DP			
	9603853	Feb 07, 2033	U-727			
	9662338	Feb 07, 2033	DP			
<u>VILOXAZINE HYDROCHLORIDE - OELBREE</u>						
N 211964	002 11324753	Sep 04, 2029	U-727		NCE	Apr 02, 2026
	11458143	Sep 04, 2029	U-727		NPP	Apr 29, 2025
	9358204	Feb 07, 2033	DP			
	9603853	Feb 07, 2033	U-727			
	9662338	Feb 07, 2033	DP			
<u>VILOXAZINE HYDROCHLORIDE - OELBREE</u>						
N 211964	003 11324753	Sep 04, 2029	U-727		NCE	Apr 02, 2026
	11458143	Sep 04, 2029	U-727		NPP	Apr 29, 2025
	9358204	Feb 07, 2033	DP			
	9603853	Feb 07, 2033	U-727			
	9662338	Feb 07, 2033	DP			
<u>VILTOLARSEN - VILTEPSO</u>						
N 212154	001 10870676	Aug 31, 2031	DS DP U-3039		NCE	Aug 12, 2025
	9079934	Aug 31, 2031	DS DP		ODE-280	Aug 12, 2027
<u>VISMODEGIB - ERIVEDGE</u>						
N 203388	001 7888364	Nov 11, 2028	DS DP			
	9278961	Dec 15, 2028	U-1825			
	9790183	Sep 02, 2025	U-3109			
<u>VOCLOSPORIN - LUPKYNIS</u>						
N 213716	001 10286036	Dec 07, 2037	U-3056		NCE	Jan 22, 2026
	11622991	Dec 07, 2037	U-3056			

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<u>VOCLOSPORIN - LUPKYNIS</u>						
N 213716	001 7332472	Oct 17, 2024	DS DP U-3056			
<u>VONOPRAZAN FUMARATE - VOQUEZNA</u>						
N 215151	001 7977488	Aug 11, 2028	DS		NCE	May 03, 2027
	9186411	Aug 11, 2030	DP		NP	Nov 01, 2026
<u>VONOPRAZAN FUMARATE - VOQUEZNA</u>						
N 215151	002 7977488	Aug 11, 2028	DS		NCE	May 03, 2027
	9186411	Aug 11, 2030	DP		NP	Nov 01, 2026
<u>VORAPAXAR SULFATE - ZONTIVITY</u>						
N 204886	001 7304078	Dec 23, 2027	DS DP U-1512			
	7713999	May 30, 2024	DS DP U-2291			
<u>VORINOSTAT - ZOLINZA</u>						
N 021991	001 7399787	Feb 09, 2025		U-892		
	7456219	Mar 11, 2027	DS			
	7851509	Feb 21, 2024	DP U-892			
	8093295	May 16, 2026	DP			
	8450372	Mar 18, 2028		U-892		
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447	001 11458134	Jun 15, 2027	DP U-3463		M-187	Jan 22, 2024
	11458134*PED	Dec 15, 2027			M-232	Aug 23, 2026
	7144884	Jun 17, 2026	DS DP U-1439		PED	Feb 23, 2027
	7144884*PED	Dec 17, 2026				
	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027		U-1668		
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027		U-2309		
	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027		U-2309		
	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027		U-2309		
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027		U-1668		
	9227946*PED	Dec 15, 2027				
	9278096	Mar 21, 2032		U-2436		
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027		U-1668		
	9861630*PED	Dec 15, 2027				
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447	002 11458134	Jun 15, 2027	DP U-3463		M-187	Jan 22, 2024
	11458134*PED	Dec 15, 2027			M-232	Aug 23, 2026
	7144884	Jun 17, 2026	DS DP U-1439		PED	Feb 23, 2027
	7144884*PED	Dec 17, 2026				
	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027		U-1668		
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027		U-2309		

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<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 002	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027	U-2309			
	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027	U-2309			
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027	U-1668			
	9227946*PED	Dec 15, 2027				
	9278096	Mar 21, 2032	U-2436			
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027	U-1668			
	9861630*PED	Dec 15, 2027				
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 003	11458134	Jun 15, 2027	DP U-3463		M-187	Jan 22, 2024
	11458134*PED	Dec 15, 2027			M-232	Aug 23, 2026
	7144884	Jun 17, 2026	DS DP U-1439		PED	Feb 23, 2027
	7144884*PED	Dec 17, 2026				
	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027	U-1668			
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027	U-2309			
	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027	U-2309			
	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027	U-2309			
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027	U-1668			
	9227946*PED	Dec 15, 2027				
	9278096	Mar 21, 2032	U-2436			
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027	U-1668			
	9861630*PED	Dec 15, 2027				
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447 004	11458134	Jun 15, 2027	DP U-3463		M-187	Jan 22, 2024
	11458134*PED	Dec 15, 2027			M-232	Aug 23, 2026
	7144884	Jun 17, 2026	DS DP U-1439		PED	Feb 23, 2027
	7144884*PED	Dec 17, 2026				
	8722684	Jun 30, 2031	DS DP			
	8722684*PED	Dec 30, 2031				
	8969355	Jun 15, 2027	U-1668			
	8969355*PED	Dec 15, 2027				
	9125908	Jun 15, 2027	U-2309			
	9125908*PED	Dec 15, 2027				
	9125909	Jun 15, 2027	U-2309			
	9125909*PED	Dec 15, 2027				
	9125910	Jun 15, 2027	U-2309			
	9125910*PED	Dec 15, 2027				
	9227946	Jun 15, 2027	U-1668			
	9227946*PED	Dec 15, 2027				

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APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u>						
N 204447	004 9278096	Mar 21, 2032	U-2436			
	9278096*PED	Sep 21, 2032				
	9861630	Jun 15, 2027	U-1668			
	9861630*PED	Dec 15, 2027				
<u>VOSORITIDE - VOXZOGO</u>						
N 214938	001 10646550	Aug 01, 2036	DP U-3256		NCE	Nov 19, 2026
	8198242	Jun 11, 2030	DS DP U-3256		NPP	Oct 20, 2026
	9907834	Aug 01, 2036	DP		ODE-387	Nov 19, 2028
	RE48267	May 20, 2030	U-3256		ODE-449	Oct 20, 2030
<u>VOSORITIDE - VOXZOGO</u>						
N 214938	002 10646550	Aug 01, 2036	DP U-3256		NCE	Nov 19, 2026
	8198242	Jun 11, 2030	DS DP U-3256		NPP	Oct 20, 2026
	9907834	Aug 01, 2036	DP		ODE-387	Nov 19, 2028
	RE48267	May 20, 2030	U-3256		ODE-449	Oct 20, 2030
<u>VOSORITIDE - VOXZOGO</u>						
N 214938	003 10646550	Aug 01, 2036	DP U-3256		NCE	Nov 19, 2026
	8198242	Jun 11, 2030	DS DP U-3256		NPP	Oct 20, 2026
	9907834	Aug 01, 2036	DP		ODE-387	Nov 19, 2028
	RE48267	May 20, 2030	U-3256		ODE-449	Oct 20, 2030
<u>VOXELOTOR - OXBRYTA</u>						
N 213137	001 10017491	Dec 28, 2032	DP		NCE	Nov 25, 2024
	10034879	Dec 28, 2032	DS DP		ODE-281	Nov 25, 2026
	10493035	Oct 12, 2037	DP		ODE-394	Dec 17, 2028
	10722502	Feb 06, 2035	DP			
	10806733	Dec 28, 2032	DS			
	11020382	Dec 02, 2036	U-3133			
	11020382	Dec 02, 2036	U-3134			
	11452720	Feb 06, 2035	U-3459			
	9018210	Nov 25, 2033	DS DP			
	9248199	Jan 29, 2034	U-2676			
	9248199	Jan 29, 2034	U-2715			
	9447071	Feb 06, 2035	DS DP			
<u>VOXELOTOR - OXBRYTA</u>						
N 213137	002 10017491	Dec 28, 2032	DP		NCE	Nov 25, 2024
	10034879	Dec 28, 2032	DS DP		ODE-394	Dec 17, 2028
	10493035	Oct 12, 2037	DP			
	10722502	Feb 06, 2035	DP			
	10806733	Dec 28, 2032	DS			
	11020382	Dec 02, 2036	U-3133			
	11020382	Dec 02, 2036	U-3134			
	11452720	Feb 06, 2035	U-3459			
	9018210	Nov 25, 2033	DS DP			
	9248199	Jan 29, 2034	U-2676			
	9248199	Jan 29, 2034	U-2715			
	9447071	Feb 06, 2035	DS DP			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

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<u>VOXELOTOR - OXBRYTA</u>						
N 216157 001	10017491	Dec 28, 2032	DP		NCE	Nov 25, 2024
	10034879	Dec 28, 2032	DS DP		ODE-394	Dec 17, 2028
	10722502	Feb 06, 2035	DP			
	10806733	Dec 28, 2032	DS			
	11020382	Dec 02, 2036		U-3133		
	11020382	Dec 02, 2036		U-3134		
	11452720	Feb 06, 2035		U-3459		
	9018210	Nov 25, 2033	DS DP			
	9248199	Jan 29, 2034		U-2676		
	9248199	Jan 29, 2034		U-2715		
	9447071	Feb 06, 2035	DS DP			
<u>VUTRISIRAN SODIUM - AMVUTTRA</u>						
N 215515 001	10131907	Aug 24, 2028	DS DP U-3396		NCE	Jun 13, 2027
	10208307	Jul 28, 2036	DS DP U-3396		ODE-212	Jun 13, 2029
	10570391	Nov 16, 2032	DS DP U-3396			
	10612024	Aug 14, 2035	DS DP U-3396			
	10683501	Jul 28, 2036	DS DP U-3396			
	10806791	Dec 04, 2028	DS			
	11286486	Jul 28, 2036	DS DP U-3396			
	11401517	Aug 14, 2035	DS DP U-3396			
	8106022	Dec 12, 2029	DS DP U-3396			
	8828956	Dec 04, 2028	DS DP U-3396			
	9370581	Dec 04, 2028	DS DP U-3396			
	9399775	Nov 16, 2032	DS DP U-3396			
<u>XENON XE-129 HYPERPOLARIZED - XENOVUE</u>						
N 214375 001	10583205	Feb 20, 2035	DP		NCE	Dec 23, 2027
	11052161	Dec 29, 2035	DP			
<u>ZANUBRUTINIB - BRUKINSA</u>						
N 213217 001	10570139	Apr 22, 2034		U-1745	I-817	Jan 19, 2026
	10570139	Apr 22, 2034		U-2145	I-871	Aug 31, 2024
	10570139	Apr 22, 2034		U-2537	I-874	Sep 14, 2024
	10570139	Apr 22, 2034		U-2666	NCE	Nov 14, 2024
	10570139	Apr 22, 2034		U-3063	ODE-274	Jan 19, 2030
	10570139	Apr 22, 2034		U-3486	ODE-276	Nov 14, 2026
	10927117	Aug 15, 2037	DS DP		ODE-370	Sep 14, 2028
	11142528	Apr 22, 2034	DP U-1745		ODE-371	Aug 31, 2028
	11142528	Apr 22, 2034	DP U-2145			
	11142528	Apr 22, 2034	DP U-2537			
	11142528	Apr 22, 2034	DP U-2666			
	11142528	Apr 22, 2034	DP U-3063			
	11142528	Apr 22, 2034	DP U-3486			
	11591340	Aug 15, 2037		U-1745		
	11591340	Aug 15, 2037		U-2145		
	11591340	Aug 15, 2037		U-2537		
	11591340	Aug 15, 2037		U-2666		
	11591340	Aug 15, 2037		U-3063		
	11591340	Aug 15, 2037		U-3486		
	11786531	Jan 19, 2043		U-3715		

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<u>ZANUBRUTINIB - BRUKINSA</u>						
N 213217	001	11786531	Jan 19, 2043	U-3716		
		11786531	Jan 19, 2043	U-3717		
		11786531	Jan 19, 2043	U-3718		
		11786531	Jan 19, 2043	U-3719		
		11786531	Jan 19, 2043	U-3720		
		9447106	Apr 22, 2034	DS DP U-1745		
		9447106	Apr 22, 2034	DS DP U-2145		
		9447106	Apr 22, 2034	DS DP U-2537		
		9447106	Apr 22, 2034	DS DP U-2666		
		9447106	Apr 22, 2034	DS DP U-3063		
		9447106	Apr 22, 2034	DS DP U-3486		
<u>ZAVEGEPANT HYDROCHLORIDE - ZAVZPRET</u>						
N 216386	001	7220862	Jan 21, 2024	DS DP	NCE	Mar 09, 2028
		8481546	Oct 07, 2031	DS DP U-3555		
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	001	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	002	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	003	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZICONOTIDE ACETATE - PRIALT</u>						
N 021060	004	8653033	Oct 01, 2024	U-48		
		8653033	Oct 01, 2024	U-55		
		8765680	Oct 01, 2024	U-48		
		8765680	Oct 01, 2024	U-55		
		9707270	Oct 01, 2024	U-2084		
<u>ZILUCOPLAN SODIUM - ZILBRYSO</u>						
N 216834	001	10106579	Jun 12, 2035	DS DP U-3735	NCE	Oct 17, 2028
		10208089	Jun 12, 2035	DS U-3735	ODE-446	Oct 17, 2030
		10435438	Jun 12, 2035	DS U-3735		
		10562934	Jun 12, 2035	DS		
		10835574	Jun 12, 2035	DP		
		11014965	Jun 12, 2035	DS U-3735		
		11535650	Jun 12, 2035	DP		
		11752190	Jun 12, 2035	DP U-3735		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ZILUCOPLAN SODIUM - ZILBRYSO</u>						
N 216834 002	10106579	Jun 12, 2035	DS DP U-3735		NCE	Oct 17, 2028
	10208089	Jun 12, 2035	DS U-3735		ODE-446	Oct 17, 2030
	10435438	Jun 12, 2035	DS U-3735			
	10562934	Jun 12, 2035	DS			
	10835574	Jun 12, 2035	DP			
	11014965	Jun 12, 2035	DS U-3735			
	11535650	Jun 12, 2035	DP			
	11752190	Jun 12, 2035	DP U-3735			
<u>ZILUCOPLAN SODIUM - ZILBRYSO</u>						
N 216834 003	10106579	Jun 12, 2035	DS DP U-3735		NCE	Oct 17, 2028
	10208089	Jun 12, 2035	DS U-3735		ODE-446	Oct 17, 2030
	10435438	Jun 12, 2035	DS U-3735			
	10562934	Jun 12, 2035	DS			
	10835574	Jun 12, 2035	DP			
	11014965	Jun 12, 2035	DS U-3735			
	11535650	Jun 12, 2035	DP			
	11752190	Jun 12, 2035	DP U-3735			
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 001					M-232	Jan 28, 2025
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 002					M-232	Jan 28, 2025
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 003					M-232	Jan 28, 2025
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>						
N 020825 004					M-232	Jan 28, 2025
<u>ZOLEDRONIC ACID - ZOMETA</u>						
N 021223 002	8324189	May 29, 2025	U-1308			
	8324189	May 29, 2025	U-1309			
	8324189	May 29, 2025	U-53			
<u>ZOLEDRONIC ACID - ZOMETA</u>						
N 021223 003	7932241	Feb 05, 2028	DP			
	8324189	May 29, 2025	U-1308			
	8324189	May 29, 2025	U-1309			
	8324189	May 29, 2025	U-53			
<u>ZOLEDRONIC ACID - RECLAST</u>						
N 021817 001	7932241	Feb 05, 2028	DP			
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997 001	9265720	Feb 25, 2031	U-674			
	9597281	Apr 06, 2027	U-674			
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
N 021997 002	9265720	Feb 25, 2031	U-674			
	9597281	Apr 06, 2027	U-674			
<u>ZOLPIDEM TARTRATE - ZOLPIMIST</u>						
N 022196 001	8236285	Aug 07, 2032	DS DP U-70			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>ZOLPIDEM TARTRATE - INTERMEZZO</u>						
N 022328	001	7658945	Apr 15, 2027	DP U-1194		
		7682628	Feb 16, 2025	U-1194		
		8242131	Aug 20, 2029	U-1266		
		8252809	Feb 16, 2025	DP		
<u>ZOLPIDEM TARTRATE - INTERMEZZO</u>						
N 022328	002	7658945	Apr 15, 2027	DP U-1194		
		7682628	Feb 16, 2025	U-1194		
		8242131	Aug 20, 2029	U-1266		
		8252809	Feb 16, 2025	DP		
<u>ZONISAMIDE - ZONISADE</u>						
N 214273	001	11478456	Aug 18, 2038	U-3458		
		11529333	Aug 18, 2038	DP		
<u>ZURANOLONE - ZURZUVAE</u>						
N 217369	001	10172871	Apr 17, 2034	U-2552	NCE	Oct 31, 2028
		10342810	Apr 17, 2034	U-2552		
		11236121	Aug 23, 2037	DS		
		9512165	Apr 17, 2034	DS DP		
<u>ZURANOLONE - ZURZUVAE</u>						
N 217369	002	10172871	Apr 17, 2034	U-2552	NCE	Oct 31, 2028
		10342810	Apr 17, 2034	U-2552		
		11236121	Aug 23, 2037	DS		
		9512165	Apr 17, 2034	DS DP		
<u>ZURANOLONE - ZURZUVAE</u>						
N 217369	003	10172871	Apr 17, 2034	U-2552	NCE	Oct 31, 2028
		10342810	Apr 17, 2034	U-2552		
		11236121	Aug 23, 2037	DS		
		9512165	Apr 17, 2034	DS DP		

PATENT AND EXCLUSIVITY TERMS

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PATENT & EXCLUSIVITY ABBREVIATIONS

CGT	COMPETITIVE GENERIC THERAPY
D	NEW DOSING SCHEDULE (SEE INDIVIDUAL REFERENCES)
GAIN	GAIN EXCLUSIVITY
I	NEW INDICATION (SEE INDIVIDUAL REFERENCES)
M	MISCELLANEOUS EXCLUSIVITY CODES (SEE INDIVIDUAL REFERENCES)
NC	NEW COMBINATION
NCE	NEW CHEMICAL ENTITY
NCE*	NEW CHEMICAL ENTITY (AN ENANTIOMER OF PREVIOUSLY APPROVED RACEMIC MIXTURE. SEE SECTION 505(U) OF THE FEDERAL FOOD AND DRUG COSMETIC ACT).
NDF	NEW DOSAGE FORM
NE	NEW ESTER OR SALT OF AN ACTIVE INGREDIENT
NP	NEW PRODUCT
NP*	NEW PRODUCT (MINT FLAVORED)
NPP	NEW PATIENT POPULATION
NR	NEW ROUTE
NS	NEW STRENGTH
ODE	ORPHAN DRUG EXCLUSIVITY (SEE INDIVIDUAL REFERENCES)
ODE*	FDA has not recognized Orphan-Drug Exclusivity (ODE) for this drug, but it contains the same active moiety or moieties as another drug(s) that was eligible for ODE, and also shares ODE-protected use(s) or indication(s) with that drug(s). An application seeking approval for the same active moiety or moieties, including an ANDA that cites this NDA as its basis of submission, may not be approved for such ODE-protected use(s) and indication(s)
PC	PATENT CHALLENGE
PED	PEDIATRIC EXCLUSIVITY
RTO	RX TO OTC SWITCH OR OTC USE
RTO*	OTC USE FOR WOMEN AGES 15 AND 16
RTO**	OTC USE FOR WOMEN 14 AND BELOW
U	PATENT USE CODE (SEE INDIVIDUAL REFERENCES)
W	EXCLUSIVITY ON THIS APPLICATION EXPIRING ON THIS DATE HAS BEEN WAIVED BY SPONSOR - SEE SECTION 1.8 OF ORANGE BOOK PREFACE WAIVED EXCLUSIVITY

EXCLUSIVITY DOSING SCHEDULE

D-1	ONCE A DAY APPLICATION
D-2	ONCE DAILY DOSING
D-3	SEVEN DAYS/SEVEN DAYS/SEVEN DAYS DOSING SCHEDULE
D-4	SEVEN DAYS/FOURTEEN DAYS DOSING SCHEDULE
D-5	TEN DAYS/ELEVEN DAYS DOSING SCHEDULE
D-6	SEVEN DAYS/NINE DAYS/FIVE DAYS DOSING SCHEDULE
D-7	BID DOSING
D-8	INTRAVENOUS, EPIDURAL AND INTRATHECAL DOSING
D-9	NARCOTIC OVERDOSE IN ADULTS
D-10	NARCOTIC OVERDOSE IN CHILDREN
D-11	POSTOPERATIVE NARCOTIC DEPRESSION IN CHILDREN
D-12	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE DUODENAL ULCER
D-13	INCREASED MAXIMUM DAILY DOSAGE RECOMMENDATION
D-14	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
D-15	SINGLE DAILY DOSE OF 25MG/37.5MG
D-16	CONTINUOUS INTRAVENOUS INFUSION
D-17	400MG EVERY 12 HOURS FOR THREE DAYS FOR UNCOMPLICATED URINARY TRACT INFECTIONS
D-18	LOWER RECOMMENDED STARTING DOSE GUIDELINES
D-19	BOLUS DOSING GUIDELINES
D-20	SINGLE 32MG DOSE
D-21	ALTERNATIVE DOSAGE OF 300MG ONCE DAILY AFTER THE EVENING MEAL

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-22 REDUCTION IN INFUSION TIME FROM 24 TO 4 HOURS FOR THE 60MG DOSE
- D-23 INCREASE MAXIMUM DOSE AND VARIATIONS IN THE DOSING REGIMEN
- D-24 FOR OVARIAN CANCER THE RECOMMENDED REGIMEN IS 135MG/M2 OR 175MG/M2 INTRAVENOUSLY OVER THREE HOURS EVERY THREE WEEKS
- D-25 ADDITIONAL DOSAGE REGIMEN EQUAL TO HALF THE ORIGINAL DOSING REGIMEN
- D-26 ONCE WEEKLY APPLICATION
- D-27 BID DOSING IN PATIENTS 12 YEARS OF AGE AND OLDER FOR PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH MODERATE EMETOGENIC CANCER CHEMOTHERAPY
- D-28 USE OF ISOVUE-370 IN EXCRETORY UROGRAPHY AT EQUIVALENT GRAMS OF IODINE TO THE CURRENTLY APPROVED ISOVUE-250 AND ISOVUE-300
- D-29 INCREASE OF CUMULATIVE DOSE TO 0.3MMOL/KG FOR MRI OF CNS IN ADULTS
- D-30 5000 IU DOSE FOR PROPHYLAXIS AGAINST DEEP VEIN THROMBOSIS
- D-31 CHANGE IN RECOMMENDED TOTAL DAILY DOSE TO 80MG (40MG BID)
- D-32 REMOVAL OF THE RESTRICTIONS LIMITING TREATMENT TO TWO CONSECUTIVE WEEKS AND TO SMALL AREAS
- D-33 ONCE DAILY DOSING FOR PLAQUE PSORIASIS
- D-34 EVERY FOUR MONTHS DOSAGE REGIMEN
- D-35 FOR A ONE WEEK DOSING OF INTERDIGITAL TINEA PEDIS
- D-36 FOR A SINGLE 2MG DOSE AS AN ALTERNATIVE TO THE 1MG DOSE GIVEN TWICE DAILY
- D-37 DOSING REGIMEN FOR ADMINISTRATION EITHER ONCE DAILY (QD) OR TWICE DAILY (BID)
- D-38 CONTINUOUS INFUSION AS AN ALTERNATE METHOD OF ADMINISTRATION
- D-39 CHANGE IN TIME TO TAKE THE DRUG PRIOR TO A MEAL TO PREVENT MEAL-INDUCED HEARTBURN SYMPTOMS FROM ".1/2 TO 1 HOUR BEFORE EATING" TO ".. RIGHT BEFORE EATING OR UP TO 60MIN BEFORE CONSUMING..."
- D-40 ONCE-A-DAY DOSING REGIMEN
- D-41 DRUG MAY BE DOSED RIGHT BEFORE A MEAL OR ANY TIME UP TO 30MIN BEFORE EATING OR DRINKING FOOD AND BEVERAGES THAT WOULD BE EXPECTED TO CAUSE SYMPTOMS
- D-42 TEN DAY DOSING REGIMEN FOR TRIPLE THERAPY, PREVACID IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN, FOR THE ERADICATION OF H.PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- D-43 INITIATION OF TREATMENT WITH 900MG/DAY BY DELETION OF THE REQUIREMENT TO TITRATE TO 900MG/DAY OVER A 3-DAY PERIOD
- D-44 IN A CLINICAL TRIAL, FEWER DISCONTINUATIONS DUE TO ADVERSE EVENTS, ESPECIALLY DIZZINESS AND VERTIGO, WERE OBSERVED WHEN TITRATING THE DOSE IN INCREMENTS OF 50MG/DAY EVERY 3 DAYS UNTIL AN EFFECTIVE DOSE (NOT EXCEEDING 400MG/DAY) WAS REACHED
- D-45 ONCE DAILY DOSING FOR MAINTENANCE ONLY
- D-46 NEW DOSING REGIMEN OF 80MG DAILY
- D-47 PREVENTION OF HEARTBURN SYMPTOMS WHEN ADMINISTERED FROM 15 MINUTES UP TO, BUT NOT INCLUDING, 1 HOUR PRIOR TO A PROVOCATIVE MEAL
- D-48 ADMINISTRATION OF CISATRACURIUM A NEUROMUSCULAR BLOCKING AGENT AT DOSES OF 3 AND 4X THE ED95 OF CISATRACURIUM FOLLOWING INDUCTION WITH THIOPENTAL
- D-49 PEDIATRIC DOSING GUIDELINES
- D-50 INFORMATION FOR USE OF CORVERT IN POST-CARDIAC SURGERY PATIENTS
- D-51 OPTIONAL STARTING DOSE OF 40MG/DAY
- D-52 ALTERNATE DOSING REGIMEN OF 1250MG TWICE DAILY
- D-53 USE IN PEDIATRIC PATIENTS FROM 1 MONTH TO 16 YEARS OF AGE
- D-54 USE OF ZYBAN FOR MAINTENANCE THERAPY. TREATMENT UP TO 6 MONTHS WAS SHOWN EFFICACIOUS
- D-55 ADDITION OF A HIGHER DOSE OF NUTROPIN FOR PUBERTAL PATIENTS (PUBERTAL DOSE LESS THAN OR EQUAL TO 0.7MG/KG/WEEK)
- D-56 ADDITION OF POSTPRANDIAL DOSING

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-57 3-HOUR INFUSION OF TAXOL GIVEN EVERY THREE WEEKS AT A DOSE OF 175MG/M2 FOLLOWED BY CISPLATIN AT A DOSE OF 75MG/M2 FOR THE FIRST-LINE TREATMENT OF ADVANCED OVARIAN CANCER
- D-58 CHANGE IN DOSING INTERVAL TO ONCE-DAILY ADMINISTRATION
- D-59 REDUCTION OF ELEVATED LDL-C IN A NEW, HIGHER STRENGTH TABLET, 0.8MG, AND FOR EXTENSION OF THE DOSAGE RANGE TO 0.8MG DAILY
- D-60 ADDITION OF A POST-OPERATIVE DOSING REGIMEN
- D-61 ONCE WEEKLY DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-62 ONCE WEEKLY DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-63 TO ALLOW A TITRATION DOSING REGIMEN USING A 25MG DOSE
- D-64 INCREASING DOSAGE FOR NERVE BLOCK ANESTHESIA USING NAROPIN 7.5MG/ML AND FOR EXTENDING THE DURATION OF TREATMENT FOR POSTOPERATIVE ANALGESIA USING NAROPIN 2MG/ML
- D-65 CHANGE DOSING AND ADMINISTRATION TO INDICATE MAINTENANCE OF WEIGHT LOSS OVER AN 18 MONTH PERIOD THUS EXTENDING THE USE OF THIS DRUG FROM ONE TO TWO YEARS
- D-66 DOSING RECOMMENDATIONS FOR PATIENTS UNDERGOING PCI
- D-67 SHORTER TREATMENT COURSE OF THREE DAYS IN THE TREATMENT OF RECURRENT EPISODES OF GENITAL HERPES
- D-68 CHANGE OF ADMIN RATE FOR INFUSION OF AREDIA FOR TREATMENT OF MODERATE AND SEVERE HYPERCALCEMIA OF MALIGNANCY FROM 24 HOURS TO 2 HOURS UP TO BUT NOT INCLUDING 24 HOURS
- D-69 SHORTENED DOSING REGIMEN TO 5 DAYS FOR THE TREATMENT OF ACUTE EXACERBATION OF CHRONIC BRONCHITIS
- D-70 80MG ONCE DAILY DOSING REGIMEN
- D-71 EIGHT WEEK DOSING REGIMEN
- D-72 INFORMATION REGARDING INCREASED RATE OF INFUSION FOR DEPACON
- D-73 ONCE A WEEK DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-74 ONCE A WEEK DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-75 INTERMITTENT DOSING REGIMEN, STARTING DAILY DOSE 14 DAYS PRIOR TO THE ANTICIPATED ONSET OF MENSTRUATION THROUGH THE FIRST FULL DAY OF MENSES AND REPEATING WITH EACH NEW CYCLE
- D-76 FOR USE ON AN "AS NEEDED" OR PRN BASIS FOR THE MANAGEMENT OF NASAL SYMPTOMS IN PATIENTS FOR WHOM THE DRUG IS INDICATED
- D-77 ADDITION OF 20MG AND 40MG DAILY AS OPTIONAL STARTING DOSES WITH 40MG INTENDED FOR PATIENTS WHO REQUIRE A LARGE REDUCTION IN LDL-C (MORE THAN 45%)
- D-78 USE OF FLEXERIL 5MG FOR THE RELIEF OF MUSCLE SPASM ASSOCIATED WITH ACUTE, PAINFUL, MUSCULOSKELETAL CONDITIONS
- D-79 NEW LOWER STARTING DOSE FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS AND/OR MODERATE TO SEVERE SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED W/ THE MENOPAUSE
- D-80 CHANGE OF DOSING SCHEDULE FOR LANTUS FROM ONCE DAILY AT BEDTIME TO FLEXIBLE DAILY DOSING
- D-81 NEW LOWER STARTING DOSE FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPORSIS
- D-82 USE OF PREMARIN 0.3 MG AND 0.45 MG FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-83 750 MG, ONCE DAILY FOR 5 DAYS FOR COMMUNITY ACQUIRED PNEUMONIA (CAP)
- D-84 ONCE-A-DAY DOSING OF FLOXACIN OTIC FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS (AGES 6 MO & OLDER) W/ OTITIS EXTERNA CAUSED BY SUSCEPTIBLE STRAINS OF E.COLI, P.AERUGINOSA AND S.AUREUS
- D-85 LOWER RECOMMENDED STARTING DOSE GUIDELINES FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE
- D-86 FOR USE IN SELECT EXTERNAL INSULIN PUMPS
- D-87 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-88 NEW DOSING RANGE OF 200-400MG PER DAY IN TWO DIVIDED DOSES FOR ADULTS WITH PARTIAL SEIZURES
- D-89 USE OF REYATAZ 300 MG/RITONAVIR 100 MG ONCE DAILY FOR TREATMENT IN HIV-INFECTED ANTIRETROVIRAL-EXPERIENCED PATIENTS
- D-90 ADDITION OF DAYTIME ADMINISTRATION TO TREAT VULVOVAGINAL CANDIDIASIS
- D-91 ALTERNATE INTERMITTENT DOSING REGIMEN
- D-92 ALTERNATIVE DOSAGE OF 1000MG ONCE DAILY AT BEDTIME
- D-93 ALTERNATE TWO OR THREE TIMES DAILY DOSING REGIMENS
- D-94 NEW MAXIMUM DOSAGE OF 72 MG/DAY IN ADOLESCENTS 13-17 YEARS OF AGE WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- D-95 BROADENED INITIAL STARTING DOSE FOR HYPERTENSION FROM 50 MG TO 100 MG TO 25 MG TO 100 MG DOSE RANGE
- D-96 ONCE-MONTHLY TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS WITH BONIVA (IBANDRONATE SODIUM) 150 MG TABLETS
- D-97 PED CANCER PT POPULATION EXPANDED TO INCLUDE PTS 6 MOS UP TO BUT NOT INCLUDING 4 YRS AND DOSING INSTRUCTIONS TO ADMIN 30 MIN BEFORE CHEMO WITH SECOND AND THIRD DOSES 4 & 8 HOURS AFTER FIRST DOSE
- D-98 DOSING FOR PED SURGICAL PTS EXPANDED TO INCLUDE PTS 1 MONTH UP TO BUT NOT INCLUDING 2 YEARS OF AGE
- D-99 ONCE DAILY ADMINISTRATION FOR THE TREATMENT OF HIV INFECTION IN THERAPY NAIVE ADULT PATIENTS
- D-100 750 MG ONCE DAILY FOR FIVE DAYS FOR THE TREATMENT OF ACUTE BACTERIAL SINUSITIS
- D-101 ONCE DAILY IN CHRONIC IDIOPATHIC URTICARIA FOR ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- D-102 NEW DOSING REGIMEN OF ONE SPRAY TWICE DAILY FOR SEASONAL ALLERGIC RHINITIS IN PATIENTS 12 YRS OF AGE AND OLDER
- D-103 NEW DOSING RECOMMENDATION FOR THE TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT PATIENTS, SPECIFICALLY A REDUCTION IN COURSE OF THERAPY FROM FAMCICLOVIR 125 MG TWICE-A-DAY FOR 5 DAYS TO 1000 MG TWICE-A-DAY FOR 1 DAY.
- D-104 0.5MG/0.1MG FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE IN WOMEN WHO HAVE A UTERUS
- D-105 USE OF ACTONEL 75MG TWO CONSECUTIVE DAYS PER MONTH FOR THE PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-106 FIVE DAY TREATMENT OF SELECTED SUSCEPTIBLE STRAINS OF STREPTOCOCCUS PNEUMONIAE, HAEMOPHILUS INFLUENZA, MYCOPLASMA PNEUMONIAE, AND CHLAMYDIA PNEUMONIAE FOR COMMUNITY-ACQUIRED PNEUMONIA
- D-107 PROVIDES FOR THE COMBINATION TABLET OF 70MG ALENDRONATE AND 5600 IU OF VITAMIN D3 FOR THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- D-108 TREATMENT OF COMPLICATED URINARY TRACT INFECTION AND ACUTE PYELONEPHRITIS WITH LEVAQUIN 750MG ONCE DAILY FOR FIVE DAYS
- D-109 PROVIDE FOR THE USE OF A LOWER DOSE FOR THE TREATMENT OF ADULTS WITH CHRONIC PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB MESYLATE
- D-110 TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGED 13-17
- D-111 PROVIDES FOR ONCE DAILY USE OF CIALIS, 2.5 MG AND 5 MG, FOR THE TREATMENT OF ERECTILE DYSFUNCTION
- D-112 PROVIDES FOR PEDIATRIC PUMP USE
- D-113 ONCE DAILY DOSING REGIMEN FOR PATIENTS WHO BECOME CONSTIPATED ON TWICE DAILY REGIMEN
- D-114 NEW DOSING RECOMMENDATIONS FOR USE OF SIROLIMUS IN COMBINATION WITH CYCLOSPORINE FOR THE PROPHYLAXIS OF REJECTION IN HIGH-RISK RENAL TRANSPLANT RECIPIENTS
- D-115 STARTING DOSE OF 15MG/DAY FOR MONOTHERAPY IN ACUTE TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED
- D-116 ALTERNATIVE DOSING REGIMEN ATAZANAVIR SULFATE CO-ADMINISTERED WITH RITONAVIR FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT NAIVE PATIENTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

D-117	50 MG TABLET FOR INITIATION OF DOSE TITRATION FOR BIPOLAR DISORDER
D-118	TWO 400MG TABLETS ONCE DAILY, CO-ADMINISTERED WITH 100MG RITONAVIR
D-119	DOSING RECOMMENDATIONS FOR HIV INFECTED PEDIATRIC PATIENTS 6 TO LESS THAN 18 YEARS OF AGE
D-120	DOSING REGIMEN ADJUSTMENTS
D-121	CHANGE TO REMOVE 20 MG MAXIMUM DOSAGE RESTRICTION
D-122	USE OF VAGIFEM 10 MCG FOR THE TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE
D-123	ALTERNATIVE DOSING REGIMEN DOSE OF 20 MG/METER SQUARE BY CONTINUOUS INTRAVENOUS INFUSION OVER 1 HOUR REPEATED DAILY FOR 5 DAYS
D-124	ONCE DAILY DOSING REGIMEN IN ADULT PATIENTS WITH LESS THAN THREE LOPINAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
D-125	EXTEND CURRENT DOSING REGIMEN TO 900MG (2-450MG TABLETS) ONCE A DAY WITHIN 10 DAYS OF TRANSPLANTATION UNTIL 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT PATIENTS AT HIGH RISK.
D-126	CHANGE DOSAGE REGIMEN FROM 250MG TO 500MG
D-127	DOSING REGIMEN FOR ADULT PATIENTS WITH CHRONIC HEPATITIS B (CHB) AND DECOMPENSATED LIVER DISEASE
D-128	SINGLE IV DOSE OF FOSAPREPITANT 150MG, DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID, FOR PREVENTION OF ACUTE & DELAYED NAUSEA & VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY EMETOGENIC CANCER CHEMO
D-129	800/100 MG DARUNAVIR/RITONAVIR, ONCE DAILY, IN TREATMENT-EXPERIENCED HIV-1 INFECTED PATIENTS WITH NO DARUNAVIR RESISTANCE ASSOCIATED SUBSTITUTIONS
D-130	DOSING RECOMMENDATIONS FOR TREATMENT OF HIV-1 INFECTION DURING PREGNANCY BASED ON DATA FROM STUDY AI424-182, A STUDY OF ATAZANAVIR/RITONAVIR IN COMBINATION WITH ZIDOVUDINE/LAMIVUDINE IN HIV INFECTED PREGNANT WOMEN
D-131	EVERY 6 TO 8 WEEKS FOR THE 120MG STRENGTH FOR PATIENTS WHO ARE CONTROLLED ON SOMATULINE DEPOT 60MG OR 90MG
D-132	45MG FOR 6 MONTH ADMINISTRATION
D-133	NEW EFFICACY DATA AND DOSING REGIMEN FOR PREGNANCY IN NORMAL OVULATORY WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION AS PART OF AN IVF OR INTRACYTOPLASMIC SPERM INJECTION (ICSI) CYCLE
D-134	INCREASING MAXIMUM DOSING OF PATIENTS WITH SCHIZOPHRENIA TO 160 MG/DAY
D-135	UPDATE LABELING WITH ONCE DAILY DOSING IN HIV-1 INFECTED, TREATMENT-NAIVE PEDIATRIC PATIENTS 12 TO LESS THAN 18 YEARS OF AGE
D-136	ALTERNATE DOSING REGIMEN FOR UNCOMPLICATED URETHRAL OR ENDOCERVICAL INFECTION CAUSED BY CHLAMYDIA TRACHOMATIS, ADMINISTER 200 MG BY MOUTH ONCE-A-DAY FOR 7 DAYS
D-137	NEW LOWER DOSING REGIMEN FOR REVATIO IN THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP 1) IN ADULTS
D-138	80 MG DOSING REGIMEN FOR THE RISK REDUCTION OF REBLEEDING OF GASTRIC AND DUODENAL ULCERS IN THE FIRST 72 HOURS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS
D-139	ADDITIONAL INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE LABELING REGARDING THE ADMINISTRATION OF BRAVELLE AND MENOPUR IN THE SAME SYRINGE TO OVULATORY WOMEN AS PART OF AN ART CYCLE
D-140	REVISED DOSING SCHEDULE TO ADMINISTER AVANAFIL 15 MINUTES PRIOR TO SEXUAL ACTIVITY
D-141	DOSING INFORMATION IN PREVIOUSLY UNTREATED MANTLE CELL LYMPHOMA
D-142	DOSE MODIFICATION GUIDELINES FOR BORTEZOMIB WHEN GIVEN IN COMBINATION WITH RITUXIMAB, CYCLOPHOSPHAMIDE, DOXORUBICIN, AND PREDNISONE
D-143	INITIATION OF VIMPAT THERAPY WITH A LOADING DOSE OF 200MG
D-144	LOWER LIMIT OF 15 MINUTES FOR THE INFUSION DURATION
D-145	UPDATES TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING TO REFLECT THE RESULTS OF TWO SHORT TERM STUDIES EVALUATING THE SAFETY AND EFFICACY OF INTUNIV IN CHILDREN AND ADOLESCENTS AGES 6 TO 17 WITH ADHD.

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

D-146	CHANGE IN TARGET DOSING TO 20MG TO 40MG ORALLY ONCE DAILY
D-147	ONCE DAILY DOSING IN PEDIATRIC PATIENTS 3 MONTHS OF AGE AND OLDER IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION
D-148	EXTENDED THE DURATION OF THE DOSING REGIMEN FROM 100 DAYS TO 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CMV DISEASE IN PEDIATRIC KIDNEY TRANSPLANT
D-149	DOSING INFORMATION ADDED TO THE LABELING REGARDING PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH ITP
D-150	1600MG DAILY FOR PATIENTS ON ADJUNCTIVE THERAPY WHO DID NOT ACHIEVE A SATISFACTORY RESPONSE ON 1200MG DAILY DOSE
D-151	DOSING RECOMMENDATIONS FOR THE TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS CO-INFECTED WITH HIV-1
D-152	DOSING RECOMMENDATIONS AS NECESSARY FOR FEVER AND PAIN FOR AGES 6MO TO LESS THAN 12 YEARS AND 12 TO 17 YEARS.
D-153	IN COMBINATION WITH RIBAVIRIN FOR 12 WEEKS, FOR THE TREATMENT OF GENOTYPE 1, CHRONIC HEPATITIS C TREATMENT EXPERIENCED PATIENTS WITH COMPENSATED CIRRHOSIS BASED UPON THE RESULTS OF THE SIRIUS STUDY
D-154	ADDITION OF A 1500MG-SINGLE-DOSE REGIMEN FOR THE TREATMENT OF ADULT PATIENTS WITH ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSI)
D-155	SINGLE IV DOSE OF FOSAPREPITANT 150MG DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID FOR PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
D-156	DOSING INFORMATION ADDED TO THE LABELING PROVIDING INFORMATION ON TRANSITIONING FROM SUBCUTANEOUS OR INTRAVENOUS ROUTES OF ADMINISTRATION OF TREPROSTINIL
D-157	UPDATED INFORMATION ADDED TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING PROVIDING DOSAGE RECOMMENDATIONS FOR INTERRUPTIONS AND DISCONTINUATION OF THERAPY
D-158	REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 1 HCV INFECTION
D-159	REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 4 HCV INFECTION
D-160	REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE PATIENTS WITH DECOMPENSATED CIRRHOSIS WITH GENOTYPE 1 HCV INFECTION
D-161	DOSAGE RECOMMENDATIONS ADDED TO INCLUDE TREATMENT OF HCV GENOTYPE 3 SUBJECTS CO-INFECTED WITH HIV-1
D-162	DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1 INFECTION WITH COMPENSATED (CHILD-PUGH A) OR DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS AND TREATMENT OF CHRONIC HCV GENOTYPE 3 INFECTION IN SUBJECTS WITH DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS
D-163	DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1A INFECTION WITH COMPENSATED (CHILD-PUGH A) CIRRHOSIS AND GENOTYPE 1B WITH OR WITHOUT COMPENSATED (CHILD-PUGH A) CIRRHOSIS
D-164	UPDATES TO THE DOSAGE AND ADMINISTRATION, DOSE MODIFICATIONS SECTION OF THE LABELING
D-165	DOSING RECOMMENDATION ADDED TO THE LABELING FOR IMBRUVICA USE IN COMBINATION WITH BENDAMUSTINE AND RITUXIMAB FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)/SMALL LYMPHOCYTIC LEUKEMIA (SLL)
D-166	BROADEN INITIAL STARTING DOSE FOR BIPOLAR I DISORDER TO 5-10MG TWICE DAILY
D-167	ADDITION OF 1200 MG ONCE DAILY DOSING FOR TREATMENT-NAIVE PATIENTS OR PATIENTS WHO ARE VIROLOGICALLY SUPPRESSED ON AN INITIAL REGIMEN OF RALTEGRAVIR FILM-COATED TABLETS 400 MG TWICE DAILY
D-168	NEW DOSING REGIMEN OF 10 MG ONCE DAILY FOR THE REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR DVT AND/OR PE AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
D-169	ONCE-DAILY DOSING FOR PATIENTS 5 YEARS OF AGE AND OLDER WHO HAVE UNDETECTABLE SERUM AND URINE SUCCINYLACETONE CONCENTRATIONS AFTER A MINIMUM OF 4 WEEKS ON A STABLE DOSAGE OF NITISINONE

PATENT AND EXCLUSIVITY TERMS

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EXCLUSIVITY DOSING SCHEDULE

D-170	TO ALLOW WITHDRAWAL THERAPY OF PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA IN CHRONIC PHASE WHO HAVE ACHIEVED A SUSTAINED MOLECULAR RESPONSE ON NILOTINIB THERAPY FOR A MINIMUM OF ONE YEAR PRIOR TO DISCONTINUATION
D-171	REVISED DOSING TO INCLUDE UP-TITRATION AS A STRATEGY TO IMPROVE TOLERABILITY AND THEREBY REDUCE TREATMENT DISCONTINUATION FOR ROFLUMILAST MAINTENANCE DOSAGE OF 500 MCG DAILY
D-172	ADDITION OF A ONCE WEEKLY DOSING REGIMEN FOR CARFILZOMIB IN COMBINATION WITH DEXAMETHASONE FOR PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
D-173	DOSING RECOMMENDATION FOR THE USE OF ELVITEGRAVIR/COBICISTAT/EMTRICITABINE/TENOFOVIR ALAFENAMIDE FIXED DOSE COMBINATION IN HIV-1 INFECTED ADULT PATIENTS WITH END-STAGE-RENAL DISEASE WHO ARE RECEIVING CHRONIC HEMODIALYSIS
D-174	MODIFICATIONS TO THE EXISTING DOSING REGIMEN TO ALLOW FOR TREATMENT INTERRUPTIONS OF UP TO 8 WEEKS FOR INTOLERABLE ADVERSE REACTIONS
D-175	EIGHT-WEEK DOSING REGIMEN FOR THE TREATMENT OF GENOTYPES 1, 2, 3, 4, 5, AND 6, CHRONIC HEPATITIS C VIRUS INFECTION IN TREATMENT-NAIVE SUBJECTS WITH COMPENSATED CIRRHOSIS BASED ON THE RESULTS FROM THE EXPEDITION-8 STUDY
D-176	IBRUTINIB IN COMBINATION WITH RITUXIMAB
D-177	INFORMATION ADDED TO THE DOSING SECTION IN REGARD TO THE TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION IN PATIENTS WITH SEVERE RENAL IMPAIRMENT INCLUDING PATIENTS WITH END STAGE RENAL DISEASE ON DIALYSIS
D-181	DOSING REGIMEN EXTENDING THE CONTRACEPTION USE FROM 5 YEARS TO UP TO 6 YEARS
D-182	NEW DOSING REGIMEN FOR THE PREVENTION AND MANAGEMENT OF NERATINIBASSOCIATED DIARRHEA
D-183	3-DAY DOSING REGIMEN FOR THE TREATMENT OF COVID-19 IN ADULTS AND PEDIATRIC PATIENTS (>12 YEARS AND WEIGHING AT LEAST 40 KG) WITH POSITIVE RESULTS OF DIRECT SARS-COV-2 VIRAL TESTING, WHO ARE NOT HOSPITALIZED AND HAVE MILD-TO-MODERATE COVID-19, AND ARE AT HIGH RISK FOR PROGRESSION TO SEVERE COVID-19, INCLUDING HOSPITALIZATION OR DEATH
D-184	NEW DOSING SCHEDULE FOR CABOTEGRAVIR/RILPIVRINE INJECTION EVERY 2 MONTHS
D-185	ADDITION OF A 3RD MAINTENANCE DOSE OF SEMAGLUTIDE
D-186	ADDITION OF A 3-DAY FOSAPREPITANT FOR INJECTION INTRAVENOUS DOSING REGIMEN IN PEDIATRIC PATIENTS FOR THE CURRENTLY APPROVED PREVENTION OF CHEMOTHERAPY-INDUCED NAUSEA AND VOMITING
D-187	ADDITION OF SECOND DOSE FOR TREATMENT OF PRESBYOPIA IN ADULTS
D-188	USE OF ALTERNATE INITIAL DOSING REGIMEN FOR INITIATION OF LACOSAMIDE TREATMENT IN PARTIAL ONSET SEIZURE PATIENTS ≥1 MONTH TO <17 YEARS OF AGE AND IN PRIMARY GENERALIZED TONIC-CLONIC SEIZURE PATIENTS ≥4 TO <17 YEARS
D-189	EXTENSION OF LETERMOVIR DOSING REGIMEN FROM 100 TO 200 DAYS POST-TRANSPLANT FOR THE PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE IN ADULT CMV SEROPOSITIVE RECIPIENTS (R+) OF AN ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANT WHO ARE AT RISK FOR LATE CMV INFECTION AND DISEASE
D-190	USE OF SEMAGLUTIDE 1.7 MG SUBCUTANEOUS WEEKLY AS AN ADDITIONAL MAINTENANCE DOSE
D-191	ADDITION OF A ONCE DAILY DOSING REGIMEN

EXCLUSIVITY INDICATION

I-1	DYSMENORRHEA
I-2	CHOLANGIOPANCREATOGRAPHY
I-3	INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
I-4	PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)
I-5	HYSTEOSALPINGOGRAPHY
I-6	TREATMENT OF JUVENILE ARTHRITIS
I-7	BIOPSY PROVEN MINIMAL CHANGE NEPHROTIC SYNDROME IN CHILDREN

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-8	ADULT INTRAVENOUS CONTRAST-ENHANCED COMPUTED TOMOGRAPHY OF THE HEAD AND BODY
I-9	PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
I-10	PREVENTION OF POSTOPERATIVE DEEP VENOUS THROMBOSIS AND PULMONARY EMBOLISM IN TOTAL HIP REPLACEMENT SURGERY
I-11	RELIEF OF MILD TO MODERATE PAIN
I-12	TREATMENT OF CUTANEOUS CANDIDIASIS
I-13	URINARY TRACT INFECTION (UTI) PREVENTION FOR PERIODS UP TO FIVE MONTHS IN WOMEN WITH A HISTORY OF RECURRENT UTI
I-14	SEBORRHEIC DERMATITIS
I-15	PHOTOPHERESIS IN THE PALLIATIVE TREATMENT OF SKIN MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PERSONS NOT RESPONSIVE TO OTHER TREATMENT
I-16	STIMULATE THE DEVELOPMENT OF MULTIPLE FOLLICLES/OOCYTES IN OVULATORY PATIENTS PARTICIPATING IN AN IN VITRO FERTILIZATION PROGRAM
I-17	MANAGEMENT OF CONGESTIVE HEART FAILURE
I-18	ENDOSCOPIC RETROGRADE PANCREATOGRAPHY
I-19	HERNIOGRAPHY
I-20	KNEE ARTHROGRAPHY
I-21	HIGH DOSE METHOTREXATE WITH LEUCOVORIN RESCUE IN COMBINATION WITH OTHER CHEMOTHERAPEUTIC AGENTS TO DELAY RECURRENCE IN PATIENTS WITH NONMETASTATIC OSTEOSARCOMA WHO HAVE UNDERGONE SURGICAL RESECTION OR AMPUTATION FOR THE PRIMARY TUMOR
I-22	RESCUE AFTER HIGH-DOSE METHOTREXATE THERAPY IN OSTEOSARCOMA
I-23	SHORT-TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
I-24	TREATMENT OF RHEUMATOID ARTHRITIS
I-25	ADULT INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY OF THE HEAD, NECK, ABDOMINAL, RENAL AND PERIPHERAL VESSELS
I-26	TREATMENT OF LIVER FLUKES
I-27	ADJUNCTIVE THERAPY TO DIET TO REDUCE THE RISK OF CORONARY ARTERY DISEASE
I-28	SELECTIVE ADULT VISCERAL ARTERIOGRAPHY
I-29	METASTATIC BREAST CANCER IN PREMENOPAUSAL WOMEN AS AN ALTERNATIVE TO OOPHORECTOMY OR OVARIAN IRRADIATION
I-30	TREATMENT OF TINEA PEDIS
I-31	CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS IN THE SPINE AND ASSOCIATED TISSUES
I-32	PEDIATRIC MYELOGRAPHY
I-33	ORAL USE OF DILUTED OMNIPAQUE INJECTION IN ADULTS FOR CONTRAST ENHANCED COMPUTED TOMOGRAPHY OF THE ABDOMEN
I-34	ORAL USE IN ADULTS FOR PASS-THROUGH EXAMINATION OF THE GASTROINTESTINAL TRACT
I-35	PEDIATRIC CONTRAST ENHANCEMENT OF COMPUTED TOMOGRAPHIC HEAD IMAGING
I-36	ARTHROGRAPHY OF THE SHOULDER JOINTS IN ADULTS
I-37	RADIOGRAPHY OF THE TEMPOROMANDIBULAR JOINT IN ADULTS
I-38	CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS OF THE CENTRAL NERVOUS SYSTEM IN CHILDREN (2 YEARS OF AGE AND OLDER)
I-39	TREATMENT OF ACUTE MYOCARDIAL INFARCTION
I-40	PRIMARY NOCTURNAL ENURESIS
I-41	MIGRAINE HEADACHE PROPHYLAXIS
I-42	HERPES ZOSTER
I-43	HERPES SIMPLEX ENCEPHALITIS
I-44	MAINTENANCE THERAPY IN HEALED DUODENAL ULCER PATIENTS AT DOSE OF 1 GRAM TWICE DAILY
I-45	ACUTE TREATMENT OF VARICELLA ZOSTER VIRUS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-46 USE IN PEDIATRIC COMPUTED TOMOGRAPHIC HEAD AND BODY IMAGING

I-47 TREATMENT OF PEDIATRIC PATIENTS WITH SYMPTOMATIC HUMAN IMMUNODEFICIENCY VIRUS (HIV) DISEASE

I-48 PEDIATRIC ANGIOCARDIOGRAPHY

I-49 TREATMENT OF TRAVELERS' DIARRHEA DUE TO SUSCEPTIBLE STRAINS OF ENTEROTOXIGENIC ESCHERICHIA COLI

I-50 FOR USE IN WOMEN WITH AXILLARY NODE-NEGATIVE BREAST CANCER

I-51 TREATMENT OF PRIMARY DYSMENORRHEA AND FOR THE TREATMENT OF IDIOPATHIC HEAVY MENSTRUAL BLOOD LOSS

I-52 PEDIATRIC EXCRETORY UROGRAPHY

I-53 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA

I-54 RENAL CONCENTRATION CAPACITY TEST

I-55 HYPERTENSION

I-56 EROSION GASTROESOPHAGEAL REFLUX DISEASE

I-57 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER

I-58 INITIAL TREATMENT OF ADVANCED OVARIAN CARCINOMA IN COMBINATION WITH OTHER APPROVED CHEMOTHERAPEUTIC AGENTS

I-59 ENDOSCOPICALLY DIAGNOSED ESOPHAGITIS, INCLUDING EROSION AND ULCERATIVE ESOPHAGITIS, AND ASSOCIATED HEARTBURN DUE TO GASTROESOPHAGEAL REFLUX DISEASE

I-60 SINGLE APPLICATION TREATMENT OF HEAD LICE IN CHILDREN TWO MONTHS TO TWO YEARS IN AGE

I-61 FEMALE ANDROGENETIC ALOPECIA

I-62 PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS

I-63 ONCE DAILY TREATMENT AS INITIAL THERAPY IN THE TREATMENT OF HYPERTENSION

I-64 PREVENTION OF SUPRAVENTRICULAR TACHYCARDIAS

I-65 PREVENTION OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS

I-66 UNCOMPLICATED GONORRHEA

I-67 TREATMENT OF ACUTE ASTHMATIC ATTACKS IN CHILDREN SIX YEARS OF AGE AND OLDER

I-68 CENTRAL PRECOCIOUS PUBERTY

I-69 SHORT TERM TREATMENT OF PATIENTS WITH SYMPTOMS OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), AND FOR THE SHORT TERM TREATMENT OF ESOPHAGITIS DUE TO GERD INCLUDING ULCERATIVE DISEASE DIAGNOSED BY ENDOSCOPY

I-70 USE IN COMBINATION WITH 5-FLUOROURACIL TO PROLONG SURVIVAL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED COLORECTAL CANCER

I-71 VARICELLA INFECTIONS (CHICKENPOX)

I-72 PREVENTION OF CMV DISEASE IN TRANSPLANT PATIENTS AT RISK FOR CMV DISEASE

I-73 INITIATE AND MAINTAIN MONITORED ANESTHESIA CARE (MAC) SEDATION DURING DIAGNOSTIC PROCEDURES

I-74 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY

I-75 TREATMENT OF ENDOSCOPICALLY DIAGNOSED EROSION ESOPHAGITIS

I-76 PREVENTION OF OSTEOPOROSIS

I-77 DERMAL INFECTIONS-TINEA PEDIS, TINEA CORPORIS, TINEA CRURIS DUE TO EPIDERMOPHYTON FLOCCOSUM

I-78 CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY AND INTRAVENOUS EXCRETORY UROGRAPHY

I-79 MANAGEMENT OF CHRONIC STABLE ANGINA AND ANGINA DUE TO CORONARY ARTERY SPASM

I-80 DIAGNOSIS AND LOCALIZATION OF ISCHEMIA AND CORONARY HEART DISEASE

I-81 PROPHYLAXIS IN DESIGNATED IMMUNOCOMPROMISED CONDITIONS TO REDUCE THE INCIDENCE OF OROPHARYNGEAL CANDIDIASIS

I-82 TREATMENT OF TRAVELERS' DIARRHEA

I-83 ANGIOCARDIOGRAPHY, CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY IN CHILDREN

I-84 INTRAOPERATIVE AND POSTOPERATIVE TACHYCARDIA AND/OR HYPERTENSION

I-85 TREATMENT OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS

I-86 TREATMENT OF SECONDARY CARNITINE DEFICIENCY

I-87 RENAL IMAGING AGENT FOR USE IN CHILDREN

I-88 MANAGEMENT OF ENDOMETRIOSIS

I-89 EPIDURAL USE IN LABOR AND DELIVERY AS AN ANALGESIC ADJUNCT TO BUPIVACAINE

I-90 INTENSIVE CARE UNIT SEDATION

I-91 MONOTHERAPY USE FOR HYPERTENSION

I-92 ADJUNCTIVE THERAPY IN THE MANAGEMENT OF HEART FAILURE

I-93 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN CHILDREN AGES 4-11 YEARS

I-94 USE WITH MRI IN ADULTS TO PROVIDE CONTRAST ENHANCEMENT AND FACILITATE VISUALIZATION OF LESIONS IN THE BODY [EXCLUDING THE HEART]

I-95 TREATMENT OF LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION

I-96 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA

I-97 ORAL OR RECTAL USE IN CHILDREN FOR THE EXAMINATION OF THE GASTROINTESTINAL TRACT

I-98 TREATMENT OF CHILDREN WHO HAVE GROWTH FAILURE ASSOCIATED WITH CHRONIC RENAL INSUFFICIENCY

I-99 PEDIATRIC ANESTHESIA IN CHILDREN 3 YEARS AND OLDER

I-100 TO DECREASE THE INCIDENCE OF CANDIDIASIS IN PATIENTS UNDERGOING BONE MARROW TRANSPLANTATION WHO RECEIVE CYTOTOXIC CHEMOTHERAPY AND/OR RADIATION THERAPY

I-101 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN-DEPENDENT DIABETES MELLITUS AND RETINOPATHY

I-102 TREATMENT OF OBSESSIVE-COMPULSIVE DISORDER

I-103 PROPHYLAXIS AGAINST PNEUMOCYSTIS CARINII PNEUMONIA IN INDIVIDUALS WHO ARE IMMUNOCOMPROMISED AND CONSIDERED TO BE AT RISK OF DEVELOPING PNEUMOCYSTIS CARINII PNEUMONIA

I-104 TREATMENT OF PULMONARY AND EXTRAPULMONARY ASPERGILLOSIS IN PATIENTS WHO ARE INTOLERANT OF OR WHO ARE REFRACTORY TO AMPHOTERICIN B THERAPY

I-105 TREATMENT OF METASTATIC CARCINOMA OF THE BREAST AFTER FAILURE OF FIRST-LINE OR SUBSEQUENT CHEMOTHERAPY

I-106 TREATMENT OF ACROMEGALY

I-107 VAGINAL CANDIDIASIS

I-108 EXPANDED USE-FOR ICU PATIENTS UNDERGOING LONG-TERM INFUSION DURING MECHANICAL VENTILATION

I-109 TYPHOID FEVER

I-110 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIOTHERAPY

I-111 TREATMENT OF PAGET'S DISEASE OF BONE

I-112 MANAGEMENT OF MODERATE TO SEVERE PAIN

I-113 TREATMENT OF PROSTATITIS

I-114 USE IN CHILDREN TO VISUALIZE LESIONS WITH ABNORMAL VASCULARITY IN THE BRAIN (INTRACRANIAL LESIONS), SPINE, AND ASSOCIATED TISSUE

I-115 USE IN MRI IN ADULTS TO VISUALIZE LESIONS IN THE HEAD AND NECK

I-116 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS

I-117 TO SLOW THE PROGRESSION FO CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE

I-118 PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM FOLLOWING KNEE REPLACEMENT SURGERY

I-119 TREATMENT OF ANEMIA CAUSED BY UTERINE LEIOMYOMATA IN WOMEN WHO FAIL IRON THERAPY

I-120 MAINTENANCE THERAPY FOR GASTRIC ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING ACUTE ULCERS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-121	EXPANDED PATIENT POPULATION -- USE IN ICU PATIENTS
I-122	PSORIASIS OF THE SCALP
I-123	RELIEF OF MILD TO MODERATE PAIN IN PATIENTS AGED 6 MONTHS AND OLDER
I-124	LEUKOCYTE LABELED SCINTIGRAPHY AS AN ADJUNCT IN THE LOCALIZATION OF INTRA-ABDOMINAL INFECTION AND INFLAMMATORY BOWEL DISEASE
I-125	EXPANSION OF CONSCIOUS SEDATION INDICATION TO INCLUDE SHORT THERAPEUTIC PROCEDURES
I-126	ADJUNCT TO THALLIUM- 201 MYOCARDIAL PERFUSION IN PATIENTS UNABLE TO EXERCISE ADEQUATELY
I-127	TREATMENT OF ACYCLOVIR-RESISTANT HERPES IN IMMUNOCOMPROMISED PATIENTS
I-128	IN PT W/ CH DISEASE AND HYPERCHOLESTEROLEMIA; REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH; REDUCE RISK NON-FATAL MI; REDUCE RISK UNDERGOING MYOCARDIAL REVASCLARIZATION PROCEDURES; REDUCTION ELEVATED TOTAL AND LDL CHOL LEVELS...
I-129	TREATMENT OF ALCOHOL DEPENDENCE
I-130	MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS
I-131	PERIPHERAL ARTERIOGRAPHY
I-132	TREATMENT OF MANIC PHASE OF BIPOLAR DISORDER
I-133	MANAGEMENT OF CHRONIC STABLE ANGINA
I-134	HEART FAILURE POST MYOCARDIAL INFARCTION
I-135	BONE METASTASES ASSOCIATED WITH MULTIPLE MYELOMA
I-136	IDIOPATHIC CHRONIC URTICARIA
I-137	PREVENTION OF METAL-INDUCED HEART BURN, ACID INDIGESTION, AND SOUR STOMACH WHEN TAKEN 30 MINUTES PRIOR TO CONSUMING FOOD OR BEVERAGES
I-138	TREATMENT OF ACUTE RECURRENT GENITAL HERPES
I-139	PALLIATIVE TREATMENT OF ADVANCED BREAST CANCER IN PRE- AND PERIMENOPAUSAL WOMEN
I-140	PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN INDIVIDUALS WITH HIV INFECTION AT RISK FOR DEVELOPING CMV DISEASE
I-141	TREATMENT OF HEMODYNAMICALLY STABLE PATIENTS WITHIN 24 HOURS OF ACUTE MYOCARDIAL INFARCTION TO IMPROVE SURVIVAL
I-142	LOCALIZE MYOCARDIAL ISCHEMIA(REVERSIBLE DEFECT) AND INFARCTION (NON-REVERSIBLE DEFECTS) IN EVALUATING MYOCARDIAL FUNCTION
I-143	EPISODIC TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
I-144	ENHANCEMENT OF MRI OF THE ADULT BODY INTERNAL ORGANS
I-145	0.1MMOL/KG AS A SINGLE INTRAVENOUS BOLUS FOR MRI OF THE CNS IN CHILDREN
I-146	CONTRAST ENHANCEMENT AND FACILITATION OF VISUALIZATION OF EXTRACRANIAL HEAD AND NECK LESIONS
I-147	PREVENTION OF GALLSTONE FORMATION IN OBESE PATIENTS EXPERIENCING RAPID WEIGHT LOSS
I-148	TREATMENT OF ACUTE PNEUMOCYSTIS CARINI PNEUMONIA (PCP) IN HIV-INFECTED PATIENTS WHOSE ALVEOLAR-ARTERIAL OXYGEN DIFFERENCE (AaDO ₂) IS LESS THAN OR EQUAL TO 55 TORR
I-149	TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER
I-150	TREATMENT OF OBSESSIVE COMPULSIVE DISORDER AND PANIC DISORDER
I-151	PREVENTION OF AND PREVENTION OF FURTHER POSTOPERATIVE NAUSEA AND VOMITING IN PEDIATRIC PATIENTS RECEIVING GENERAL ANESTHESIA
I-152	SLOWING THE PROGRESSION OF CORONARY ATHEROSCLEROSIS AND REDUCING THE RISK OF ACUTE CORONARY EVENTS
I-153	MANAGEMENT OF SEVERE SPASTICITY [ENCOMPASSES SPINAL AND CEREBRAL ORIGIN]
I-154	PATIENT POPULATION ALTERED TO INCLUDE PEDIATRIC USE
I-155	TREATMENT OF ONYCHOMYCOSIS DUE TO DERMATOPHYTES (TINEA UNGUIUM) OF THE TOENAIL WITH OR WITHOUT FINGERNAIL INVOLVEMENT

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-156	ADDITIONAL DATA REGARDING THE SAFE USE OF NORVASC IN PATIENTS WITH HEART FAILURE
I-157	TREATMENT OF ACUTE UNCOMPLICATED CYSTITIS IN FEMALES
I-158	TREATMENT OF OSTEOLYTIC BONE METASTASES OF BREAST CANCER
I-159	FOR HYPERCHOLESTEROLEMIC PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE REDUCE THE RISK OF MYOCARDIAL INFARCTION, REVASCULARIZATION, AND DEATH DUE TO CARDIOVASCULAR CAUSES WITH NO INCREASE IN DEATH FROM NON-CARDIOVASCULAR CAUSES
I-160	TREATMENT OF BACTERIAL CORNEAL ULCERS
I-161	TREATMENT OF ADULT-ONSET OR CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENCY
I-162	FOR USE IN PATIENTS 6-11 YEARS OF AGE
I-163	TREATMENT OF PHOTOPHOBIA
I-164	CHRONIC BACTERIAL PROSTATITIS
I-165	MANAGEMENT OF ADULTS WITH ACTIVE, CLASSIC AND DEFINITIVE RHEUMATOID ARTHRITIS WHO HAVE HAD INSUFFICIENT THERAPEUTIC RESPONSE TO OR ARE INTOLERANT OF AN ADEQUATE TRIAL OF FULL DOSES OF ONE OR MORE NON-STEROIDAL ANTI-INFLAMMATORY DRUGS
I-166	TREATMENT OF BULIMIA
I-167	COMPLICATED INTRA-ABDOMINAL INFECTIONS (USED IN COMBINATION WITH METRONIDAZOLE) CAUSED BY MIXED AEROBIC/ANAEROBIC PATHOGENS
I-168	MANAGEMENT OF LOCALLY CONFINED STAGE B2-C METASTATIC CARCINOMA OF THE PROSTATE (IN COMBINATION WITH LHRH AGONISTS)
I-169	USE IN COMBINATION WITH CORTICOSTEROIDS AS INITIAL CHEMOTHERAPY FOR THE TREATMENT OF PATIENTS WITH PAIN RELATED TO ADVANCED HORMONE-REFRACTORY PROSTATE CANCER
I-170	PROPHYLACTIC USE DURING HEAD LICE EPIDEMICS
I-171	RELIEF OF SYMPTOMS OF THE COMMON COLD
I-172	TREATMENT OF INITIAL EPISODE OF GENITAL HERPES
I-173	PREOPERATIVELY FOR THE PREVENTION OF INFECTION IN TRANSRECTAL PROSTATE BIOPSY
I-174	PELVIC INFLAMMATORY DISEASE
I-175	TREATMENT OF TINEA CORPORIS AND TINEA CRURIS
I-176	TREATMENT OF POSTOPERATIVE INFLAMMATION IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION
I-177	TX OF MODERATE ACNE VULGARIS IN FEMALES, GREATER OR EQUAL TO 15YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, DESIRE CONTRACEPTION, HAVE ACHIEVED MENARCHE AND ARE UNRESPONSIVE TO TOPICAL ANTI-ACNE MEDICATIONS
I-178	TREATMENT OF ONYCHOMYCOSIS OF THE FINGERNAIL WITHOUT CONCOMITANT ONCHOMYCOSIS OF THE TOENAIL WITH A PULSE DOSING REGIMEN
I-179	NOSOCOMIAL PNEUMONIA-MILD TO MODERATE AND SEVERE CAUSED BY HAEMOPHILUS INFLUENZAE OR KLEBSIELLA PNEUMONIAE
I-180	TREATMENT OF PLANTAR TINEA PEDIS (MOCCASIN TYPE)
I-181	TREATMENT OF PATIENTS WITH COMPLEX PARTIAL SEIZURES WITH AND WITHOUT SECONDARY GENERALIZATION
I-182	TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME
I-183	MAINTENANCE THERAPY IN THE MANAGEMENT OF MILD TO MODERATE ASTHMA IN PEDIATRIC PATIENTS AGES 6-11
I-184	TREATMENT OF PANIC DISORDER AT A RECOMMENDED DOSE RANGE OF 1 TO 2MG/DAY (MAXIMUM OF 4MG)
I-185	PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
I-186	TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR CAUSED BY OR PRESUMED TO BE CAUSED BY PITYROSPORUM ORBICULARE (ALSO KNOWN AS MALASSEZIA FURFUR OR M. ORBICULARE)
I-187	PREVENTION OF FRACTURES IN THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
I-188	TREATMENT OF ACUTE SINUSITIS AND ACUTE EXACERBATION OF CHRONIC SINUSITIS
I-189	TREATMENT OF ACUTE OTITIS MEDIA IN PEDIATRIC PATIENTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-190	PLANAR IMAGING AS A SECOND LINE DIAGNOSTIC DRUG AFTER MAMMOGRAPHY TO ASSIST IN THE EVALUATION OF BREAST LESIONS IN PATIENTS WITH AN ABNORMAL MAMMOGRAM OR A PALPABLE BREAST MASS
I-191	ENDOMETRIAL THINNING AGENT PRIOR TO ENDOMETRIAL ABLATION FOR DYSFUNCTIONAL UTERINE BLEEDING
I-192	THE PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS AND A NEW DOSAGE REGIMEN, 40MG ONCE DAILY, FOR THIS INDICATION
I-193	TREATMENT OF PANIC DISORDER IN A RECOMMENDED DOSE RANGE OF 50 TO 200MG/DAY
I-194	CONGESTIVE HEART FAILURE
I-195	FOR USE OF LANSOPRAZOLE IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF DUODENAL ULCER
I-196	ACUTE TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
I-197	MAINTENANCE OF HEALING OF DUODENAL ULCER
I-198	FOR THE USE OF LANSOPRAZOLE IN COMBINATION WITH AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF A DUODENAL ULCER
I-199	MONOTHERAPY AND COMBINATION THERAPY WITH SULFONYLUREA IN THE TREATMENT OF TYPE II DIABETES
I-200	TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR
I-201	EMPIRICAL THERAPY FOR FEBRILE NEUTROPENIC PATIENTS
I-202	SECOND-LINE TREATMENT OF AIDS-RELATED KAPOSI'S SARCOMA
I-203	MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
I-204	USE IN PEDIATRIC PATIENTS BETWEEN THE AGES OF 6 AND 11 FOR THE TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
I-205	INITIAL ANTICONVULSANT TREATMENT OF STATUS EPILEPTICUS
I-206	TREATMENT OF EDEMA ASSOCIATED WITH CHRONIC RENAL FAILURE
I-207	FOR THE SUPPRESSION OF RECURRENT EPISODES OF GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
I-208	TREATMENT OF OBSESSIVE COMPULSIVE DISORDER IN THE PEDIATRIC POPULATION
I-209	PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA (PSVT)
I-210	TO SLOW THE PROGRESSION OF CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL AND LDL CHOLESTEROL TO TARGET LEVELS
I-211	FOR USE IN PEDIATRIC POPULATION
I-212	TREATMENT OF SYMPTOMS OF DRY MOUTH IN PATIENTS WITH SJOGREN'S SYNDROME
I-213	TEMPORARY RELIEF OF PAIN AND PHOTOPHOBIA IN PATIENTS UNDERGOING CORNEAL REFRACTIVE SURGERY
I-214	TREATMENT OF OSTEOPOROSIS
I-215	PRE-PROCEDURAL APPLICATION TO ADULT MALE GENITAL SKIN PRIOR TO SITE-SPECIFIC SUBCUTANEOUS INFILTRATION WITH LIDOCAINE FOR THE REMOVAL OF GENITAL WARTS
I-216	FOR THE LONG-TERM TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
I-217	PREVENTION (DURING AND FOLLOWING HOSPITALIZATION) OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
I-218	USE OF LIPITOR AS AN ADJUNCTIVE THERAPY TO DIET FOR THE TREATMENT OF PATIENTS WITH ELEVATED SERUM TRIGLYCERIDE LEVELS (FREDERICKSON TYPE IV)
I-219	USE OF LIPITOR BY PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDERICKSON TYPE III) WHO DO NOT RESPOND ADEQUATELY TO DIET
I-220	TREATMENT OF EPISODIC- HEARTBURN, ACID INDIGESTION AND SOUR STOMACH
I-221	TREATMENT OF BENIGN PROSTATIC HYPERPLASIA (BPH) IN MEN WITH AN ENLARGED PROSTATE TO IMPROVE SYMPTOMS, REDUCE THE RISK OF ACUTE URINARY RETENTION AND REDUCE THE RISK OF THE NEED OF SURGERY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-222 PREVENTION OF ISCHEMIC COMPLICATIONS OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION, WHEN CONCURRENTLY ADMINISTERED WITH ASPIRIN
- I-223 USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH ALLERGIC AND NONALLERGIC-PERENNIAL RHINITIS IN CHILDREN AGE 6-11 YEARS
- I-224 FOR THE USE IN PEDIATRIC PATIENTS 4 TO 11 YEARS OF AGE FOR THE MANAGEMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- I-225 USE IN PATIENTS WITH PREVIOUS MI AND NORMAL CHOLESTEROL LEVELS, TO REDUCE RISK OF RECURRENT MI, MYOCARDIAL REVASCULARIZATION, AND CEREBROVASCULAR DISEASE EVENTS
- I-226 FIRST-LINE THERAPY FOR THE TREATMENT OF ADVANCED CARCINOMA OF THE OVARY IN COMBINATION WITH CISPLATIN
- I-227 SHORT-TERM TREATMENT OF SYMPTOMATIC GASTROESOPHAGEAL REFLUX DISEASE (GERD)
- I-228 PREVENTION OF MEAL INDUCED HEARTBURN AT A DOSE OF 75MG TAKEN 30-60MIN PRIOR TO A MEAL
- I-229 PRILOSEC (OMEPRAZOLE), AMOXICILLIN, AND CLARITHROMYCIN FOR THE ERADICATION OF H. PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- I-230 IN COMBINATION WITH CIS-PLATIN, FOR THE FIRST LINE TREATMENT OF NON-SMALL CELL LUNG CANCER IN PATIENTS WHO ARE NOT CANDIDATES FOR POTENTIALLY CURATIVE SURGERY AND/OR RADIATION
- I-231 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY
- I-232 TREATMENT OF RECURRENT MUCOCUTANEOUS HERPES SIMPLEX INFECTIONS IN HIV-AFFECTED PATIENTS AT A DOSE OF 500MG TWICE DAILY
- I-233 PROPHYLACTIC USE TO REDUCE PERIOPERATIVE BLOOD LOSS AND THE NEED FOR BLOOD TRANSFUSION IN PATIENTS UNDERGOING CARDIOPULMONARY BYPASS IN THE COURSE OF CORONARY ARTERY BYPASS GRAFT SURGERY
- I-234 FOR USE IN COMBINATION WITH CISPLATIN FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED (STAGE IIIA OR IIIB) OR METASTATIC (STAGE IV) NON-SMALL CELL LUNG CANCER
- I-235 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 12 YEARS OF AGE AND OLDER
- I-236 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-237 MAINTENANCE TREATMENT OF ASTHMA AND PREVENTION OF BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-238 ADJUNCTIVE TREATMENT OF LENNOX-GASTAUT SYNDROME IN PEDIATRIC AND ADULT PATIENTS
- I-239 TREATMENT OF PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- I-240 MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM AND RESULTANT METABOLIC BONE DISEASE IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL FAILURE (CCR 15 TO 55ML/MIN) NOT YET ON DIALYSIS
- I-241 USE IN PHOTODYNAMIC THERAPY (PDT) FOR REDUCTION OF OBSTRUCTION AND PALLIATION OF SYMPTOMS IN PATIENTS WITH COMPLETELY OR PARTIALLY OBSTRUCTING ENDOBRONCHIAL NONSMALL CELL LUNG CANCER
- I-242 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE AND IN THE TREATMENT OF VULVAR AND VAGINAL ATROPHY IN WOMEN WITH AN INTACT UTERUS
- I-243 USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH THE COMMON COLD IN CHILDREN AGE 5 TO 11 YEARS
- I-244 REDUCE THE INCIDENCE OF BREAST CANCER IN WOMEN AT HIGH RISK FOR BREAST CANCER
- I-245 TREATMENT OF ACUTE SINUSITIS
- I-246 TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS
- I-247 USE IN CONVERSION TO MONOTHERAPY IN ADULTS WITH PARTIAL SEIZURES WHO ARE RECEIVING TREATMENT WITH A SINGLE ENZYME-INDUCING ANTIEPILEPTIC DRUG
- I-248 INPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITH/WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM AND OUTPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM
- I-249 TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS WITH COMPENSATED LIVER DISEASE PREVIOUSLY UNTREATED WITH ALPHA INTERFERON THERAPY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-250	PRIMARY PREVENTION OF CORONARY HEART DISEASE IN PATIENTS WITHOUT SYMPATOMATIC CARDIOVASCULAR DISEASE WHO HAVE AVERAGE TO MODERATELY ELEVATED TOTAL-C AND LDL-C AND BELOW AVERAGE HDL-C
I-251	TREATMENT OF GENERALIZED ANXIETY DISORDER
I-252	NEW COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS METFORMIN
I-253	COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS INSULIN
I-254	PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS (LOSS OF BONE MASS)
I-255	PREVENTION OF PNEUMOCYSTIS CARINII PNEUMONIA (PCP)
I-256	USE IN TREATMENT OF SMALL CELL LUNG CANCER SENSITIVE DISEASE AFTER FAILURE OF FIRST-LINE CHEMOTHERAPY
I-257	TREATMENT OF CHRONIC HEPATITIS B ASSOCIATED WITH EVIDENCE OF HEPATITIS B VIRAL REPLICATION AND ACTIVE LIVER INFLAMMATION
I-258	FOR PERENNIAL NONALLERGIC RHINITIS FOR AGES 4 AND ABOVE
I-259	PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
I-260	EXPANDED PEDIATRIC USE IN CHILDREN YOUNGER THAN ONE MONTH OF AGE TO BIRTH (WITH A GESTATIONAL AGE OF 37 WEEKS OR GREATER)
I-261	TREATMENT OF SOCIAL ANXIETY DISORDER
I-262	TREATMENT OR PREVENTION OF BRONCHOSPASM WITH REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE AND FOR THE PREVENTION OF EXERCISE INDUCED BRONCHOSPASM IN CHILDREN AGES 4-12
I-263	TREATMENT OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION FOR THE PREVENTION OF ISCHEMIC COMPLICATIONS IN PATIENTS ON CONCURRENT ASPIRIN THERAPY
I-264	PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIATION, INCLUDING TOTAL BODY IRRADIATION (TBI) AND FRACTIONATED ABDOMINAL RADIATION
I-265	TREATMENT OF ATOPIC DERMATITIS IN PEDIATRIC PATIENTS 6 YEARS AND OLDER
I-266	USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN PEDIATRIC PATIENTS AGES 2-16 YEARS WITH PARTIAL ONSET SEIZURES
I-267	USE IN PEDIATRIC PATIENTS 3 MONTHS OLD AND OLDER - FOR CORTICOSTEROID-RESPONSIVE DERMATOSES
I-268	PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 7-11 YEARS OF AGE
I-269	PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HIGHLY EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING CISPLATIN
I-270	ADJUVANT TREATMENT OF NODE-POSITIVE BREAST CANCER ADMINSTRERED SEQUENTIALLY TO STANDARD DOXORUBICIN-CONTAINING COMBINATION CHEMOTHERAPY
I-271	TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
I-272	TREATMENT OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN MEN AND WOMEN RECEIVING GLUCOCORTICOID IN A DAILY DOSE EQUIVALENT TO 7.5MG OR GREATER OF PREDNISONE AND WHO HAVE LOW BONE MINERAL DENSITY
I-273	ADJUNCT TO DIET TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NON FAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
I-274	USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
I-275	USE IN COMBINATION WITH METFORMIN AND SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES
I-276	USE OF REZULIN IN COMBINATION WITH METFORMIN AND SULFONYLUREAS IN PATIENTS WITH TYPE 2 DIABETES
I-277	TREATMENT OF TYPE III HYPERLIPOPROTEINEMIA
I-278	TREATMENT OF PATIENTS WITH ISOLATED HYPERTRIGLYCERIDEMIA (FREDERICKSON TYPE IV)
I-279	TREATMENT OF POST-TRAUMATIC STRESS DISORDER
I-280	USE OF CARNITOR INJECTION FOR THE PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-281	INCREASING HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NONFAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
I-282	TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
I-283	TO REDUCE THE INCIDENCE OF MODERATE TO SEVERE XEROSTOMIA IN PATIENTS UNDERGOING POST-OPERATIVE RADIATION TREATMENT FOR HEAD AND NECK CANCER, WHERE THE RADIATION PORT INCLUDES A SUBSTANTIAL PORTION OF THE PAROTID GLANDS
I-284	TO REDUCE THE NUMBER OF ADENOMATOUS COLORECTAL POLYPS IN FAMILIAL ADENOMATOUS POLYPOSIS PATIENTS AS AN ADJUNCT TO USUAL CARE
I-285	TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN ADULTS AND CHILDREN 3 YEARS OF AGE AND OLDER
I-286	TREATMENT OF PATIENTS WITH FREDERICKSON TYPE III
I-287	USE OF PRAVASTATIN IN PATIENTS WITH EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF TOTAL MORTALITY BY REDUCING CORONARY DEATH
I-288	CHANGES IN SEVERAL SECTIONS OF THE INSERT TO INCORPORATE STATEMENTS CONCERNING THE USE OF HIGH DOSES OF LISINAPRIL TO REDUCE THE RISK OF THE COMBINED OUTCOMES OF MORTALITY AND HOSPITALIZATION IN PATIENTS WITH CONGESTIVE HEART FAILURE
I-289	USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN DIET AND EXERCISE WITH EITHER SINGLE AGENT DOES NOT ACHIEVE ADEQUATE GLYCEMIC CONTROL
I-290	PREVENTION OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
I-291	PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
I-292	TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
I-293	TREATMENT OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
I-294	TREATMENT OF UNCOMPLICATED ACUTE ILLNESS DUE TO INFLUENZA A AND B IN PEDIATRIC PATIENTS 7 YEARS AND OLDER WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 2 DAYS
I-295	PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR WOMEN WITH AN INTACT UTERUS
I-296	LONG-TERM INTRAVENOUS TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH THE SCLERODERMA SPECTRUM OF DISEASE IN NYHA CLASS III AND CLASS IV PATIENTS WHO DO NOT RESPOND TO CONVENTIONAL THERAPY
I-297	SHORT-TERM TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
I-298	TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IIA AND IIB HYPERLIPOPROTEINEMIA
I-299	USE OF CAMPTOSAR AS A COMPONENT OF FIRST-LINE THERAPY IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR PATIENTS WITH METASTATIC CARCINOMA OF THE COLON OR RECTUM
I-300	PROPHYLAXIS FOR ASTHMA IN CHILDREN 2-5 YEARS OF AGE
I-301	TREATMENT OF SIGNS AND SYMPTOMS OF ALLERGIC CONJUNCTIVITIS
I-302	TREATMENT OF PEDIATRIC PATIENTS WITH PRADER-WILLI SYNDROME
I-303	INCREASING HDL-CHOLESTEROL IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIAS
I-304	TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IV
I-305	TREATMENT OF LEVOFLOXACIN SUSCEPTIBLE STRAINS OF PENICILLIN-RESISTANT STREPTOCOCCUS PNEUMONIAE IN PATIENTS WITH COMMUNITY ACQUIRED PNEUMONIA
I-306	INDUCTION OF SPERMATOGENESIS IN MEN WITH PRIMARY AND SECONDARY HYPOGONADOTROPIC HYPOGONADISM IN WHOM THE CAUSE OF INFERTILITY IS NOT DUE TO PRIMARY TESTICULAR FAILURE
I-307	NEW COMBINATION USE OF METFORMIN AND INSULIN IN TYPE 2 DIABETES
I-308	TREATMENT OF PEDIATRIC PATIENTS WITH POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS WHO RESPONDED INADEQUATELY TO SALICYLATES OR OTHER NSAIDS
I-309	USE OF ACTONEL 35MG ONCE A WEEK TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
I-310	REDUCTION IN RISK OF MYOCARDIAL INFARCTION, STROKE, AND DEATH FROM CARDIOVASCULAR CAUSES
I-311	ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS AGE 3 TO 12 YEARS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-312	FIRST LINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER
I-313	EXTENSION OF INDICATION TO PROVIDE FOR MAINTENANCE OF RESPONSE
I-314	TOPICAL ANESTHETIC FOR SUPERFICIAL MINOR SURGERY OF GENITAL MUCOUS MEMBRANES AND AS AN ADJUNCT FOR LOCAL INFILTRATION ANESTHESIA IN GENITAL MUCOUS MEMBRANES
I-315	THROMBOPROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS
I-316	TREATMENT OF NSAID-ASSOCIATED GASTRIC ULCER PATIENTS WHO CONTINUE NSAID USE AND REDUCING RISK OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS WITH HISTORY OF DOCUMENTED GASTRIC ULCER WHO REQUIRE USE OF AN NSAID
I-317	PROPHYLAXIS OF INFLUENZA IN ADULTS AND ADOLESCENTS 13 YEARS AND OLDER
I-318	FIRSTLINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER
I-319	USE FOR SUSPECTED OR CONFIRMED METHANOL POISONING, EITHER ALONE OR IN COMBINATION WITH HEMODIALYSIS
I-320	TREATMENT OF TYPE 2 DIABETES IN PEDIATRIC PATIENTS (AGES 10-16 YEARS)
I-321	JUVENILE RHEUMATOID ARTHRITIS
I-322	USE OF DIPRIVAN IN PATIENTS 3 MONTHS TO 16 YEARS
I-323	COLORECTAL CANCER
I-324	REDUCING NEUROLOGIC DISABILITY AND/OR FREQUENCY OF CLINICAL RELAPSES IN PATIENTS WITH SECONDARY (CHRONIC) PROGRESSIVE, PROGRESSIVE RELAPSING, OR WORSENING RELAPSING-REMITTING MULTIPLE SCLEROSIS
I-325	PREVENTION OF RELAPSE AND RECURRENCE OF DEPRESSION
I-326	GENERALIZED ANXIETY DISORDER
I-327	SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN PATIENTS 5 YEARS AND OLDER
I-328	PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 5-6 YEARS OF AGE
I-329	UNCOMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS
I-330	MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND CONTROL OF DAYTIME AND NIGHTTIME HEARTBURN SYMPTOMS IN PATIENTS WITH GERD
I-331	TREATMENT OF MODERATE ACNE VULGARIS
I-332	EMPIRIC THERAPY IN FEBRILE NEUTROPENIC PATIENTS WITH SUSPECTED FUNGAL INFECTIONS (EFTN)
I-333	TOPICAL TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR DUE TO MALASSEZIA FURFUR (FORMERLY PITYROSPORUM ORBICULARE)
I-334	LONG-TERM TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE WHO FAIL TO MANIFEST CATCH-UP GROWTH BY TWO YEARS OF AGE
I-335	ADJUNCTIVE THERAPY IN PATIENTS TWO YEARS AND OLDER WITH SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME
I-336	EXPANSION OF INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH PREDOMINATELY CLASSIC SUBFOVEAL CHOROIDAL NEOVASCULARIZATION DUE TO PATHOLOGIC MYOPIA OR PRESUMED OCULAR HISTOPLASMOSIS
I-337	PATHOLOGICAL HYPERSECRETION ASSOCIATED WITH ZOLLINGER-ELLISON SYNDROME
I-338	MANAGEMENT OF ACUTE PAIN IN ADULTS AND TREATMENT OF PRIMARY DYSMENORRHEA
I-339	TREATMENT OF HEPATITIS B IN PEDIATRIC PATIENTS AGES 2-17 YEARS
I-340	ATOPIC DERMATITIS IN PEDIATRIC PATIENTS AGES 2-5
I-341	BREAST CANCER COMBINATION THERAPY
I-342	USE OF FORADIL FOR LONG-TERM, TWICE DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHO-CONSTRICTION IN PATIENTS WITH COPD INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
I-343	USE OF COREG FOR SEVERE HEART FAILURE
I-344	ACNE VULGARIS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-345 TREATMENT OF POSTTRAUMATIC STRESS DISORDER

I-346 TREATMENT OF SYMPTOMATIC GASTRO ESOPHAGEAL REFLUX DISEASE (GERD)

I-347 TREATMENT OR PREVENTION OF BRONCHOSPASM IN CHILDREN 6 YEARS OF AGE AND OLDER WITH OBSTRUCTIVE AIRWAY DISEASE

I-348 LONG-TERM, TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD (INCLUDING EMPHYSEMA AND CHRONIC BRONCHITIS)

I-349 ACUTE CORONARY SYNDROME

I-350 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND GIRLS AT LEAST ONE YEAR POSTMENARCHAL, AGES 10 TO 17 YEARS, WITH A RECOMMENDED DOSING RANGE OF 10 TO 40MG ONCE DAILY

I-351 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR ALL STRENGTHS

I-352 ANTICOAGULANT IN PATIENTS WITH OR AT RISK FOR HEPARIN-INDUCED THROMBOCYTOPENIA UNDERGOING PERCUTANEOUS CORONARY INTERVENTIONS (PCI)

I-353 TREATMENT OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS

I-354 MANAGEMENT OF POST HERPETIC NEURALGIA

I-355 PREMENSTRUAL DYSPHORIC DISORDER

I-356 TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS, INCLUDING ZOLLINGER-ELLISON SYNDROME

I-357 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS

I-358 TREATMENT OF PANIC DISORDER

I-359 TREATMENT OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE

I-360 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN CHILDREN AGES TWO UP TO AGE THREE

I-361 TREATMENT OF MULTIPLE MYELOMA AND DOCUMENTED BONE METASTASES FROM SOLID TUMORS, IN CONJUNCTION WITH STANDARD ANTINEOPLASTIC THERAPY. PROSTATE CANCER SHOULD HAVE PROGRESSED AFTER TREATMENT WITH AT LEAST ONE HORMONAL THERAPY

I-362 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA

I-363 ADJUVANT TREATMENT OF POST MENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER

I-364 TREATMENT OF COMMUNITY-ACQUIRED PNEUMONIA IN ADULTS

I-365 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV) IN PATIENTS WHO ARE INTOLERANT TO AN ACE INHIBITOR

I-366 PREVENTION OF RELAPSE FOLLOWING LONG-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER

I-367 COMBINATION THERAPY WITH THIAZOLIDINEDIONE TO LOWER BLOOD GLUCOSE IN PTS WHOSE HYPERGLYCEMIA CANNOT BE CONTROLLED BY DIET/EXERCISE PLUS MONOTHERAPY WITH ANY OF THE FOLLOWING AGENTS: METFORMIN, SULFONYLUREAS, REPAGLINIDE, OR THIAZOLIDINEDIONES

I-368 USE OF GLUCOVANCE WITH A THIAZOLIDINEDIONE WHEN GLYCEMIC CONTROL IS NOT OBTAINED WITH GLUCOVANCE ALONE

I-369 PREVENTION AND TREATMENT OF POSTOPERATIVE NAUSEA AND VOMITING

I-370 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN CHILDREN, AGES 8-13 YEARS, WITH RECOMMENDED DOSE OF 20MG ONCE DAILY AND IN ADOLESCENTS, AGES 14-18 WITH A RECOMMENDED DOSE OF 40MG ONCE DAILY

I-371 HELICOBACTER PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

I-372 NOSOCOMIAL PNEUMONIA

I-373 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY

I-374 SHORT TERM TOPICAL TREATMENT OF MILD TO MODERATE PLAQUE-TYPE PSORIASIS OF NON SCALP REGIONS

I-375 FIRST LINE THERAPY FOR THE REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION

I-376 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (CML)

I-377 USE OF BRAVELLE FOR MULTIPLE FOLLICULAR DEVELOPMENT (CONTROLLED OVARIAN

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- STIMULATION) DURING ASSISTED REPRODUCTIVE TECHNOLOGY CYCLES IN PATIENTS WHO HAVE PREVIOUSLY RECEIVED PITUITARY SUPPRESSION
- I-378 RELIEF OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-379 USE TAXOTERE IN COMBINATION WITH CISPLATIN FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHO HAVE NOT PREVIOUSLY RECEIVED CHEMOTHERAPY FOR THIS CONDITION
- I-380 TO TREAT PATIENTS WITH SCHIZOPHRENIA OR SCHIZOAFFECTIVE DISORDER AT RISK FOR EMERGENT SUICIDAL BEHAVIOR
- I-381 TREATMENT OF COLD SORES (HERPES LABIALIS) IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER
- I-382 FOR NEWLY-DIAGNOSED HIGH GRADE MALIGNANT GLIOMA PATIENTS AS AN ADJUNCT TO SURGERY AND RADIATION
- I-383 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY
- I-384 USE IN COMBINATION WITH INSULIN FOR THE TREATMENT OF PATIENTS WITH TYPE 2 DIABETES MELLITUS
- I-385 MODIFICATION OF THE INDICATION FOR COMMUNITY ACQUIRED PNEUMONIA TO ADD "INCLUDING PENICILLIN-RESISTANT STRAINS, MIC PENICILLIN \geq 2MCG/ML" TO STREPTOCOCCUS PNEUMONIAE
- I-386 RAPAMUNE (SIROLIMUS) WITHIN AN IMMUNOSUPPRESSIVE REGIMEN THAT WOULD ALLOW FOR THE WITHDRAWAL OF CYCLOSPORINE 2 TO 4 MONTHS AFTER RENAL TRANSPLANTATION IN PATIENTS CONSIDERED AT LOW TO MODERATE IMMUNOLOGIC RISK FOR RENAL TRANSPLANT REJECTION
- I-387 ADJUNCTIVE THERAPY OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE
- I-388 TREATMENT OF PATIENTS WITH LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- I-389 SUPPRESSION OF RECURRENT GENITAL HERPES IN HIV-INFECTED INDIVIDUALS
- I-390 USE IN PTS AT HIGH RISK CORONARY EVENTS DUE TO EXISTING CORONARY HEART DISEASE, DIABETES, PERIPHERAL VESSEL DISEASE, STROKE HISTORY, OTHER CV DISEASE TO REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH, REDUCE NONFATAL MI & STROKE.....
- I-391 ABLATION OF HIGH-GRADE DYSPLASIA IN BARRETT'S ESOPHAGUS PATIENTS WHO DO NOT UNDERGO ESOPHAGECTOMY
- I-392 TX OF PED PATIENTS W/PH+ CHRONIC PHASE CML DISEASE RECURRENCE AFTER STEM CELL TRANSPLANT OR RESISTANCE TO INTERFERON ALPHA THERAPY. NO CONTROLLED TRIALS DEMONSTRATING A CLINICAL BENEFIT SUCH AS IMPROVEMENT IN DISEASE RELATED SX OR INCREASED SURVIVAL
- I-393 CHRONIC BACTERIAL PROSTATITIS
- I-394 USE IN PATIENTS WITH CORONARY HEART DISEASE TO REDUCE THE RISK OF UNDERGOING CORONARY REVASCULARIZATION PROCEDURES
- I-395 TO IMPROVE PHYSICAL FUNCTION
- I-396 EXPANDED INDICATION TO INCLUDE THE ASSESSMENT OF VENTRICULAR FUNCTION IN SUBJECTS BEING EVALUATED FOR HEART DISEASE AND/OR VENTRICULAR FUNCTION
- I-397 EXTENDED PROPHYLAXIS IN PATIENTS UNDERGOING HIP FRACTURE SURGERY
- I-398 IDIOPATHIC SHORT STATURE
- I-399 TREATMENT OF CANDIDEMIA AND THE FOLLOWING CANDIDA INFECTIONS: INTRA-ABDOMINAL ABSCESES, PERITONITIS AND PLEURAL SPACE INFECTIONS
- I-400 USE OF OLANZAPINE IN COMBINATION WITH LITHIUM OR VALPROATE FOR THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-401 LONGER-TERM EFFICACY OF ARIPIRAZOLE IN THE TREATMENT OF SCHIZOPHRENIA
- I-402 DIABETIC FOOT INFECTIONS WITHOUT CONCOMITANT OSTEOMYELITIS
- I-403 USE OF VALTREX IN COMBINATION WITH SAFER SEX PRACTICES FOR THE REDUCTION OF THE RISK OF TRANSMISSION OF GENITAL HERPES DURING SUPPRESSIVE THERAPY OF THE SOURCE PARTNER IN A HETEROSEXUAL COUPLE
- I-404 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES (DEPRESSION, MANIA, HYPOMANIA, MIXED EPISODES) IN PATIENTS TREATED

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY

- I-405 TREATMENT OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) USING AN INTERMITTENT DOSING REGIMEN
- I-406 PREVENTION OF CYTOMEGALOVIRUS DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)
- I-407 IMPROVE SURVIVAL OF STABLE PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (EJECTION FRACTION<=40%) AND CLINICAL EVIDENCE OF CONGESTIVE HEART FAILURE AFTER AN ACUTE MYOCARDIAL INFARCTION
- I-408 STIMULATION OF PANCREATIC SECRETIONS TO FACILITATE THE IDENTIFICATION OF THE AMPULLA OF VATER AND ACCESSORY PAPILLA DURING ENDOSCOPIC RETROGRADE CHOLANGIO-PANCREATOGRAPHY (ERCP)
- I-409 ESOPHAGEAL CANDIDIASIS
- I-410 USE OF ADVAIR DISKUS 250/50 FOR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) ASSOCIATED WITH CHRONIC BRONCHITIS
- I-411 EXPANDED INDICATION FOR USE IN COMBINATION WITH ANTIDIABETIC DRUGS IN THE THIAZOLIDINEDIONE CLASS
- I-412 MONOTHERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-413 ADJUNCTIVE THERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-414 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM (PE) IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS
- I-415 SEVERE HYPERTENSION WHEN THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY
- I-416 THE USE OF CIPRO XR FOR COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
- I-417 USE IN THE LONG TERM TREATMENT OF BIPOLAR I DISORDER
- I-418 ADJUNCTIVE THERAPY W/ MOOD STABILIZERS (LITHIUM OR DIVALPROEX) IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDERS
- I-419 MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-420 TOPICAL TREATMENT OF CLINICALLY TYPICAL, NONHYPERKERATOTIC, NONHYPERTROPHIC ACTINIC KERATOSES ON THE FACE OR SCALP IN IMMUNOCOMPETENT ADULTS
- I-421 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND PYELONEPHRITIS DUE TO E.COLI FOR PED PATIENTS (1-17) NOT AS FIRST CHOICE
- I-422 INDICATED FOR THE IN-HOSPITAL SHORT-TERM (UP TO 4 HOURS) REDUCTION IN BLOOD PRESSURE IN PEDIATRIC PATIENTS
- I-423 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS
- I-424 MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL INSUFFICIENCY NOT YET ON DIALYSIS
- I-425 ELOXATIN IN COMBINATION WITH INFUSIONAL 5-FLUOROURACIL (5-FU) AND LEUCOVORIN (LV) FOR THE TREATMENT OF PATIENTS PREVIOUSLY UNTREATED FOR ADVANCED COLORECTAL CANCER
- I-426 TREATMENT OF ACUTE PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-427 TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-428 FOR USE IN COMBINATION WITH PACLITAXEL FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR ANTHRACYCLINE CONTAINING ADJUVANT CHEMOTHERAPY UNLESS ANTHRACYCLINES WERE CLINICALLY CONTRAINDICATED
- I-429 FOR USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER
- I-430 FOR USE IN THE RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS
- I-431 NOSOCOMIAL PNEUMONIA AND COMMUNITY-ACQUIRED PNEUMONIA CAUSED BY STREPTOCOCCUS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- PNEUMONIAE INDICATION EXPANDED TO INCLUDE MULTI-DRUG RESISTANT STRAINS
- I-432 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA CAUSED BY MULTI-DRUG RESISTANT STREPTOCOCCUS PNEUMONIAE
- I-433 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA IN IMMUNOCOMPETENT ADULTS, WITH A MAXIMUM TUMOR DIAMETER OF 2.0CM, LOCATED ON THE TRUNK (EXCLUDING ANOGENITAL SKIN), NECK, OR EXTREMITIES (EXCLUDING HANDS AND FEET)
- I-434 PREVENTION OF CARDIOVASCULAR DISEASE IN ADULT PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE, BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE TO REDUCE RISK OF MI AND RISK FOR REVASCULARIZATION PROCEDURES AND ANGINA
- I-435 CHRONIC IDIOPATHIC CONSTIPATION
- I-436 FOR USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR THE ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- I-437 TREATMENT OF ACUTE MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-438 EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS
- I-439 USED TO TREAT ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-440 FOR THE REPLACEMENT OF ENDOGENOUS GROWTH HORMONE IN ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-441 USE COMBINATION WITH INFUSIONAL 5-FU/LV FOR ADJUVANT TREATMENT STAGE III COLON CANCER PTS WHO HAVE UNDERGONE COMPLETE RESECTION PRIMARY TUMOR-BASED ON IMPROVEMENT IN DISEASE FREE SURVIVAL, NO DEMONSTRATED BENEFIT OVERALL SURVIVAL AFTER 4YRS
- I-442 USED FOR CANDIDEMIA IN NONNEUTROPENIC PATIENTS AND THE FOLLOWING CANDIDA INFECTIONS: DISSEMINATED INFECTIONS IN SKIN & INFECTIONS IN ABDOMEN, KIDNEY, BLADDER WALL, AND WOUNDS
- I-443 TREATMENT OF NASAL POLYPS IN PATIENTS 18 YEARS OF AGE AND OLDER
- I-444 USE OF PROTONIX IV FOR INJECTION AS STAND ALONE THERAPY FOR THE SHORT-TERM TREATMENT OF PATIENTS HAVING GASTROESOPHAGEAL REFLUX (GERD) WITH A HISTORY OF EROSIVE ESOPHAGITIS
- I-445 TO IMPROVE (COMPARED TO 4.25% DEXTROSE) LONG-DWELL ULTRAFILTRATION AND CLEARANCE OF CREATININE AND UREA NITROGEN IN PATIENTS WITH HIGH AVERAGE OR GREATER TRANSPORT CHARACTERISTICS, AS DEFINED USING THE PERITONEAL EQUILIBRATION TEST (PET)
- I-446 EXTENDED ADJUVANT TREATMENT OF EARLY BREAST CANCER IN POSTMENOPAUSAL WOMEN WHO HAVE RECEIVED 5 YRS ADJUVANT TAMOXIFEN THERAPY-EFFECTIVENESS BASED ON AN ANALYSIS OF DISEASE FREE SURVIVAL IN PATIENTS TREATED FOR A MEDIAN 24 MONTHS
- I-447 USE OF COPEGUS (RIBAVIRIN) FOR TREATMENT OF CHRONIC HEPATITIS C IN ADULT PATIENTS COINFECTED WITH HIV IN COMBINATION WITH PEGASYS (PEGINTERFERON ALFA-2A)
- I-448 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV AND EJECTION FRACTION <=40%) TO REDUCE THE RISK OF DEATH FROM CARDIOVASCULAR CAUSES AND TO REDUCE HOSPITALIZATIONS FOR HEART FAILURE
- I-449 TO IMPROVE WAKEFULNESS IN TWO NEW PATIENT POPULATIONS WITH EXCESSIVE SLEEPINESS: THOSE WITH OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME AND THOSE WITH SHIFT WORK SLEEP DISORDER
- I-450 TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED HIGH GRADE GLIOMAS CONCOMITANTLY WITH RADIOTHERAPY AND THEN AS ADJUVANT TREATMENT
- I-451 MANAGEMENT OF ENDOMETRIOSIS ASSOCIATED PAIN
- I-452 EXPANDED INDICATION TO INCLUDE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY
- I-453 USE IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN WHEN DIET, EXERCISE AND BOTH AGENTS DO NOT RESULT IN ADEQUATE GLYCEMIC CONTROL (TRIPLE THERAPY)
- I-454 MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON FOR UP TO 3 MONTHS
- I-455 MODIFIED HEART FAILURE INDICATION TO INCLUDE TREATMENT OF HEART FAILURE IN PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (NYHA CLASS II-IV; EJECTION FRACTION LESS THAN OR EQUAL TO 40%)
- I-456 TO REDUCE CARDIOVASCULAR DEATH AND TO REDUCE HEART FAILURE HOSPITALIZATIONS.

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

INCLUDES ADDITIONAL INFORMATION ON THE ADDED EFFECT ON THESE OUTCOMES WHEN USED WITH AN ACE INHIBITOR

- I-457 TREATMENT OF PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS
- I-458 USE OF BIVALIRUDIN FOR INJECTION WITH PROVISIONAL USE OF GLYCOPROTEIN IIB/IIIA INHIBITOR (GPI) AS LISTED IN THE CLINICAL TRIALS REPLACE-2 SECTION FOR USE AS AN ANTICOAGULANT IN PATIENTS UNDERGOING PERCUTANEOUS CORONARY INTERVENTION (PCI)
- I-459 NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE (NDD-CKD) PATIENTS RECEIVING OR NOT RECEIVING AN ERYTHROPOIETIN
- I-460 TREATMENT OF DIARRHEA CAUSED BY CRYPTOSPORIDIUM PARVUM IN NON-HIV INFECTED PATIENTS 12 YEARS OF AGE AND OLDER
- I-461 USE AS A SINGLE AGENT FOR ADJUVANT TREATMENT IN PATIENTS WITH DUKES' C COLON CANCER WHO HAVE UNDERGONE COMPLETE RESECTION OF THE PRIMARY TUMOR WHEN TREATMENT WITH FLUOROPYRIMIDINE THERAPY ALONE IS PREFERRED
- I-462 LONG TERM TREATMENT OF IDIOPATHIC SHORT STATURE
- I-463 TREATMENT OF PATIENTS POST MYOCARDIAL INFARCTION
- I-464 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME
- I-465 PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 6 MONTHS OF AGE AND OLDER
- I-466 FOR RELIEF OF THE SIGNS AND SYMPTOMS OF ANKYLOSING SPONDYLITIS
- I-467 USE OF TOPIRAMATE AS INITIAL MONOTHERAPY IN PATIENTS 10 YEARS OF AGE AND OLDER WITH PARTIAL ONSET OR PRIMARY GENERALIZED TONIC CLONIC SEIZURES
- I-468 USE IN PATIENTS WITH STABLE CORONARY ARTERY DISEASE TO REDUCE THE RISK OF CARDIOVASCULAR MORTALITY OR NON-FATAL MYOCARDIAL INFARCTION
- I-469 RELIEF OF THE SIGNS AND SYMPTOMS OF PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS IN PATIENTS 2 YEARS OF AGE AND OLDER
- I-470 DIABETIC PERIPHERAL NEUROPATHIC PAIN
- I-471 INDICATED TO REDUCE THE RISK OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH TYPE 2 DIABETES AND WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE
- I-472 USE IN PATIENTS WITH ANGIOGRAPHICALLY DOCUMENTED CORONARY ARTERY DISEASE
- I-473 USE IN COMBINATION WITH GEMCITABINE FOR THE FIRST LINE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER
- I-474 TREATMENT OF IRON DEFICIENCY ANEMIA IN PERITONEAL DIALYSIS DEPENDANT CHRONIC KIDNEY DISEASE IN PATIENTS RECEIVING AN ERYTHROPOIETIN
- I-475 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- I-476 TREATMENT OF DIABETIC FOOT INFECTIONS WITHOUT OSTEOMYELITIS
- I-477 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY METHICILLIN SUSCEPTIBLE STAPHYLOCOCCUS AUREUS, ESCHERICHIA COLI, KLEBSIELLA PNEUMONIAE, OR ENTEROBACTER CLOACAE
- I-478 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN CHILDREN WITH EPILEPSY AGED 2-4 YEARS
- I-479 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS CAUSED BY E.COLI, B. FRAGILIS, S.ANGINOSUS, S.CONSTELLATUS, E. FAECALIS, P. MIRABILIS, C. PERFRINGENS, B. THETA IOTAOMICRON OR PEPTOSTREPTOCOCCUS SPECIES
- I-480 PROPHYLAXIS OF INFLUENZA FOR PATIENTS BETWEEN 1-12 YEARS OF AGE
- I-481 INDICATED FOR THE ADJUVANT TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER
- I-482 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH OR WITHOUT PSYCHOTIC FEATURES
- I-483 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- I-484 FOR THE RISK REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS
- I-485 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-486	ANGIOMAX IS INDICATED FOR PATIENTS WITH, OR AT RISK OF, HIT/HITTS UNDERGOING PCI
I-487	INDICATED FOR THE RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YRS OF AGE OR OLDER
I-488	MAINTENANCE THERAPY IN BIPOLAR I DISORDER
I-489	FOR USE IN PEDIATRIC PATIENTS WITH TYPE I DIABETES
I-490	FOR USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE TREATMENT OF PATIENTS WITH ADVANCED GASTRIC ADENOCARCINOMA, INCLUDING ADENOCARCINOMA OF GASTROESOPHAGEAL JUNCTION, WHO HAVE NOT RECEIVED PRIOR CHEMOTHERAPY FOR ADVANCED DISEASE
I-491	INFLUENZA PROPHYLAXIS
I-492	MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC OR MIXED EPISODES IN BIPOLAR I DISORDER, WITH OR WITHOUT PSYCHOTIC FEATURES
I-493	ADMINISTERED IN COMBINATION WITH FENOFIBRATE, AS ADJUNCTIVE THERAPY TO DIET FOR THE REDUCTION OF ELEVATED TOTAL-C, LDL-C, APO B, AND NON-HDL-C IN PATIENTS WITH MIXED HYPERLIPIDEMIA
I-494	CLINICAL DATA IN SUPPORT OF AVANDAMET AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH DUAL ROSIGLITAZONE AND METFORMIN THERAPY IS APPROPRIATE
I-495	ADJUVANT TX OF POSTMENOPAUSAL WOMEN WITH ESTROGEN-RECEPTOR POSITIVE EARLY BREAST CANCER WHO HAVE RECEIVED 2 TO 3 YRS OF TAMOXIFEN AND ARE SWITCHED TO AROMASIN FOR COMPLETION OF A TOTAL OF 5 CONSECUTIVE YRS OF ADJUVANT HORMONAL THERAPY
I-496	LONG TERM TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME IN PATIENTS WHO HAVE OPEN EPIPHYSES
I-497	PREVENTION OF SEASONAL MAJOR DEPRESSIVE EPISODES IN PATIENTS WITH SEASONAL AFFECTIVE DISORDER
I-498	PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
I-499	USE OF GEMZAR IN COMBINATION WITH CARBOPLATIN FOR THE TREATMENT OF PATIENTS WITH ADVANCED OVARIAN CANCER THAT HAS RELAPSED AT LEAST 6 MONTHS AFTER COMPLETION OF PLATINUM-BASED THERAPY
I-500	FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
I-501	TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN IMMUNOCOMPETENT PATIENTS WITH A SINGLE DOSE OF FAMCICLOVIR 1500 MG.
I-502	FOR PTS WITH ST-SEGMENT ELEVATION ACUTE MYOCARDIAL INFARCTION, PLAVIX TO REDUCE RATE OF DEATH FROM ANY CAUSE AND THE RATE OF A COMBINED ENDPOINT OF DEATH, REINFARCTION OR STROKE. NOT KNOWN TO PERTAIN TO PTS WHO RECEIVE PRIMARY ANGIOPLASTY
I-503	TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER
I-504	TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME
I-505	TREATMENT OF STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA), INCLUDING THOSE WITH RIGHT SIDED INFECTIVE ENDOCARDITIS, CAUSED BY METHICILLIN-SUSCEPTIBLE AND METHICILLIN-RESISTANT ISOLATES
I-506	ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 12 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY
I-507	ADJUNCT TO DIET TO REDUCE TOTAL-C, LDL-C AND APO B LEVELS IN ADOLESCENT BOYS AND GIRLS WHO ARE AT LEAST ONE YEAR POST-MENARCHE, 10-16 YEARS OF AGE, WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
I-508	PREMENSTRUAL DYSPHONIC DISORDER
I-509	TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
I-510	ADULT DERMATOFIBROSARCOMA PROTUBERANS (DFSP)
I-511	ADULT MYELODYSPLASTIC SYNDROME/MYELOPROLIFERATIVE DISEASES (MDS/MDP)
I-512	ADULT PH+ ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) MONOTHERAPY
I-513	ADULT AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM)
I-514	ADULT HYPEREOSINOPHILIC SYNDROME/CHRONIC EOSINOPHILIC LEUKEMIA (HES/CEL)

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-515 PROPHYLAXIS OF SURGICAL SITE INFECTION FOLLOWING ELECTIVE COLORECTAL SURGERY

I-516 PRIMARY GENERALIZED TONIC CLONIC SEIZURES IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER

I-517 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEG SYNDROME (RLS)

I-518 TREATMENT OF SHORT STATURE OR GROWTH FAILURE IN CHILDREN WITH SHOX (SHORT STATURE HOMEBOX CONTAINING GENE) DEFICIENCY WHOSE EPIPHYSES ARE NOT CLOSED

I-519 USE OF TAXOTERE (DOCETAXEL) INJECTION CONCENTRATE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN)

I-520 USE OF EXENATIDE IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE USING A THIAZOLIDINEDIONE ALONE OR IN COMBINATION WITH METFORMIN BUT HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL

I-521 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 YEAR PRIOR THERAPY

I-522 TREATMENT OF MODERATE ACNE VULGARIS IN WOMEN AT LEAST 14 YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, AND HAVE ACHIEVED MENARCHE, IF THE PATIENT DESIRES AN ORAL CONTRACEPTIVE FOR BIRTH CONTROL.

I-523 USE IN ADULT PATIENTS WITH CLINICALLY EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF NONFATAL MYOCARDIAL INFARCTION, FATAL AND NONFATAL STROKE, ANGINA, REVASCULARIZATION PROCEDURES AND HOSPITALIZATION FOR CONGESTIVE HEART FAILURE

I-524 GENERALIZED ANXIETY DISORDER (GAD)

I-525 USE OF 0.5MG/0.1MG FOR PREVENTION OF POST-MENOPAUSAL OSTEOPOROSIS

I-526 TREATMENT OF HYPONATREMIA IN HOSPITALIZED PATIENTS

I-527 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY

I-528 TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE

I-529 TREATMENT OF DEMENTIA OF THE ALZHEIMER'S TYPE IN PATIENTS WITH SEVERE ALZHEIMER'S DISEASE

I-530 PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION IN PATIENTS 15 YEARS OF AGE AND OLDER

I-531 MAINTENANCE TREATMENT OF SCHIZOPHRENIA

I-532 TREATMENT OF BACTERIAL VAGINOSIS IN NON-PREGNANT FEMALES

I-533 ACUTE ST-SEGMENT ELEVATION MYOCARDIAL INFARCTION (STEMI)

I-534 EXTENDED TREATMENT OF SYMPTOMATIC VENOUS THROMBOEMBOLISM (VTE) AND/OR PULMONARY EMBOLISM TO REDUCE THE RECURRENCE OF VTE IN PATIENTS WITH CANCER

I-535 MANAGEMENT OF FIBROMYALGIA

I-536 FOR THE TREATMENT OF SHORT STATURE IN CHILDREN WITH NOONAN SYNDROME

I-537 LONG TERM TREATMENT OF PANIC DISORDER

I-538 SHORT TERM TREATMENT OF PANIC DISORDER

I-539 REDUCTION IN RISK OF INVASIVE BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS OR AT HIGH RISK FOR INVASIVE BREAST CANCER

I-540 TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGES 13-17

I-541 TREATMENT OF BIPOLAR I DISORDER IN CHILDREN AGES 10-12 AND ADOLESCENTS AGES 13-17

I-542 EXPANSION OF PATIENT POPULATION FOR HEAD AND NECK CANCER FROM "INOPERABLE" PATIENTS TO ALL PATIENTS

I-543 USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION TREATMENT OF PATIENTS WITH LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN)

I-544 ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 16 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY

I-545 ADJUNCTIVE TREATMENT TO TREAT PATIENTS WITH MAJOR DEPRESSIVE DISORDER

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-546 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA

I-547 ADJUNCTIVE THERAPY TO DIET TO SLOW THE PROGRESSION OF ARTEROSCLEROSIS IN ADULT PATIENTS AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL-C AND LDL-C TO TARGET LEVELS

I-548 SEASONAL ALLERGIC RHINITIS IN PATIENTS 6 THROUGH LESS THAN 12 YEARS OF AGE

I-549 USE OF AVALIDE TABLETS AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS

I-550 TREATMENT OF HYPERTENSION IN PEDIATRIC PATIENTS 6-16 YEARS OF AGE

I-551 TREATMENT OF SHORT STATURE IN CHILDREN WITH TURNER'S SYNDROME

I-552 ADJUNCTIVE TREATMENT FOR RADIOIODINE ABLATION OF THYROID TISSUE REMNANTS IN PATIENTS WHO HAVE UNDERGONE THYROIDECTOMY FOR WELL-DIFFERENTIATED THYROID CANCER AND WHO DO NOT HAVE EVIDENCE OF METASTATIC THYROID CANCER

I-553 FOR USE AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

I-554 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESES

I-555 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER IN PEDIATRIC PATIENTS AGED 10-17 YEARS

I-556 PREVENTION OF POST OPERATIVE NAUSEA AND VOMITING FOR UP TO 24 HOURS FOLLOWING SURGERY

I-557 USE OF AMITIZA (LUBIPROSTONE) 8 MCG TWICE DAILY FOR TREATMENT OF IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN WOMEN GREATER THAN OR EQUAL TO 18 YEARS OLD

I-558 MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION AND REDUCING EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA

I-559 ADJUNCTIVE THERAPY ADDED TO LITHIUM OR VALPROATE IN SHORT TERM TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED

I-560 MAINTENANCE TREATMENT FOR BIPOLAR I DISORDER, AS ADJUNCTIVE THERAPY TO LITHIUM OR DIVALPROEX

I-561 LONG-TERM TREATMENT OF SOCIAL ANXIETY DISORDER

I-562 MAINTENANCE TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD) IN CHILDREN AND ADOLESCENTS

I-563 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND CHILDREN 16 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY

I-564 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA

I-565 USE OF DUTASTERIDE IN COMBINATION WITH TAMSULOSIN FOR THE TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA (BPH)

I-566 MANAGEMENT OF FIBROMYALGIA

I-567 INITIAL THERAPY IN PATIENTS LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS

I-568 USE OF APTIVUS, CO-ADMINISTERED W/RITONAVIR, FOR COMBINATION ANTIRETROVIRAL TREATMENT OF HIV-1 INFECTED PED (AGE 2-18 YRS) PATIENTS WHO ARE TREATMENT-EXPERIENCED AND INFECTED W/HIV-1 STRAINS RESISTANT TO MORE THAN ONE PROTEASE INHIBITOR

I-569 TREATMENT OF CHRONIC HEPATITIS B

I-570 TREATMENT OF CHICKEN POX IN IMMUNOCOMPETENT PEDIATRIC PATIENTS 2 TO <18 YEARS OF AGE

I-571 NON-SMALL CELL LUNG CANCER IN COMBINATION WITH CISPLATIN AND AS SINGLE AGENT FOR NONSQUAMOUS NON-SMALL CELL LUNG CANCER

I-572 TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE (SGA) WITH NO CATCH-UP BY AGE 2-4 YRS.

I-573 TO TREAT PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDRICKSON TYPE III HYPERLIPOPROTEINEMIA) AS AN ADJUNCT TO DIET

I-574 MONOTHERAPY IN THE TREATMENT OF BIPOLAR DEPRESSION

I-575 MONOTHERAPY IN THE TREATMENT OF BIPOLAR MANIA

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-576 ADJUNCTIVE THERAPY IN THE TREATMENT OF BIPOLAR MANIA

I-577 SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES

I-578 EXPANSION OF INDICATION TO INCLUDE TREATMENT OF HIV IN TREATMENT NAIVE ADULTS

I-579 TREATMENT OF MODERATE TO SEVERE DYSpareunia, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE AND NEW TWICE WEEKLY DOSING REGIMEN FOR THIS INDICATION

I-580 INDOLENT B-CELL NON-HODGKINS LYMPHOMA (NHL) THAT HAS PROGRESSED DURING OR WITHIN SIX MONTHS OF TREATMENT WITH RITUXIMAB OR A RITUXIMAB CONTAINING REGIMEN

I-581 TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

I-582 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

I-583 ADJUVANT TREATMENT OF ADULT PATIENTS FOLLOWING COMPLETE GROSS RESECTION OF KIT (CD117) POSITIVE GASTROINTESTINAL STROMAL TUMORS (GIST)

I-584 TREATMENT AND PREVENTION OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN PATIENTS EXPECTED TO BE ON GLUCOCORTICIDS FOR AT LEAST 12 MONTHS

I-585 TREATMENT OF SHORT STATURE IN PEDIATRIC PATIENTS SMALL FOR GESTATIONAL AGE WHO DO NOT MANIFEST CATCH UP GROWTH BY AGE 2 TO 4 YEARS

I-586 COMMUNITY ACQUIRED BACTERIAL PNEUMONIA

I-587 ADDITIONAL PATHOGENS TO COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS INDICATION

I-588 ADDITIONAL PATHOGENS TO COMPLICATED INTRA-ABDOMINAL INFECTIONS INDICATION

I-589 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH OLANZAPINE

I-590 ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH OLANZAPINE)

I-591 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH FLUOXETINE

I-592 ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH FLUOXETINE)

I-593 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD)

I-594 INDICATION EXPANDED TO INCLUDE PATIENTS WHO HAVE EXPERIENCED A FIRST CLINICAL EPISODE AND HAVE MRI FEATURES CONSISTENT WITH MULTIPLE SCLEROSIS

I-595 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

I-596 USE AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER

I-597 MONOTHERAPY FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER

I-598 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION INDICATION EXPANDED TO INCLUDE DELAY IN CLINICAL WORSENING

I-599 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS

I-600 FOR USE AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS

I-601 MAINTENANCE TREATMENT IN PATIENTS WITH ADVANCED OR METASTATIC NONSQUAMOUS NON-SMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST LINE CHEMOTHERAPY

I-602 TREATMENT OF MEN AND WOMEN WITH OSTEOPOROSIS ASSOCIATED WITH SUSTAINED SYSTEMIC GLUCOCORTICOID THERAPY AT HIGH RISK FOR FRACTURE

I-603 GOUT FLARES

I-604 PREVENTION OF CMV DISEASE IN KIDNEY AND HEART TRANSPLANT PATIENTS 4 MONTHS TO 16 YEARS AT HIGH RISK

I-605 ADJUNCT TO MOOD STABILIZERS AND/OR ANTIDEPRESSANTS FOR SCHIZOAFFECTIVE DISORDER

I-606 TREATMENT OF SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY

I-607 INDICATION EXPANDED TO INCLUDE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP I) IN PATIENTS WITH CLASS II SYMPTOMS

I-608 REDUCE LDL-C LEVELS IN BOYS AND POSTMENARCHAL GIRLS, 10 TO 17 YEARS OF AGE, WITH

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA AS MONOTHERAPY OR IN COMBINATION WITH A STATIN AFTER FAILING AN ADEQUATE TRIAL OF DIET THERAPY
- I-610 TREATMENT OF HEAVY MENSTRUAL BLEEDING FOR WOMEN WHO CHOOSE TO USE INTRAUTERINE CONTRACEPTION AS THEIR METHOD OF CONTRACEPTION
- I-611 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND POSTMENARCHAL GIRLS, AGES 10 TO 17 YEARS, WITH A RECOMMENDATION DOSING RANGE OF 5 TO 20 MG ONCE DAILY
- I-612 MICARDIS 80 MG FOR REDUCTION OF THE RISK OF MYOCARDIAL INFARCTION, STROKE, OR DEATH FROM CARDIOVASCULAR CAUSES IN PATIENTS 55 YEARS OF AGE OR OLDER AT HIGH RISK OF DEVELOPING MAJOR CARDIOVASCULAR EVENTS WHO ARE UNABLE TO TAKE ACE INHIBITORS
- I-613 MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE TO LESS THAN 18 YEARS OF AGE
- I-614 SHORT TERM TREATMENT OF EROSIIVE ESOPHAGITIS ASSOCIATED WITH GERD IN PEDIATRIC PATIENTS AGES FIVE YEARS AND OLDER
- I-615 MAINTENANCE TREATMENT OF BIPOLAR DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE
- I-616 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17 YEARS OF AGE
- I-617 MAINTENANCE OF GENERALIZED ANXIETY DISORDER (GAD)
- I-618 ADJUNCTIVE THERAPY IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- I-619 INTRAVENOUS CONTRAST ENHANCED COMPUTER TOMOGRAPHY OF THE HEAD AND BODY
- I-620 FOR USE IN COMBINATION WITH LETROZOLE FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER THAT OVEREXPRESSES THE HER2 RECEPTOR FOR WHOM HORMONAL THERAPY IS INDICATED
- I-621 PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE, BASED ON THE RESULTS OF JUSTIFICATION FOR THE USE OF STATINS IN PRIMARY PREVENTION; AN INTERVENTION TRIAL EVALUATING ROSUVASTATIN (JUPITER)
- I-622 ADJUNCTIVE THERAPY FOR PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS THIRTEEN YEARS OF AGE AND OLDER
- I-623 TREATMENT OF SIGNS AND SYMPTOMS OF ADVANCED IDIOPATHIC PARKINSON'S DISEASE
- I-624 MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST-LINE CHEMOTHERAPY
- I-625 PANCREATIC INSUFFICIENCY DUE TO CHRONIC PANCREATITIS AND PANCREATECTOMY
- I-626 RELIEF OF NASAL CONGESTION ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-627 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH & CML) IN CHRONIC PHASE.
- I-628 MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS
- I-629 ADJUNCTIVE THERAPY WITH EITHER LITHIUM OR VALPROATE FOR THE ACUTE TREATMENT OF MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-630 TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS (TS) WHO REQUIRE THERAPEUTIC INTERVENTION BUT ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION.
- I-631 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE FOLLOWING OPIOID DETOXIFICATION
- I-632 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN
- I-633 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE
- I-634 TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY
- I-635 ADJUNCTIVE TREATMENT WITH LONG-ACTING ORAL PSYCHOSTIMULANTS FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- I-636 TREATMENT OF EXTERNAL GENITAL AND PERIANAL WARTS/CONDYLOMA ACUMINATA IN PATIENTS 12 YEARS OR OLDER
- I-637 USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER

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I-638	FOR PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC.
I-639	TREATMENT OF PROGRESSIVE, WELL-DIFFERENTIATED PANCREATIC NEUROENDOCRINE TUMORS IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC DISEASE
I-640	MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
I-641	TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
I-642	TREATMENT OF ERECTILE DYSFUNCTION (ED) AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
I-643	REDUCE THE RISK OF STROKE AND SYSTEMIC EMBOLISM IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION.
I-644	MONOTHERAPY IN PATIENTS 13 YEARS OF AGE AND OLDER WITH PARTIAL SEIZURES WHO ARE RECEIVING THERAPY WITH A SINGLE ANTIEPILEPTIC DRUG (AED)
I-645	MAINTENANCE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS
I-646	SIGNS AND SYMPTOMS OF ADVANCED PARKINSON'S DISEASE (APD)
I-647	SIGNS AND SYMPTOMS OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)
I-648	TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
I-649	TREATMENT OF PATIENTS WITH ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
I-650	TREATMENT OF ADULTS WITH RENAL ANGIOMYOLIPOMA AND TUBEROUS SCLEROSIS COMPLEX (TSC), NOT REQUIRING IMMEDIATE SURGERY
I-651	MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH SPINAL CORD INJURY
I-652	MANAGEMENT OF POSTHERPETIC NEURALGIA
I-653	TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
I-654	MAGNETIC RESONANCE ANGIOGRAPHY (MRA) TO EVALUATE ADULTS WITH KNOWN OR SUSPECTED RENAL OR AORTO-ILIO-FEMORAL OCCLUSIVE VASCULAR DISEASE
I-655	TREATMENT OF POSTMENOPAUSAL WOMEN WITH ADVANCED HORMONE RECEPTOR-POSITIVE, HER2-NEGATIVE BREAST CANCER (ADVANCED HR+BC) IN COMBINATION WITH EXEMESTANE, AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
I-656	MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY (DPN) IN ADULTS WHEN A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME
I-657	PLAQUE PSORIASIS OF THE SCALP
I-658	FIRST-LINE TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER, IN COMBINATION WITH CARBOPLATIN, IN PATIENTS WHO ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION THERAPY
I-659	PLAQUE PSORIASIS OF THE BODY
I-660	TREATMENT OF DEEP VEIN THROMBOSIS
I-661	TREATMENT OF PULMONARY EMBOLISM
I-662	REDUCTION IN RISK FOR DEEP VEIN THROMBOSIS AND THE REDUCTION IN RISK FOR PULMONARY EMBOLISM
I-663	IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
I-664	TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY
I-665	TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS OF AGE AND OLDER WITH (NTDT)SYNDROMES AND WITH A (LIC) OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT (MG FE/G DW) AND SERUM FERRITIN GREATER THAN 300MCG/L
I-666	TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) IN COMBINATION WITH CHEMOTHERAPY
I-667	TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE
I-668	PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT

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- I-669 SCINTIGRAPHIC ASSESSMENT OF SYMPATHETIC INNERVATION OF THE MYOCARDIUM BY MEASUREMENT OF THE HEART TO MEDIASTINUM (H/M) RATIO OF RADIOACTIVITY UPTAKE IN PATIENTS WITH NYHA CLASS II OR CLASS III HEART FAILURE AND LVEF LESS THAN 35%
- I-670 TREATMENT OF OPIOID-INDUCED CONSTIPATION (OIC) IN ADULTS WITH CHRONIC, NON-CANCER PAIN
- I-671 FIRSTLINE TREATMENT OF PATIENTS WITH METASTATIC NON- SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21(L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-672 USE IN PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- I-673 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA/VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) CAUSED BY SUSCEPTIBLE ISOLATES OF S. AUREUS (INCLUDING METHICILLIN-SUSCEPTIBLE AND RESISTANT ISOLATES) WHEN ALTERNATIVE TREATMENTS ARE NOT SUITABLE
- I-674 TREATMENT OF PATIENTS WITH DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION) AS MONOTHERAPY AND AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE
- I-675 MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER
- I-676 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS, IN COMBINATION WITH GEMCITABINE
- I-677 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DTC) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT
- I-678 TRAMETINIB, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-679 RISK REDUCTION OF REBLEEDING OF GASTRIC OR DUODENAL ULCERS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS
- I-680 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-681 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT) WHICH MAY LEAD TO PULMONARY EMBOLISM (PE), IN ADULT PATIENTS WHO HAVE UNDERGONE HIP OR KNEE REPLACEMENT
- I-682 TREATMENT OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM (PE) IN PATIENTS WHO HAVE BEEN TREATED WITH A PARENTERAL ANTICOAGULANT FOR 5-10 DAYS
- I-683 TO REDUCE THE RISK OF RECURRENCE OF DVT AND PE IN PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED
- I-684 PREVENTION OF ACUTE NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING HIGHLY EMETOGENIC CANCER CHEMOTHERAPY IN PEDIATRIC PATIENTS AGED 1 MONTH TO LESS THAN 17 YEARS
- I-685 EXPANDED INDICATION OF RASAGILINE AS AN ADD-ON THERAPY TO STABLE DOSES OF DOPAMINE AGONISTS IN THE TREATMENT OF EARLY PARKINSON'S DISEASE
- I-686 INDICATED FOR THE TREATMENT OF DIABETIC MACULAR EDEMA IN PATIENTS WHO ARE PSEUDOPHAKIC OR ARE PHAKIC AND SCHEDULED FOR CATARACT SURGERY
- I-687 GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY
- I-688 GADAVIST IS INDICATED WITH MRI TO DETECT THE PRESENCE AND EXTENT OF MALIGNANT BREAST DISEASE
- I-689 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION
- I-690 INDICATED FOR THE TREATMENT OF DEEP VEIN THROMBOSIS (DVT)
- I-691 INDICATED TO REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM (PE) FOLLOWING INITIAL THERAPY
- I-692 INDICATED FOR MANAGEMENT OF OSTEOARTHRITIS PAIN.
- I-693 TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC)
- I-694 TREATMENT OF PATIENTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY

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I-695	REVISED INDICATION FOR BORTEZOMIB IN THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA
I-696	USE AS MONOTHERAPY IN THE TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGE 17 YEARS AND OLDER
I-697	FOR USE IN COMBINATION WITH SOFOSBUVIR FOR THE TREATMENT OF PATIENTS WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1 INFECTION
I-698	SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS
I-699	FOR TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA
I-700	TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S DISORDER (6-18 YEARS)
I-701	FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL-OR MODERATELY-DIFFERENTIATED, LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS) TO IMPROVE PROGRESSION FREE SURVIVAL
I-702	FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM MACROGLOBULINEMIA
I-703	MODERATE TO SEVERE BINGE EATING DISORDER (BED)
I-704	EXPANDED INDICATION TO INCLUDE PATIENTS WHO ARE VIROLOGICALLY-SUPPRESSED (HIV-1 RNA <50 COPIES/ML) ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TREATMENT FAILURE IN ORDER TO REPLACE THEIR CURRENT REGIMEN
I-705	TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE
I-706	EXPANDED INDICATION FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA
I-707	POMALYST, IN COMBINATION WITH DEXAMETHASONE, IS INDICATED FOR PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
I-708	DAILY TREATMENT OF ASTHMA IN PATIENTS AGED 18 YEARS AND OLDER
I-709	TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS
I-710	ADJUNCTIVE THERAPY FOR THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC (PG TC) SEIZURES IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE OR OLDER.
I-711	INCLUSION OF PEDIATRIC PATIENTS AGES 6 YRS AND OLDER FOR THE TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC ITP WHO HAVE HAD AN INSUFFICIENT RESPONSE TO CORTICOSTEROIDS, IMMUNOGLOBULINS, OR SPLENECTOMY.
I-712	EXPANDED INDICATION FOR USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE PRIOR LINES OF THERAPY
I-713	REVISIONS TO THE LABELING TO PERMIT THE USE OF ZUBSOLV AS INITIAL ("INDUCTION") TREATMENT OF OPIOID DEPENDENCE
I-714	EXTENDS THE 2011 APPROVAL OF BRILINTA FOR USE BEGINNING WITH ACS TO USE BEGINNING MORE REMOTE FROM MYOCARDIAL INFARCTION
I-715	FOR THE ADDITION OF THE INDICATION FOR MONOTHERAPY TREATMENT IN PARTIAL-ONSET SEIZURES IN ADULTS.
I-716	REVISED INDICATION TO INCLUDE LANGUAGE ABOUT THE BENEFITS OF USING LETAIRIS IN COMBINATION WITH TADALAFIL TO REDUCE THE RISK OF DISEASE PROGRESSION AND HOSPITALIZATION FOR WORSENING PAH AND TO IMPROVE EXERCISE ABILITY, BASED ON THE AMBITION STUDY
I-717	EXPANDED INDICATION TO INCLUDE THE TREATMENT OF CHRONIC HEPATITIS C GENOTYPE 4
I-718	EXPANDED INDICATION TO INCLUDE SUBJECTS INFECTED WITH CHRONIC HEPATITIS C, GENOTYPE 6 VIRUS INFECTION BASED UPON THE RESULTS OF THE ELECTRON- 2 STUDY
I-719	EXPANDED INDICATION TO INCLUDE THE TREATMENT OF SUBJECTS WITH GENOTYPE 5 CHRONIC HEPATITIS C VIRUS INFECTION BASED ON THE RESULTS FROM STUDY GS-US-337-119.
I-720	EXPANDED INDICATION TO INCLUDE TREATMENT OF GENOTYPE 4, CHRONIC HEPATITIS C VIRUS INFECTION BASED UPON THE RESULTS FROM STUDIES ION-4 AND GS-US-337-119.
I-721	TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN.
I-722	REVISED INDICATION FOR USE IN COMBINATION WITH DEXAMETHASONE OR WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR

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- REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY.
- I-723 AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE OR MORE LINES OF THERAPY
- I-724 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL DIFFERENTIATED, NON-FUNCTIONAL NEUROENDOCRINE TUMORS (NET) OF GI OR LUNG ORIGIN WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- I-725 TREATMENT OF HORMONE RECEPTOR (HR)-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION THERAPY WITH PALBOCICLIB AND FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY.
- I-726 EXPANSION OF THE PATIENT POPULATION TO INCLUDE PATIENTS WITH RECURRENCE OF HEPATITIS C VIRUS (HCV) GENOTYPE 1 OR 3 AFTER LIVER TRANSPLANTATION
- I-727 EXPANSION OF THE INDICATION TO INCLUDE TREATMENT OF SUBJECTS WITH GENOTYPE-1 CHRONIC HEPATITIS C VIRUS INFECTION, INCLUDING SUBJECTS WHO ARE CO-INFECTED WITH THE HUMAN IMMUNODEFICIENCY VIRUS (HIV-1) BASED ON THE RESULTS FROM THE ALLY-2 CLINICAL TRIAL
- I-728 EXPANDED INDICATION FOR USE IN ULTRASONOGRAPHY OF THE LIVER FOR CHARACTERIZATION OF FOCAL LIVER LESIONS IN ADULT AND PEDIATRIC PATIENTS
- I-729 PROVIDES FOR THE FRONTLINE INDICATION FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA
- I-730 NEW INDICATION FOR THE TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSING AFTER PLATINUM-BASED CHEMOTHERAPY
- I-731 FOR USE IN MAGNETIC RESONANCE ANGIOGRAPHY IN ADULT AND PEDIATRIC PATIENTS (INCLUDING TERM NEONATES) TO EVALUATE KNOWN OR SUSPECTED SUPRA-AORTIC OR RENAL ARTERY DISEASE
- I-732 TREATMENT OF PEDIATRIC PATIENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA TO REDUCE LDL-C, TOTAL C, NONHDL-C AND APOB AS AN ADJUNCT TO DIET, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS
- I-733 USE OF CANAGLIFLOZIN FOR INITIAL THERAPY IN COMBINATION WITH METFORMIN
- I-734 EXPANDED INDICATION FOR THE USE OF LENVIMA IN COMBINATION WITH EVEROLIMUS FOR THE TREATMENT OF PATIENTS WITH ADVANCED RCC FOLLOWING ONE PRIOR ANTI-ANGIOGENIC THERAPY.
- I-735 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH CANAGLIFLOZIN AND METFORMIN IS APPROPRIATE
- I-736 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- I-737 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL) WITH 17P DELETION
- I-738 REVISIONS TO THE INDICATIONS AND USAGE SECTION WITH RESPECT TO COMPLICATED INTRA-ABDOMINAL INFECTIONS
- I-739 TO REDUCE THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-740 EXPANDED INDICATION FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 YEARS AND OLDER TO INCLUDE THE G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, OR S549R MUTATION IN THE CFTR GENE
- I-741 TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY
- I-742 TREATMENT OF NODAL MARGINAL ZONE LYMPHOMA
- I-743 INFORMATION ADDED TO THE LABELING FOR THE ADDITION OF THE TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 4 (GT4) INFECTED PATIENTS WITH COMPENSATED CIRRHOSIS BASED ON RESULTS FROM STUDY M11-665
- I-744 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
- I-745 MEKINIST, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST.
- I-746 NEW INDICATION OF MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR I DISORDER IN ADULTS

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- I-747 FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED WITH HIGH-DOSE BUSULFAN AND CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENEIC HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION FOR PEDIATRIC PATIENTS WITH CLASS 3 BETA-THALASSEMIA
- I-748 TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS FIVE YEARS OF AGE AND OLDER
- I-749 MONOTHERAPY FOR THE TREATMENT OF HORMONE RECEPTOR (HR) POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY
- I-750 REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-751 TREATMENT OF TARDIVE DYSKINESIA
- I-752 CORONARY COMPUTED TOMOGRAPHY ANGIOGRAPHY (CCTA) TO ASSIST DIAGNOSTIC EVALUATION OF PATIENTS WITH SUSPECTED CORONARY ARTERY DISEASE
- I-753 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- I-754 TO REDUCE THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY WHEN USED FOR THE TREATMENT OF ADULTS WITH CARCINOID SYNDROME
- I-755 ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RENAL CELL CARCINOMA (RCC) FOLLOWING NEPHRECTOMY
- I-756 EXPANDED THE APPROVED INDICATION BY REMOVING THE RESTRICTION FOR USE ONLY IN PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- I-757 TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION
- I-758 FOR USE WITH RILPIVIRINE AS A COMPLETE REGIMEN TO REPLACE THE CURRENT ARV REGIMEN IN VIROLOGICALLY SUPPRESSED PATIENTS ON A STABLE ARV REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TX FAILURE OR KNOWN SUBSTITUTIONS ASSOC. WITH RESISTANCE TO EITHER ARV
- I-759 TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+CML)
- I-760 FOR THE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- I-761 TREATMENT OF ADULT PATIENTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO METHOTREXATE OR OTHER NON-BIOLOGIC DISEASE-MODIFYING ANTIRHEUMATIC DRUGS
- I-762 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE METASTATIC BREAST CANCER WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT OR METASTATIC SETTING
- I-763 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS HAVE NON-RESISTANT EPIDERMAL GROWTH FACTOR RECEPTOR MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-764 TREATMENT IN ADULT PATIENTS FOR IRRITABLE BOWEL SYNDROME WITH CONSTIPATION (IBS-C)
- I-765 ABIRATERONE ACETATE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC HIGH-RISK CASTRATION-SENSITIVE PROSTATE CANCER
- I-766 TREATMENT OF MINIMALLY TO MODERATELY THICK ACTINIC KERATOSIS OF THE UPPER EXTREMITIES IN CONJUNCTION WITH A BLUE LIGHT PHOTODYNAMIC THERAPY ILLUMINATOR
- I-767 TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON
- I-768 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- I-769 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- I-770 TREATMENT OF ACUTE OTITIS EXTERNA IN PATIENTS 6 MONTHS OF AGE AND OLDER DUE TO PSEUDOMONAS AERUGINOSA AND STAPHYLOCOCCUS AUREUS
- I-771 REVISION OF THE INDICATION SECTION OF THE PACKAGE INSERT REGARDING AN INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK TO PRODUCE POSTSURGICAL REGIONAL ANALGESIA

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-772 FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- I-773 FOR THE ADJUNCTIVE TREATMENT OF ADULT AND PEDIATRIC PATIENTS AGE 2 YEARS AND OLDER WITH TUBEROUS SCLEROSIS COMPLEX (TSC)-ASSOCIATED PARTIAL-ONSET SEIZURES
- I-774 TO ALLOW FOR FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS, AS DETECTED BY AN FDA APPROVED TEST
- I-775 REVISED INDICATION FOR FIXED-DOSE COMBINATION OF FLUTICASONE FUROATE, UMECLIDINIUM, AND VILANTEROL TO TREAT AIRFLOW OBSTRUCTION IN CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) AND TO REDUCE COPD EXACERBATIONS IN PTS WITH HISTORY OF EXACERBATIONS
- I-776 FIRSTLINE MAINTENANCE TX IN PTS W/ DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE, SOMATIC BRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CA WHO ARE IN COMPLETE OR PARTIAL RESPONSE TO FIRSTLINE PLATINUM-BASED CHEMOTHERAPY
- I-777 CO-ADMINISTRATION THERAPY OF MIRABEGRON WITH SOLIFENACIN SUCCINATE FOR TREATMENT OF OVERACTIVE BLADDER WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- I-778 DABRAFENIB IN COMBINATION WITH TRAMETINIB FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- I-779 USE OF TOLVAPTAN TO SLOW KIDNEY FUNCTION DECLINE IN ADULTS AT RISK OF RAPIDLY PROGRESSING AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)
- I-780 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC)
- I-781 DABRAFENIB IN COMBINATION WITH TRAMETINIB FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- I-782 REVISIONS TO INDICATION FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-783 EXPANDED INDICATION TO INCLUDE RIBOCICLIB WITH AN AROMATASE INHIBITOR IN PRE/PERIMENOPAUSAL WOMEN WITH HORMONE RECEPTOR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE-BASED THERAPY
- I-784 RIBOCICLIB WITH FULVESTRANT FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- I-785 TREATMENT OF PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- I-786 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
- I-787 FIRST-LINE TREATMENT OF PATIENTS WITH UNRESECTABLE HEPATOCELLULAR CARCINOMA (HCC)
- I-788 NEW INDICATION FOR CANAGLIFLOZIN TO REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, NONFATAL MYOCARDIAL INFARCTION AND NONFATAL STROKE) IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE (CVD)
- I-789 VENETOCLAX IN COMBO WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE FOR THE TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- I-790 USE OF FERRIC CITRATE FOR THE TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WITH CKD NOT ON DIALYSIS
- I-791 TREATMENT OF PEDIATRIC PATIENTS ONE YEAR OF AGE AND OLDER WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE (PH+) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN COMBINATION WITH CHEMOTHERAPY
- I-792 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-793	TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE
I-794	TREATMENT OF ADULT PATIENTS WITH METASTATIC GASTRIC OR GASTROESOPHAGEAL JUNCTION ADENOCARCINOMA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
I-795	VENETOCLAX IN COMBINATION WITH OBINUTUZUMAB IN PREVIOUSLY UNTREATED PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA
I-796	USED IN COMBINATION WITH A RITUXIMAB PRODUCT, ARE INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED FOLLICULAR LYMPHOMA (FL)
I-797	USED IN COMBINATION WITH A RITUXIMAB PRODUCT, ARE INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA (MZL)
I-798	TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION)
I-799	TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (GVHD) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
I-800	TREATMENT OF OCULAR INFLAMMATION FOLLOWING OPHTHALMIC SURGERY
I-801	USE IN CARDIAC MRI TO ASSESS MYOCARDIAL PERFUSION (STRESS, REST) AND LATE GADOLINIUM ENHANCEMENT IN ADULT PATIENTS WITH KNOWN OR SUSPECTED CORONARY ARTERY DISEASE (CAD)
I-802	TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
I-803	TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCETS DISEASE
I-804	EXPANDED INDICATION FOR USE AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
I-805	SLOW THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE
I-806	EXPANDED INDICATION FOR PTS WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML) ON A STABLE ARV REGIMEN WITH NO HX OF TX FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED W RESISTANCE TO DORAVIRINE, LAMIVUDINE OR TENOFOVIR DISOPROXIL FUMARATE
I-807	TREATMENT OF ADVANCED ENDOMETRIAL CARCINOMA THAT IS NOT MICROSATELLITE INSTABILITY-HIGH OR MISMATCH REPAIR DEFICIENT, WHO HAVE DISEASE PROGRESSION FOLLOWING PRIOR SYSTEMIC THERAPY AND ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION
I-808	TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC)
I-809	TO REDUCE THE RISK OF END-STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND DIABETIC NEPHROPATHY WITH ALBUMINURIA > 300 MG/DAY
I-810	PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING
I-811	TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 12 YEARS OF AGE OR OLDER, WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 48 HOURS AND ARE AT HIGH RISK OF DEVELOPING INFLUENZA-RELATED COMPLICATIONS
I-812	FOR USE IN AT RISK ADULTS AND ADOLESCENTS WEIGHING AT LEAST 35 KG FOR PRE-EXPOSURE PROPHYLAXIS TO REDUCE THE RISK OF HIV-1 INFECTION FROM SEXUAL ACQUISITION, EXCLUDING INDIVIDUALS AT RISK FROM RECEPTIVE VAGINAL SEX
I-813	TX OF ADULT PTS W/ ADV OVARIAN FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & ASSOCIATED W/ HRD DEFICIENCY POSITIVE STATUS DEFINED BY A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
I-814	TX OF ADV OVARIAN FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & ASSOCIATED W/ HRD DEFICIENCY DEFINED BY POSITIVE STATUS GENOMIC INSTABILITY & WHO HAVE PROGRESSED >6MO AFTER RESPONSE TO LAST PLATINUM-BASED CHEMO
I-815	TREATMENT OF COMMUNITY ACQUIRED BACTERIAL PNEUMONIA (CABP) CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA IN ADULTS
I-816	TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WHO ARE >= 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-817 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- I-818 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DISEASE HAS NOT PROGRESSED ON AT LEAST 16 WEEKS OF A FIRST-LINE PLATINUM-BASED CHEMOTHERAPY REGIMEN
- I-819 ADJUNCT TO MAX TOLERATED STATIN TX TO REDUCE RISK OF MI, STROKE, CORONARY REVASCULARIZATION, & UNSTABLE ANGINA REQUIRING HOSPITALIZATION IN ADULTS W/ ELEVATED TG LEVELS & ESTABLISHED CV DISEASE OR DIABETES MELLITUS & 2+ RISK FACTORS FOR CV DISEASE
- I-820 INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO DELAY DISEASE PROGRESSION
- I-821 TREATMENT OF CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESSSES WITHOUT MENINGOENCEPHALITIS AND/OR OCULAR DISSEMINATION IN PEDIATRIC PATIENTS YOUNGER THAN 4 MONTHS OF AGE
- I-822 REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, NON-FATAL MYOCARDIAL INFARCTION OR NON-FATAL STROKE) IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-823 USE IN COMBINATION WITH CAPECITABINE, FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED OR METASTATIC HER2-POSITIVE BREAST CANCER WHO HAVE RECEIVED TWO OR MORE PRIOR ANTI-HER2 BASED REGIMENS IN THE METASTATIC SETTING
- I-824 RIVAROXABAN IN COMBINATION WITH ASPIRIN, IS INDICATED TO REDUCE THE RISK OF MAJOR CV EVENTS (CV DEATH, MI, AND STROKE) IN PATIENTS WITH CHRONIC CORONARY ARTERY DISEASE (CAD) OR PERIPHERAL ARTERY DISEASE (PAD)
- I-825 TREATMENT FOR CHRONIC FIBROSING INTERSTITIAL LUNG DISEASES WITH A PROGRESSIVE PHENOTYPE
- I-826 ENCORAFENIB, IN COMBINATION WITH CETUXIMAB, FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC COLORECTAL CANCER WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, AFTER PRIOR THERAPY
- I-827 EXPANDED INDICATION FOR PATIENTS WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML) ON A STABLE ARV REGIMEN WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO DORAVIRINE
- I-828 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- I-829 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY (CPP)
- I-830 TREATMENT OF ADULT PATIENTS WITH A DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC)-ASSOCIATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE BEEN TREATED WITH ANDROGEN RECEPTOR-DIRECTED THERAPY AND A TAXANE-BASED CHEMOTHERAPY
- I-831 W/BEVACIZUMAB FOR MAINTENANCE TX OF ADULTS W/ADV. EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CA IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMO & CA ASSOCIATED W/ HOMOLOGOUS RECOMBINATION DEFICIENCY POSITIVE STATUS
- I-832 TX OF ADULT PTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC HOMOLOGOUS RECOMBINATION REPAIR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE
- I-833 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- I-834 TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE (NYHA CLASS II-IV) WITH REDUCED EJECTION FRACTION
- I-835 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL) WHOSE TUMORS ARE POSITIVE FOR AN EZH2 MUTATION AS DETECTED BY AN FDA-APPROVED TEST AND WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES
- I-836 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY (R/R) FOLLICULAR LYMPHOMA (FL) WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- I-837 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-838 TREATMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY (DPN) OF THE FEET
- I-839 TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPRESSED (HIV-1 RNA LESS THAN 50 COPIES PER ML) ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- I-840 TREATMENT OF SYMPTOMS IN ADULTS WITH MAJOR DEPRESSIVE DISORDER (MDD) WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR.
- I-841 TO REDUCE THE RISK OF HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE OR MULTIPLE CARDIOVASCULAR RISK FACTORS
- I-842 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY IN COMBINATION WITH DARATUMUMAB AND DEXAMETHASONE
- I-843 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS AGED 18 YEARS AND OLDER
- I-844 INDICATED IN PATIENTS 18 YEARS OF AGE AND OLDER FOR THE TREATMENT OF HOSPITAL ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) CAUSED BY THE FOLLOWING SUSCEPTIBLE GRAM-NEGATIVE MICROORGANISMS: ACINETOBACTER BAUMANNII COMPLEX, ESCHERICHIA COLI, ENTEROBACTER CLOACAE COMPLEX, KLEBSIELLA PNEUMONIAE, PSEUDOMONAS AERUGINOSA, AND SERRATIA MARCESCENS
- I-845 TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA-APPROVED TEST
- I-846 TREATMENT OF CATAPLEXY IN ADULT PATIENTS WITH NARCOLEPSY
- I-847 EXPANDED INDICATION OF THE TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA-APPROVED TEST
- I-848 REDUCE THE RISK OF STROKE IN PATIENTS WITH ACUTE ISCHEMIC STROKE (NIH STROKE SCALE SCORE \leq 5) OR HIGH-RISK TRANSIENT ISCHEMIC ATTACK (TIA)
- I-849 CHRONIC PHASE (CP) CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE OR INTOLERANCE TO AT LEAST TWO PRIOR KINASE INHIBITORS
- I-850 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PATIENTS 16 YEARS OF AGE AND OLDER
- I-851 TO REDUCE THE RISK OF A FIRST MYOCARDIAL INFARCTION (MI) OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE (CAD) AT HIGH RISK FOR SUCH EVENTS
- I-852 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ALK-POSITIVE
- I-853 INDICATION OF OSIMERTINIB AS ADJUVANT THERAPY AFTER TUMOR RESECTION IN ADULT PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- I-854 FOR THE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA, AS A FIRST-LINE TREATMENT IN COMBINATION WITH NIVOLUMAB
- I-855 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS 3 YEARS AND OLDER AND WEIGHING 35 KILOGRAMS OR MORE
- I-856 INDICATION FOR THE TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH INTERSTITIAL LUNG DISEASE TO IMPROVE EXERCISE ABILITY
- I-857 TO REDUCE THE RISK OF SUSTAINED EGFR DECLINE, END-STAGE KIDNEY DISEASE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH CHRONIC KIDNEY DISEASE AT RISK OF PROGRESSION
- I-858 FOR THE TOPICAL TREATMENT OF SCABIES INFESTATIONS IN ADULT AND PEDIATRIC PATIENTS 4 YEARS OF AGE AND OLDER
- I-859 TREATMENT OF PATIENTS WITH TRANSFUSIONAL IRON OVERLOAD DUE TO SICKLE CELL DISEASE OR OTHER ANEMIAS
- I-860 FOR THE TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS IN ADULT PATIENTS
- I-861 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER AND WEIGHING GREATER THAN 25 KG

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-862 TREATMENT OF VENOUS THROMBOEMBOLIC EVENTS (VTE) IN PEDIATRIC PATIENTS 8 TO LESS THAN 18 YEARS OF AGE WHO HAVE BEEN TREATED WITH A PARENTERAL ANTICOAGULANT FOR AT LEAST 5 DAYS AND TO REDUCE THE RISK OF RECURRENCE OF VTE IN PEDIATRIC PATIENTS 8 TO LESS THAN 18 YEARS OF AGE WHO HAVE BEEN PREVIOUSLY TREATED
- I-863 TREATMENT OF ADULT PATIENTS WITH ADVANCED SYSTEMIC MASTOCYTOSIS (ADVSM), INCLUDING PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM) AND SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN)
- I-864 TREATMENT OF ADULT PATIENTS WITH MAST CELL LEUKEMIA (MCL)
- I-865 FOR THE PREVENTIVE TREATMENT OF EPISODIC MIGRAINE IN ADULTS
- I-866 FOR THE TREATMENT OF TRICHOMONIASIS CAUSED BY TRICHOMONAS VAGINALIS IN ADULTS
- I-867 INDICATED TO REDUCE THE RISK OF MAJOR THROMBOTIC VASCULAR EVENTS (MYOCARDIAL INFARCTION, ISCHEMIC STROKE, ACUTE LIMB ISCHEMIA, AND MAJOR AMPUTATION OF VASCULAR ETIOLOGY) IN PATIENTS WITH PAD, INCLUDING PATIENTS WHO HAVE RECENTLY UNDERGONE A LOWER EXTREMITY REVASCLARIZATION PROCEDURE DUE TO SYMPTOMATIC PAD
- I-868 LENVATINIB IN COMBINATION WITH PEMBROLIZUMAB, IS INDICATED FOR THE FIRST-LINE TREATMENT OF ADULT PATIENTS WITH ADVANCED RENAL CELL CARCINOMA (RCC)
- I-869 REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE AND REDUCED EJECTION FRACTION
- I-870 INDICATED FOR THE TREATMENT OF IDIOPATHIC HYPERSOMNIA (IH) IN ADULTS
- I-871 TREATMENT OF ADULT PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA (WM)
- I-872 ADDITION OF THE INDICATION OF TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- I-873 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC DIFFERENTIATED THYROID CANCER (DTC) THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY OR INELIGIBLE
- I-874 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
- I-875 FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH AN IDH1 MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- I-876 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS
- I-877 INDICATION FOR THE USE OF ABEMACICLIB IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, NODE-POSITIVE, EARLY CANCER (EBC) AT HIGH RISK OF RECURRENCE AND A KI-67 SCORE>20% AS DETERMINED BY AN FDA APPROVED TEST
- I-878 ADDITION OF A NEW INDICATION FOR ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-879 TREATMENT OF ADULT PATIENTS WITH ACTIVE ANKYLOSING SPONDYLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS, TO THE PRESCRIBING INFORMATION
- I-880 TREATMENT OF ADULTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- I-881 FOR THE TREATMENT OF INVASIVE ASPERGILLOSIS IN PATIENTS 13 YEARS OF AGE AND OLDER
- I-882 INDICATED FOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I OR II DISORDER (BIPOLAR DEPRESSION) IN ADULTS, AS MONOTHERAPY AND AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE
- I-883 TREATMENT OF ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH REFRACTORY, MODERATE-TO-SEVERE ATOPIC DERMATITIS WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH OTHER SYSTEMIC DRUG PRODUCTS, INCLUDING BIOLOGICS, OR WHEN USE OF THOSE THERAPIES ARE INADVISABLE
- I-884 REVISIONS TO THE LABELING TO INCLUDE DATA FOR SUBJECTS WITH MILD TO MODERATE PLAQUE PSORIASIS, AND TO ALLOW FOR AN EXPANSION OF THE INDICATION
- I-885 FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA MUTATED HER2-NEGATIVE HIGH-RISK EARLY BREAST CANCER WHO HAVE PREVIOUSLY BEEN TREATED WITH NEOADJUVANT OR ADJUVANT CHEMOTHERAPY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-886	TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
I-887	INDICATION FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) IN PATIENTS WHO ARE 2 YEARS OF AGE AND OLDER
I-888	TREATMENT OF ADULTS WITH ACTIVE ANKYLOSING SPONDYLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
I-889	TREATMENT OF PEDIATRIC PATIENTS AGED ONE MONTH AND OLDER WITH NEWLY DIAGNOSED JUVENILE MYELOMONOCYTIC LEUKEMIA (JMML)
I-890	TREATMENT OF ADULT PATIENTS WITH SEVERE ALOPECIA AREATA
I-891	TREATMENT OF COVID-19 IN HOSPITALIZED ADULTS REQUIRING SUPPLEMENTAL OXYGEN, NONINVASIVE OR INVASIVE MECHANICAL VENTILATION, OR EXTRACORPOREAL MEMBRANE OXYGENATION (ECMO)
I-892	CHRONIC WEIGHT MANAGEMENT IN ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH BARDET-BIEDL SYNDROME (BBS)
I-893	IN COMBINATION WITH AZACITIDINE OR AS MONOTHERAPY FOR THE TREATMENT OF NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST IN ADULTS 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
I-894	DABRAFENIB IS INDICATED IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH UNRESECTABLE OR METASTATIC SOLID TUMORS WITH BRAF V600E MUTATION WHO HAVE PROGRESSED FOLLOWING PRIOR TREATMENT AND HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
I-895	TRAMETINIB IS INDICATED IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH UNRESECTABLE OR METASTATIC SOLID TUMORS WITH BRAF V600E MUTATION WHO HAVE PROGRESSED FOLLOWING PRIOR TREATMENT AND HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
I-896	INDICATED FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
I-897	TREATMENT OF ADULT AND PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH UNRESECTABLE, RECURRENT OR REFRACTORY INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK) POSITIVE
I-898	FOR THE MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS
I-899	TREATMENT OF ADULTS WITH RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS (MLNS) WITH FIBROBLAST GROWTH FACTOR RECEPTOR 1 (FGFR1) REARRANGEMENT
I-900	TREATMENT OF ADULT PATIENTS WITH METASTATIC HORMONESENSITIVE PROSTATE CANCER (MHSPC) IN COMBINATION WITH DOCETAXEL
I-901	EXPANDED INDICATION TO INCLUDE LOWERING OF PLASMA OXALATE LEVELS IN ADULT AND PEDIATRIC PATIENTS WITH PRIMARY HYPEROXALURIA TYPE 1 (PH1)
I-902	TREATMENT OF ADULT PATIENTS WITH HISTIOCYTIC NEOPLASMS
I-903	REDUCTION IN THE INCIDENCE OF RECURRENT VULVOVAGINAL CANDIDIASIS (RVVC) IN ADULT AND POST-MENARCHAL PEDIATRIC FEMALES
I-904	ADJUNCTIVE THERAPY TO ANTIDEPRESSANTS FOR THE TREATMENT OF MAJOR DEPRESSIVE DISORDER IN ADULTS
I-905	ADJUNCT FOR INTRAOPERATIVE IDENTIFICATION OF MALIGNANT AND NONMALIGNANT PULMONARY LESIONS IN ADULT PATIENTS WITH KNOWN OR SUSPECTED CANCER IN THE LUNG
I-906	TUCATINIB IN COMBINATION WITH TRASTUZUMAB FOR THE TREATMENT OF ADULT PATIENTS WITH RAS WILD-TYPE, HER2-POSITIVE, UNRESECTABLE OR METASTATIC COLORECTAL CANCER THAT HAS PROGRESSED FOLLOWING TREATMENT WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-BASED CHEMOTHERAPY
I-907	TO INCREASE BONE DENSITY IN MEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE (DEFINED AS A HISTORY OF OSTEOPOROTIC FRACTURE OR MULTIPLE RISK FACTORS FOR FRACTURE), OR PATIENTS WHO HAVE FAILED OR ARE INTOLERANT TO OTHER AVAILABLE OSTEOPOROSIS THERAPY
I-908	TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA WITH A BRAF V600 MUTATION WHO REQUIRE SYSTEMIC THERAPY
I-909	PREVENTIVE TREATMENT OF MIGRAINE IN ADULTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-910 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA WITH A BRAF V600E MUTATION WHO REQUIRE SYSTEMIC THERAPY
- I-911 FOR FLUORESCENCE IMAGING OF LYMPH NODES AND DELINEATION OF LYMPHATIC VESSELS DURING LYMPHATIC MAPPING IN ADULTS WITH BREAST CANCER FOR WHICH THIS PROCEDURE IS A COMPONENT OF INTRAOPERATIVE MANAGEMENT
- I-912 TREATMENT OF ADULT PATIENTS WITH INDOLENT SYSTEMIC MASTOCYTOSIS (ISM)
- I-913 TREATMENT OF AGITATION ASSOCIATED WITH DEMENTIA DUE TO ALZHEIMER'S DISEASE
- I-914 IN COMBINATION WITH ABIRATERONE AND PREDNISONE OR PREDNISOLONE FOR THE TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS BRCA-MUTATED (BRCA) METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC)
- I-915 TREATMENT OF IRON DEFICIENCY IN ADULT PATIENTS WITH HEART FAILURE AND NEW YORK HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY
- I-916 PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT RECIPIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE [D+/R-])
- I-917 TREATMENT OF HEAVY MENSTRUAL BLEEDING FOR UP TO 5 YEARS IN PATIENTS WHO CHOOSE INTRAUTERINE CONTRACEPTION AS THEIR METHOD OF CONTRACEPTION
- I-918 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 12 MONTHS OF AGE AND OLDER WITH ALAGILLE SYNDROME (ALGS)
- I-919 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE CROHN'S DISEASE WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- I-920 USE OF TALAZOPARIB IN COMBINATION WITH ENZALUTAMIDE FOR THE TREATMENT OF ADULT PATIENTS WITH HOMOLOGOUS RECOMBINATION REPAIR (HRR) GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC).
- I-921 TREATMENT OF FUNCTIONAL CONSTIPATION IN PEDIATRIC PATIENTS 6 TO 17 YEARS OF AGE
- I-922 USE OF EMPAGLIFLOZIN TO REDUCE THE RISK OF SUSTAINED DECLINE IN EGFR, END-STAGE KIDNEY DISEASE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION IN ADULTS WITH CHRONIC KIDNEY DISEASE AT RISK OF PROGRESSION
- I-923 FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+CML), NEWLY-DIAGNOSED OR RESISTANT OR INTOLERANT TO PRIOR THERAPY
- I-924 FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES (MDS) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- I-925 TREATMENT OF ADULTS WITH CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
- I-926 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (NMCSPC) WITH BIOCHEMICAL RECURRENCE AT HIGH RISK FOR METASTASIS
- I-928 ENCORAFENIB IN COMBINATION WITH BINIMETINIB, IS INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST

EXCLUSIVITY MISCELLANEOUS

- M-1 INFORMATION REGARDING SUPERIORITY CLAIM OVER RANITIDINE FOR DAY AND NIGHT HEARTBURN ADDED TO CLINICAL STUDIES SECTION
- M-2 APPROVAL FOR ADDITION TO CLINICAL PHARMACOLOGY SECTION OF THE LABEL REGARDING (1) IMPROVEMENT IN BONE MINERAL DENSITY IN CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENT PATIENTS AND (2) INCREASES IN SERUM ALKALINE PHOSPHATASE
- M-3 ADDITION OF EFFICACY AND SAFETY INFORMATION IN WHICH FOSAMAX WAS USED CONCOMITANTLY WITH ESTROGEN ALONE OR WITH ESTROGEN PLUS PROGESTIN
- M-4 CHANGES TO PEDIATRIC USE SECTION TO PROVIDE INFORMATION REGARDING SAFETY AND EFFICACY IN PEDIATRIC PATIENTS AS YOUNG AS 2 YEARS OLD
- M-5 INFORMATION REGARDING EFFECTS IN PATIENTS WITH ASTHMA ON CONCOMITANT INHALED CORTICOSTEROIDS IN CLINICAL PHARMACOLOGY SECTION

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-6 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH GLUCOPHAGE/GLYBURIDE COMBINATION ADDED TO CLINICAL PHARMACOLOGY AND DOSING AND ADMINISTRATION
- M-7 CLINICAL PHARMACOLOGY IN PEDIATRIC PATIENTS; DOSAGE AND ADMINISTRATION INFORMATION
- M-8 ADDITIONAL INFORMATION FOR THE USE OF SONATA CAPSULES FOR UP TO 5 WEEKS (35 NIGHTS) OF TREATMENT IN A CONTROLLED TRIAL SETTING
- M-9 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING OF TEXT AND TWO TABLES CONTAINING INFORMATION FOR THE PRESCRIBING PHYSICIAN ON BLOOD PRESSURE, HEART RATE, AND HEART RATE VARIABILITY
- M-10 INFORMATION REGARDING MAINTENANCE OF AN ANTIDEPRESSANT EFFECT UP TO 1 YEAR OF DOSING
- M-11 USE FOR LONG-TERM TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- M-12 NEW LANGUAGE FOR PEDIATRIC USE
- M-13 INFORMATION FROM PEDIATRIC STUDIES ADDED TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION
- M-14 ADDITIONAL CLINICAL TRIAL INFORMATION ADDED TO PEDIATRIC USE SUBSECTION
- M-15 LONGER TERM EFFICACY INFORMATION FOR RISPERIDONE IN THE TREATMENT OF SCHIZOPHRENIA
- M-16 CHANGE IN WORDING OF THE PEDIATRIC SECTION OF THE PACKAGE INSERT
- M-17 INFORMATION REGARDING USE OF ULTANE IN PEDIATRIC PATIENTS WITH CONGENITAL HEART DISEASE
- M-18 INFORMATION DENOTING THE EFFICACY OF REMERON IN MAINTAINING A RESPONSE IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER (MDD)
- M-19 INFORMATION REGARDING USE IN PEDIATRIC PATIENTS TWO YEARS OF AGE AND OLDER
- M-20 LABELING REVISIONS RELATED TO MCCUNE ALBRIGHT SYNDROME
- M-21 COMPARISON DATA ON THE ANTIHYPERTENSIVE EFFECTS OF ATACAND AND COZAAR
- M-22 CHANGE IN TIME TO ONSET OF ACTION
- M-23 INFORMATION REGARDING ELIMINATION ADDED TO CLINICAL PHARMACOLOGY, STUDY RESULTS IN PATIENTS WITH HEPATIC AND RENAL IMPAIRMENT
- M-24 INFORMATION ON RESULTS OF A LONG TERM LONGITUDINAL GROWTH STUDY AND PEDIATRIC SAFETY INFORMATION
- M-25 ADDITIONAL SAFETY AND PHARMACOKINETICS INFORMATION IN CHILDREN 6 MONTHS TO LESS THAN 6 YEARS OF AGE ADDED TO PACKAGE INSERT
- M-26 INCORPORATION OF INFORMATION CONTAINED IN THE PEG-INTRON PACKAGE INSERT INTO THE REBETOL PACKAGE INSERT AND MEDGUIDE-PEG-INTRON WAS APPROVED FOR USE IN COMBINATION WITH REBETOL FOR TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION ON 8/7/01
- M-27 INFORMATION DESCRIBING ASPIRIN ENDOSCOPY STUDY AND THE MAXIMUM RECOMMENDED DOSE FOR PATIENTS WITH MODERATE HEPATIC INSUFFICIENCY
- M-28 INFORMATION FROM A STUDY IN PEDIATRIC PATIENTS IN ASSOCIATION WITH A NEUROLOGICAL CONDITION
- M-29 LABELING CHANGES TO PROVIDE INFORMATION IN THE MANAGEMENT OF OBESITY IN ADOLESCENTS AGED 12 TO 16 YEARS
- M-30 CHANGES TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION SECTIONS OF LABELING CONCERNING USE OF LOTENSIN IN PEDIATRIC PATIENTS WITH HYPERTENSION
- M-31 INFORMATION FOR USE IN PEDIATRIC PATIENTS WITH CHRONIC KIDNEY DISEASE STAGE 5 (END-STAGE RENAL DISEASE)
- M-32 ADDITIONAL LANGUAGE TO CLINICAL PHARMACOLOGY AND CLINICAL STUDIES
- M-33 INFORMATION FOR USE OF ADVAIR DISKUS 100/50 IN CHILDREN 4 TO 11 YEARS OF AGE WITH ASTHMA
- M-34 EXPANDED INFORMATION TO PEDIATRIC USE SUBSECTION OF LABELING IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-35 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH ACTOS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, OR INSULIN ADDED TO CLINICAL PHARMACOLOGY

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- M-36 ADDITION OF INFORMATION TO CLINICAL STUDIES REGARDING PREVENTION OF CARDIOVASCULAR DISEASE
- M-37 INFORMATION ADDED TO THE LABELING THAT DETAILS INFORMATION RELATIVE TO STUDIES DONE IN PEDIATRIC POPULATIONS IN THE CLINICAL PHARMACOLOGY AND PEDIATRIC USE SUBSECTIONS
- M-38 SAFETY AND IOP-LOWERING EFFECTS OF TRUSOPT HAVE BEEN DEMONSTRATED IN PEDIATRIC PATIENTS IN A 3 MONTH, MULTI-CENTER DOUBLE MASKED ACTIVE-TREATMENT-CONTROLLED TRIAL
- M-39 FOR LABELING CHANGES BASED ON RESULTS OF THE SPD422-202 CLINICAL STUDY REPORT (CSR) SUBMITTED IN RESPONSE TO THE WRITTEN REQUEST
- M-40 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED IN PEDIATRIC PATIENTS WITH LEUKEMIA ADDED TO PRECAUTIONS
- M-41 REVISION TO THE PEDIATRIC USE PRECAUTIONS OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM THE CAPPS-169 STUDY ENTITLED "THE EFFECT OF ORTHO TRICYCLEN ON BONE MINERAL DENSITY IN PEDIATRIC SUBJECTS WITH ANOREXIA NERVOSA"
- M-42 ADDITION OF A GERIATRIC USE SUBSECTION TO THE PRECAUTIONS SECTION OF THE PACKAGE INSERT AND GERIATRIC DOSING INFORMATION
- M-43 INCLUSION OF RESULTS OF STUDY "PLACEBO-CONTROLLED STUDY TO EVALUATE SAFETY AND PILOT EFFICACY OF ILOPROST AS ADD ON THERAPY WITH BOSENTAN IN SUBJECTS WITH PULMONARY ARTERIAL HYPERTENSION"
- M-44 CLINICAL INFORMATION ADDED TO THE PEDIATRIC USE SUBSECTION OF PRECAUTIONS REGARDING THE USE OF NOVOLOG IN ADOLESCENTS WITH TYPE I DIABETES AGE 6 TO 18
- M-45 INFORMATION ADDED TO CLINICAL TRIALS SECTION OF LABELING, "EFFECTS OF HUMATROPE TREATMENT IN ADULTS WITH GROWTH HORMONE DEFICIENCY"
- M-46 PROVISION OF RESULTS OF STUDY AND PROPOSED REVISIONS TO PACKAGE INSERT SEE SECTION ON CARDIAC ELECTROPHYSIOLOGY
- M-47 PROVIDES FOR USE OF ANTARA WITHOUT REGARD TO MEALS
- M-48 CHANGES TO THE LABELING DESCRIBING THE RESULTS OF A STUDY OF THE USE OF NOVOLOG MIX 70/30 WITH ORAL ANTIDIABETIC AGENTS IN PATIENTS WITH TYPE 2 DIABETES
- M-49 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING EFFECT OF SINGULAIR ON GROWTH RATES IN PREPUBERTAL CHILDREN
- M-50 NEW INFO TO THE CLINICAL STUDIES, ADULT GROWTH HORMONE DEFICIENCY (GHD) SUBSECTION OF THE NUTROPIN AQ PACKAGE INSERT DESCRIBING THE EFFECTS OF SOMATROPIN ON VISCERAL ADIPOSE TISSUE IN THE ADULT GROWTH HORMONE DEFICIENT PATIENT POPULATION
- M-51 INFORMATION ADDED TO LABELING REGARDING OSTEOGENESIS IMPERFECTA STUDY
- M-52 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY/CLINICAL STUDIES SECTION REGARDING THE USE OF RISEDRONATE ADMINISTERED ONCE A WEEK IN THE PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- M-53 FOR LABELING CHANGES TO THE QUALITY OF LIFE (QOL) STATEMENT IN THE APPROVED PACKAGE INSERT
- M-54 INFORMATION FROM PEDIATRIC STUDIES ADDED TO LABEL
- M-55 INFORMATION ON RESULTS OF A STUDY OF THE USE OF SANDOSTATIN LAR DEPOT IN PEDIATRIC PATIENTS WITH HYPOTHALAMIC OBESITY.
- M-56 INFORMATION ADDED TO CLINICAL TRIAL SECTION WITH INFORMATION ON "GEMINI" TRIAL
- M-57 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING THE PHARMACOKINETICS OF EZETIMIBE IN ASIAN SUBJECTS
- M-58 CHANGES TO THE CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, VYTORIN SUBSECTION OF THE PACKAGE INSERT TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR AN ATORVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PRMTRS
- M-59 RESULTS OF THE T20-310 STUDY WHICH EVALUATED THE PHARMACOKINETICS, SAFETY, AND ANTIVIRAL ACTIVITY OF FUZEON IN TREATMENT EXPERIENCED PEDIATRIC SUBJECTS AND ADOLESCENTS WAS ADDED TO THE PEDIATRIC SUBSECTION OF PRECAUTIONS
- M-60 CHANGES TO CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR A ROSUVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PARAMETERS IN PATIENTS WTH HYPERCHOLESTEROLEMIA
- M-61 REVISIONS TO LABELING BASED ON DATA SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

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- M-62 CLINICAL INFORMATION FROM ONE CLINICAL STUDY INVESTIGATING THE USE OF AVANDAMET PLUS INSULIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL WITH PREVIOUS ANTI-DIABETIC THERAPIES
- M-63 DETAILED INFORMATION ON AN INCONCLUSIVE PEDIATRIC STUDY
- M-64 CHANGES TO CLINICAL PHARMACOLOGY DETAILING STUDY RESULTS
- M-65 ADDITION OF INFORMATION TO LABEL TO INCLUDE INFORMATION REGARDING USE IN PATIENTS WITH HIV-ASSOCIATED ADIPOSE REDISTRIBUTION SYNDROME (HARS)
- M-66 USE IN SPECIFIC POPULATIONS - PATIENTS WITH CONCOMITANT ILLNESS SUBSECTION OF THE LABELING REGARDING USE OF STRATTERA IN PATIENTS WITH ADHD WHO HAVE COMORBID TIC DISORDER
- M-67 INDICATION EXPANDED TO INCLUDE PATIENTS ON PERITONEAL DIALYSIS
- M-68 DESCRIPTION OF RESULTS OF STUDY OF INITIAL THERAPY IN COMBINATION WITH METFORMIN WHEN DIET AND EXERCISE DO NOT PROVIDE GLYCEMIC CONTROL
- M-69 RESULTS OF STUDY OF COMBINATION THERAPY AND NON-INFERIORITY STUDY
- M-70 PROVISION OF INFORMATION OF THE RESULTS OF A PHASE 2 RANDOMIZED TRIAL OF SPRYCEL 70MG TWICE DAILY OR IMATINIB 800MG DAILY
- M-71 REVISIONS TO PROVIDE FOR RESULTS OF MAINTENANCE DATA IN ADULT PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- M-72 INFORMATION ABOUT USE OF INSPRA (EPLERENONE) FOR HYPERTENSION IN PEDIATRIC PATIENTS
- M-73 NEW INFORMATION ADDED REGARDING THE TUMOR SHRINKING POTENTIAL OF SANDOSTATIN LAR DEPOT INJECTION ON GH - SECRETING PITUITARY ADENOMAS
- M-74 REVISIONS TO CLINICAL STUDIES - CHILDREN AND ADOLESCENTS BASED ON CLINICAL TRIAL DATA TO SUPPORT A DURATION OF ACTION CLAIM UP TO 12 HOURS
- M-75 PROVISION FOR USE OF ARGATROBAN IN CERTAIN PEDIATRIC PATIENTS WITH HEPARIN-INDUCED THROMBOCYTOPENIA (HIT) OR HEPARIN-INDUCED THROMBOCYTOPENIA WITH THROMBOSIS (HITTS)
- M-76 REMOVAL OF SCREEN REQUIREMENT IN PTS WITH G6PD DEFICIENCY PRIOR TO INITIATING ACZONE TREATMENT; REMOVAL OF BLOOD COUNT & RETICULOCYTE MONITORING DURING TREATMENT IN G6PD DEFICIENT PTS AND IN PATIENTS WITH HISTORY OF ANEMIA
- M-77 USE IN COMBINATION WITH THE NEW AKTILITE CL128 LAMP FOR THE TREATMENT OF THIN AND MODERATELY THICK, NON-HYPERKERATOTIC, NON-PIGMENTED ACTINIC KERATOSES OF THE FACE AND SCALP IN IMMUNOCOMPETENT PATIENTS
- M-78 CLINICAL TRIAL INFO ON USE OF STRATTERA IN PATIENTS WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) AND COMORBID ANXIETY DISORDER WITHOUT CAUSING WORSENING OF ANXIETY
- M-79 LABELING REVISIONS RELATED TO SMOKING AND ERLOTINIB EXPOSURE
- M-80 ADDITIONAL TIME POINT OF 30 MINUTES (0.5 HOUR) IN CHILDREN AGED 6-12 YEARS WITH A DIAGNOSIS OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD)
- M-81 ADDITIONAL INFO FOR PEDIATRIC USE FOR CASODEX (STUDIED IN COMBINATION WITH ARIMIDEX) IN THE PEDIATRIC POPULATION, SPECIFICALLY BOYS WITH FAMILIAL MALE-LIMITED PRECOCIOUS PUBERTY (TESTOTOXICOSIS)
- M-82 LABELING REVISIONS RELATED TO CLINICAL STUDIES
- M-83 ADDITIONAL INFORMATION ADDED TO LABELING REGARDING ESTABLISHMENT OF EFFICACY IN ADDITIONAL CLINICAL TRIALS AND ONE MAINTENANCE TRIAL
- M-84 STUDY INFORMATION ADDED TO LABEL REGARDING BONE MINERAL DENSITY
- M-85 INFORMATION ADDED TO LABELING REGARDING USE OF PREVACID IN PATIENTS LESS THAN 1 YEAR WITH SYMPTOMATIC GERD
- M-86 LABELING CHANGES SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN REQUEST FOR INFANTS AGES BIRTH TO 11 MONTH INCLUSIVE REFLECTING LACK OF EFFICACY FOR GERD INDICATION FOR THIS PATIENT POPULATION
- M-87 INCLUSION OF RESULTS FROM TWO DRUG INTERACTION STUDIES WITH LIPITOR AND CRESTOR IN CLINICAL PHARMACOLOGY SECTION
- M-88 ADDITION OF INFORMATION REGARDING ABUSE POTENTIAL OF CONCERTA VERSUS IMMEDIATE-RELEASE METHYLPHENIDATE

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EXCLUSIVITY MISCELLANEOUS

- M-89 PROVIDES FOR REVISIONS TO MULTIPLE SECTIONS OF THE PACKAGE INSERT TO REFLECT RESULTS OF CLINICAL TRIALS 205.235 (UPLIFT) AND 205.266 (VA STUDY) IN SUPPORT OF EXACERBATION CLAIM
- M-90 LABELING CHANGES BASED ON DATA FROM CLINICAL STUDIES NV20235 AND NV20236 STUDIES OF SEASONAL PROPHYLAXIS OF INFLUENZA IN IMMUNOCOMPROMISED PATIENTS AND CHILDREN AGES 1-12
- M-91 UPDATED LABELING BASED UPON STUDY: A SINGLE-DOSE, SINGLE-BLIND, PLACEBO-AND MOXIFLOXACIN-CONTROLLED 2-PERIOD, RANDOMIZED, CROSSOVER, 3RD PERIOD SEQUENTIAL STUDY OF SIDE EFFECTS OF TEMSIROLIMUS ON CARDIAC REPOLARIZATION IN HEALTHY SUBJECTS
- M-92 UPDATES TO THE PACKAGE INSERT BASED UPON THE TRIAL ENTITLED "A PHASE I PHARMACOKINETIC AND PHARMACODYNAMIC STUDY OF TEMSIROLIMUS IN PATIENTS WITH ADVANCED MALIGNANCIES AND NORMAL AND IMPAIRED LIVER FUNCTION"
- M-93 EXPANSION OF LABELING TO INCLUDE INFORMATION ON SAFETY AND EFFICACY OF CREON IN PATIENTS AGES 7 YEARS THROUGH 11 YEARS WITH PANCREATIC EXOCRINE INSUFFICIENCY DUE TO CYSTIC FIBROSIS
- M-94 INFO ADDED TO LABEL RELATED TO NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE (PH+) CHRONIC MYELOID LEUKEMIA IC CHRONIC PHASE
- M-95 INFORMATION FOR TREATMENT OF CHRONIC HEPATITIS B (CHB) IN ADULT PATIENTS WITH DECOMPENSATED LIVER DISEASE BASED ON DATA FROM CLINICAL TRIAL GS-US-174-0108
- M-96 UPDATED INFORMATION IN THE CLINICAL STUDIES SECTION RELATED TO THE LOSS AND RECOVERY OF BONE MINERAL DENSITY IN ADOLESCENT GIRLS DURING AND FOLLOWING THE USE OF DEPO-PROVERA CONTRACEPTIVE INJECTION
- M-97 LABELING CHANGES IN RESPONSE TO PEDIATRIC STUDIES - NOT INDICATED FOR USE IN PEDIATRIC POPULATION
- M-98 NEW INFORMATION FROM A STUDY WHICH EVALUATED THE SAFETY AND EFFICACY OF FAMVIR IN TREATING RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT BLACK/AFRICAN AMERICAN SUBJECTS.
- M-99 ADDITION OF FINDINGS FROM A SINGLE PEDIATRIC CLINICAL TRIAL (P04292) OF NASONEX NASAL SPRAY IN THE TREATMENT OF NASAL POLYPS IN PATIENTS 6 TO <18 YEARS OF AGE TO THE PACKAGE INSERT.
- M-100 INFORMATION ADDED TO LABEL BASED UPON COMPLETED CLINICAL TRIAL REPORTS
- M-101 INCLUSION OF DATA FROM AN ADDITIONAL 19 SUBJECTS WITH HYPERCALCEMIA FROM PARATHYROID CARCINOMA TO THE INFORMATION CURRENTLY PRESENTED IN THE LABEL
- M-102 INFORMATION FROM PEDIATRIC STUDY REPORT ML16633, "INTRAVENOUS GRANISETRON (KYTRIL) IN THE PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING (PONV) IN PEDIATRIC SUBJECTS UNDERGOING TONSILLECTOMY OR ADENOTONSILLECTOMY."
- M-103 SAFETY, EFFICACY AND PHARMACOKINETIC INFO FOR FASLODEX IN THE PEDIATRIC POPULATION, SPECIFICALLY FOR GIRLS WITH PROGRESSIVE PRECOCIOUS PUBERTY ASSOCIATED WITH MCCUNE-ALBRIGHT SYNDROME ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING
- M-104 INFORMATION ADDED TO DOSING AND ADMINISTRATION REGARDING A 26 WEEK STUDY
- M-105 NEW LANGUAGE ADDED TO CLINICAL STUDIES REGARDING USE IN SMOKERS WITH CARDIOVASCULAR DISEASE, CHRONIC OBSTRUCTIVE PULMONARY DISEASE, AND USE ACCORDING TO AN ALTERNATIVE SET OF DIRECTIONS FOR SETTING A QUIT DATE
- M-106 ADDITION OF THE T1-WEIGHTED GD-ENHANCED LESION EFFICACY VARIABLE IN THE CLINICAL STUDIES SECTION 14 OF THE PACKAGE INSERT
- M-107 INFORMATION TO THE CLINICAL STUDIES SECTION OF THE LUPRON DEPOT-PED 1-MONTH BASED UPON THE PHASE 3/4 COMPLETED CLINICAL STUDY REPORT FOR STUDY M90-516 ENTITLED "STUDY OF LUPRON DEPOT IN THE TREATMENT OF CENTRAL PRECOCIOUS PUBERTY".
- M-108 CHANGES ARE BASED ON RESULTS FROM STUDY CV181057
- M-110 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL
- M-111 LABELING CHANGES BASED ON STUDY HW80-EW-GWCI ENTITLED A PLACEBO AND POSITIVE CONTROLLED STUDY OF THE ELECTROPHYSIOLOGICAL EFFECTS OF A SINGLE 10 MCG DOSE OF EXENATIDE ON THE 12 LEAD ELECTROCARDIOGRAM QT INTERVAL IN HEALTHY SUBJECTS
- M-112 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO ADD INFORMATION FROM A PEDIATRIC STUDY IN PATIENTS AGED 12 YEARS TO LESS THAN 18 YEARS OF AGE WITH RECURRENT HERPES LABIALIS

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M-113 LABELING CHANGES BASED ON STUDY H80-US-GWCO ENTITLED A RANDOMIZED TRIAL COMPARING EXENATIDE WITH PLACEBO IN SUBJECTS WITH TYPE 2 DIABETES ON INSULIN GLARGINE WITH OR WITHOUT ORAL ANTIHYPERGLYCEMIC MEDICATIONS

M-114 CHANGES IN SECTION 14 OF THE PACKAGE INSERT TO INCLUDE DATA FROM THE SWITCHMRK STUDIES (SWITCH OF SUPPRESSED SUBJECTS FROM LOPINAVIR/RITONAVIR TO RALTEGRAVIR)

M-115 REVISIONS TO THE PI BASED ON RESULTS FROM STUDY NN2211-1842, ENTITLED THE EFFECT OF INSULIN DETEMIR IN COMBINATION WITH LIRAGLUTIDE AND METFORMIN COMPARED TO LIRAGLUTIDE AND METFORMIN IN SUBJECTS WITH TYPE 2 DIABETES

M-116 LABELING CHANGES BASED ON RESULTS FROM CLINICAL STUDY 01-06-TL-OPIMET-008

M-117 ADDITION OF RESULTS OF PEDIATRIC TRIAL TO LABEL

M-118 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.36

M-119 LABELING CHANGES REGARDING MISSED DOSES

M-120 CHANGES TO CLINICAL TRIALS DETAILING STUDY RESULTS

M-121 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.43

M-122 LABELING CHANGES TO INCLUDE THE RESULTS OF THE PARAMOUNT TRIAL

M-123 UPDATED RESULTS OF OVERALL SURVIVAL FROM 'CONFIRM' STUDY

M-124 LONG TERM SAFETY AND EFFICACY DATA FROM STUDY CLDT600A2303 FOR SUBJECTS PREVIOUSLY ENROLLED IN THE ORIGINAL TWO YEAR GLOBE (NV-02B-007/CLDT600A2302) AND NV02B-015 STUDIES WHO CONTINUED TELBIVUDINE TREATMENT FOR UP TO 208 WEEKS

M-125 LABELING CHANGES TO INCLUDE LACK OF EFFICACY IN CHILDREN 6 MONTHS TO 4 YEARS OF AGE

M-126 UPDATES TO THE CLINICAL STUDIES SECTION 14, OF THE PACKAGE INSERT (PI), WITH THE RESULTS OF CLINICAL TRIAL P06086

M-127 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO REFLECT THE RESULTS FROM CLINICAL STUDY C-10-004

M-128 CLINICAL TRIAL STUDY RESULTS

M-129 RESULTS OF A CLINICAL STUDY REPORT WHICH ASSESSES THE SAFETY AND EFFICACY IN CHILDREN AGES 6 TO 12 YEARS OF AGE

M-130 ADDITION OF INFORMATION ON LONG-TERM TREATMENT WITH VPRIV IN THE CLINICAL TRIALS SECTION OF THE PACKAGE INSERT

M-131 INFORMATION FROM STUDIES CONDUCTED IN PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED NON-DISSEMINATED DIFFUSED INTRINSIC BRAINSTEM GLIOMAS

M-132 REVISIONS TO THE CLINICAL TRIALS SECTION IN THE INOMAX LABEL TO REFLECT RESULTS FROM THE PEDIATRIC STUDY REPORTS

M-133 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF SILDENAFIL TO BOSENTAN THERAPY

M-134 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED WITH SAXAGLIPTIN IN COMBINATION WITH METFORMIN AND A SULFONYLUREA ADDED TO THE LABELING

M-135 ADDITION OF INFORMATION TO THE CLINICAL STUDIES - RADIOGRAPHIC RESPONSE SECTION OF THE PACKAGE INSERT

M-136 ADDITIONAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING REGARDING POST-OPERATIVE NAUSEA AND VOMITING STUDIES IN PEDIATRIC PATIENTS

M-137 LABELING REVISIONS RESULTING FROM A MAINTENANCE TRIAL IN PEDIATRIC PATIENTS WITH IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER

M-138 INFORMATION ADDED TO THE 8.4 PEDIATRIC USE SECTION ON THE USE OF MEMANTINE IN CHILDREN AGES 6-12 YEARS WITH AUTISM SPECTRUM DISORDER

M-139 INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE PACKAGE INSERT REGARDING RETREATMENT WITH VELCADE FOR PATIENTS WITH MULTIPLE MYELOMA

M-140 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING USE OF LATISSE IN PATIENTS WHO WERE POST-CHEMOTHERAPY OR HAD ALOPECIA AREATA, AND ADOLESCENTS WHO HAD HYPERTRICHOSIS WITH NO ASSOCIATED MEDICAL CONDITION

M-141 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING TO INCORPORATE STUDY RESULTS FOR TREATMENT OF MAJOR DEPRESSIVE DISORDER IN ADOLESCENTS (AGES 12-17)

M-142 ADDITIONS TO THE LABELING DESCRIBING RESULTS FROM STUDY H6P-MC-HDAY

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- M-143 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN PATIENTS WITH CURRENT OR PAST HISTORY OF MAJOR DEPRESSIVE DISORDER
- M-144 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN PATIENTS WHO HAD BEEN PREVIOUSLY TREATED WITH VARENICLINE
- M-145 ADDITION OF INFORMATION ABOUT LONG-TERM TREATMENT OF PULMONARY ARTERIAL HYPERTENSION TO THE CLINICAL STUDIES SECTION OF THE LABELING
- M-146 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION ON INITIAL COMBINATION THERAPY WITH LINAGLIPTIN AND METFORMIN VS. LINAGLIPTIN MONOTHERAPY IN TREATMENT NAIVE PATIENTS
- M-147 OTC USE FOR TEMPORARY RELIEF OF OCULAR SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES
- M-148 LABELING CHANGES BASED ON STUDY H80-EW-GWDM
- M-149 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE MONOTHERAPY FOR ADHD
- M-150 ADDITION OF THE RESULTS OF A CONTROLLED CLINICAL STUDY TREATING ADULT PATIENTS WITH SCHIZOPHRENIA EXPERIENCING AN ACUTE RELAPSE
- M-151 REVISIONS TO THE LABELING BASED ON THE OUTCOMES OF PEDIATRIC STUDIES CONDUCTED TO ASSESS THE SAFETY AND EFFICACY OF XOPENEX IN SUBJECTS LESS THAN 6 YEARS OF AGE
- M-152 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION OF THE LABELING REGARDING A SAFETY STUDY IN PEDIATRIC SUBJECTS AGES 6 MONTHS TO 4 YEARS OF AGE WITH AN ACTIVE HEAD LICE INFESTATION
- M-153 ADDITION OF INFORMATION REGARDING THE INTRANASAL ABUSE POTENTIAL OF OXYCONTIN
- M-154 UPDATE TO THE LABELING TO REFLECT THE RESULTS OF A LONG-TERM MAINTENANCE TREATMENT STUDY OF ADHD IN CHILDREN AND ADOLESCENTS AGES 6-17.
- M-155 ADDITION OF CLINICAL FINDINGS FROM AN OBSERVATIONAL STUDY IN A PEDIATRIC AGE GROUP GREATER THAN 2 MONTHS TO 18 YEARS IN SECTION 8.4 PEDIATRIC USE OF THE PACKAGE INSERT
- M-156 UPDATE TO THE LABELING WITH INFORMATION REGARDING A CLINICAL TRIAL IN CHILDREN LESS THAN 4 YEARS OF AGE.
- M-157 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN 10MG ONCE DAILY IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND SULFONYLUREA
- M-158 UPDATES TO THE LABELING TO REFLECT SAFETY RESULTS FROM CLINICAL TRIALS IN SCHIZOPHRENIA ADOLESCENT PATIENTS AGED 12 TO 17 YEARS
- M-159 ADDITION OF PED SAFETY INFORMATION DERIVED FROM A MAINTENANCE TREATMENT STUDY OF BIPOLAR 1 DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES IN PATIENTS (> THAN OR = TO 13 YRS OF AGE) TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
- M-160 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND ACTIVE-CONTROLLED STUDY COMPARING EMPAGLIFLOZIN TO GLIMEPIRIDE IN PATIENTS WITH TYPE 2 DIABETES AND INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN TREATMENT
- M-161 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY OF EMPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS AND INSUFFICIENT GLYCEMIC CONTROL ON A MULTIPLE DAILY INJECTION INSULIN REGIMEN ALONE OR WITH METFORMIN
- M-162 INCLUSION OF EFFICACY AND SAFETY DATA TO THE PRESCRIBING INFORMATION OF BYDUREON BASED ON STUDY GWDE
- M-163 INFORMATION ADDED TO THE LABELING REGARDING PREVIOUSLY UNTREATED ALK-POSITIVE METASTATIC NON SMALL CELL LUNG CANCER (NSCLC)
- M-164 REVISES THE CLINICAL TRIALS SECTION OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM STUDY E7273-G000-401 ENTITLED "PHASE IV RANDOMIZED STUDY OF TWO DOSE LEVELS OF TARGRETIN CAPSULES IN SUBJECTS WITH REFRACTORY CUTANEOUS T-CELL LYMPHOMA"
- M-165 PROVIDES FOR UPDATES TO THE PEDIATRIC USE SECTION BASED ON THE PEDIATRIC STUDY REPORT ENTITLED, "A PHASE II PILOT TRIAL OF BORTEZOMIB IN COMBINATION WITH INTENSIVE RE-INDUCTION THERAPY IN CHILDREN WITH RELAPSED ACUTE LYMPHOBLASTIC LYMPHOMA (LL) "

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- M-166 UPDATE TO LABELING WITH WEEK 48 RESULTS FROM VIKING-4 IN ANTIRETROVIRAL THERAPY (ART) - EXPERIENCED INTEGRASE STRAND TRANSFER INHIBITOR (INSTI) - RESISTANT SUBJECTS
- M-167 APPROVED FOR REVISIONS TO THE LABELING BASED ON THE CLINICAL STUDY ENTITLED "BRONCHOPULMONARY DYSPLASIA (BPD) IN PRETERM INFANTS REQUIRING MECHANICAL VENTILATION OR POSITIVE PRESSURE SUPPORT ON DAYS 5 TO 14 AFTER BIRTH".
- M-168 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE RE-NOVATE AND RE-NOVATE LL STUDIES (PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM FOLLOWING HIP REPLACEMENT SURGERY)
- M-169 UPDATES TO LABELING DESCRIBING RESPONSE TO A REPEAT COURSE OF PICATO GEL 0.015% ON THE FACE OR SCALP IF AN INCOMPLETE RESPONSE IS OBSERVED AT A FOLLOW-UP EXAMINATION.
- M-170 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION REGARDING USE FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- M-171 UPDATES TO LABELING WITH RESULTS TO THE TIGER CLINICAL TRIAL
- M-172 UPDATES TO THE CLINICAL TRIALS SECTION OF THE LABELING TO INCLUDE RESULTS OF STUDIES PERFORMED TO EVALUATE THE BENEFIT OF ADDING INCRUSE ELLIPTA TO PATIENTS WHO ARE ON BACKGROUND THERAPY WITH BREO ELLIPTA AND ADVAIR DISKUS
- M-173 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING DESCRIBING THE EFFECTS OF STIOLTO RESPIMAT ON COPD PATIENTS
- M-174 INFORMATION ADDED TO CLINICAL STUDIES SECTION OF THE LABELING REGARDING INITIAL COMBINATION THERAPY OF EMPAGLIFLOZIN WITH METFORMIN
- M-175 INFORMATION ADDED TO THE LABELING DESCRIBING SAVOR, A PHASE IV TRIAL EVALUATING THE EFFECT OF SAXAGLIPTIN ON THE INCIDENCE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION OR ISCHAEMIC STROKE IN PATIENTS WITH TYPE 2 DIABETES
- M-176 INFORMATION ADDED TO THE LABELING DESCRIBING TRIAL NN2211-3916, A TRIAL EVALUATING THE SAFETY AND EFFICACY OF LIRAGLUTIDE IN SUBJECTS WITH TYPE 2 DIABETES AND MODERATE RENAL IMPAIRMENT
- M-177 INFORMATION ADDED TO THE LABELING DESCRIBING EXAMINE, A TRIAL EVALUATING CARDIOVASCULAR ISCHEMIC RISKS ASSOCIATED WITH ALOGLIPTIN USE IN PATIENTS WITH TYPE 2 DIABETES AT HIGH RISK OF ISCHEMIC CARDIOVASCULAR DISEASE
- M-178 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE OF REMISSION IN CROHN'S DISEASE IN PEDIATRIC PATIENTS
- M-179 UPDATES TO THE PRODUCT LABELING WITH STUDY REPORTS FROM THE OPTIMIST-1 AND OPTIMIST-2 CLINICAL TRIALS
- M-180 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF MAINTENANCE TREATMENT IN PATIENTS WITH SCHIZOPHRENIA
- M-181 UPDATE TO THE DOSAGE AND ADMINISTRATION, PATIENT SELECTION (2.1), SECTION OF THE PACKAGE INSERT TO INCLUDE THE USE OF AN FDA-APPROVED PLASMA TEST FOR THE IDENTIFICATION OF EGFR EXON 19 DELETION OR EXON 21 (L858R) SUBSTITUTION MUTATIONS
- M-182 UPDATES TO THE PRODUCT LABELING BASED ON THE RESULTS OF STUDY H7T-MC-TADO TITLED, "A PHASE 3 DOUBLE-BLIND, RANDOMIZED, MULTICENTER, EFFICACY AND SAFETY STUDY OF PRASUGREL COMPARED TO PLACEBO IN PEDIATRIC PATIENTS WITH SICKLE CELL DISEASE"
- M-183 CHANGES TO THE DOSAGE AND ADMINISTRATION AND CLINICAL STUDIES SECTIONS OF THE LABELING TO SUPPORT THE REDUCE-TO-QUIT PARADIGM
- M-184 UPDATES MADE TO THE LABELING TO INCLUDE INFORMATION FROM STUDY MO25743 ON THE ANTI-TUMOR ACTIVITY OF VEMURAFENIB IN THE TREATMENT OF PATIENTS WITH BRAF V600E MUTATION-POSITIVE MELANOMA WITH BRAIN METASTASES
- M-185 UPDATES TO THE LABELING TO INCLUDE RESULTS OF A TRIAL TO EVALUATE THE SAFETY OF MOXIFLOXACIN IN PEDIATRIC PATIENTS WITH COMPLICATED INTRA-ABDOMINAL INFECTIONS
- M-186 UPDATES TO THE PRODUCT INFORMATION REGARDING MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS BASED UPON THE RESULTS FROM STUDY 331-10-232
- M-187 ADDITION OF CLINICAL INFORMATION OBTAINED FROM A PEDIATRIC TRIAL TO SECTION 8.4 OF THE LABELING
- M-188 PROVIDES FOR DATA SUPPORTING THE SAFETY AND EFFECTIVENESS FOR THE MAINTENANCE TREATMENT OF MODERATE TO SEVERE BINGE EATING DISORDER (BED)

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- M-189 LABELING DESCRIBING THE EXPECTED REDUCTION OF ABUSE OF SINGLE-ENTITY MORPHINE BY THE INTRANASAL ROUTE OF ADMINISTRATION DUE TO PHYSICOCHEMICAL PROPERTIES
- M-190 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE LACK OF EFFICACY OF TARCEVA IN MAINTENANCE TREATMENT OF PATIENTS WITHOUT EGFR MUTATIONS
- M-191 ADDITION OF DATA BASED ON PEDIATRIC STUDIES TO FULFILL THE POSTMARKETING REQUIREMENT 1857-2
- M-192 PROVIDES FOR DATA EVALUATING THE NEUROPSYCHIATRIC SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN SUBJECTS WITH AND WITHOUT A HISTORY OF PSYCHIATRIC DISORDERS
- M-193 INFORMATION ADDED TO THE LABELING REGARDING A 15-WEEK, RANDOMIZED, DOUBLE-BLIND, PARALLEL-GROUP, PLACEBO-CONTROLLED FLEXIBLE-DOSE SAFETY AND EFFICACY STUDY OF PREGABALIN IN ADOLESCENTS (12 THROUGH 17 YEARS OLD) WITH FIBROMYALGIA
- M-194 INFORMATION ADDED TO THE LABELING REGARDING USE OF REGADENOSON ADMINISTRATION FOLLOWING AN INADEQUATE EXERCISE STRESS TEST AS COMPARED TO REGADENOSON ALONE
- M-195 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING REFLECTING LACK OF EFFICACY FOR IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17
- M-196 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM A RANDOMIZED, PLACEBO CONTROLLED, MULTICENTER STUDY OF INTRAVENOUS ACETAMINOPHEN FOR THE TREATMENT OF ACUTE PAIN IN PEDIATRIC PATIENTS TO FULFILL THE POST-MARKETING REQUIREMENT 1704-1
- M-197 NEW CLINICAL DATA ADDED TO THE PRESCRIBING INFORMATION REGARDING CANAGLIFLOZIN ADD-ON COMBINATION THERAPY WITH METFORMIN AND A DIPEPTIDYL-PEPTIDASE-4 INHIBITOR
- M-198 PACKAGE INSERT UPDATED WITH RESULTS FROM STUDY CV181168, A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED, PARALLEL GROUP, PHASE 3 TRIAL TO EVALUATE THE SAFETY AND EFFICACY OF SAXAGLIPTIN ADDED TO DAPAGLIFLOZIN AND METFORMIN
- M-199 INFORMATION ADDED TO LABELING REGARDING THE TREATMENT OF PATIENTS WITH ALK-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAD NOT RECEIVED PRIOR SYSTEMIC THERAPY FOR METASTATIC DISEASE.
- M-200 CLINICAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING.
- M-201 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM AN OPEN LABEL, MULTI-CENTER STUDY OF CABAZITAXEL IN PEDIATRIC PATIENTS WITH REFRACTORY SOLID TUMORS INCLUDING TUMORS OF THE CENTRAL NERVOUS SYSTEM.
- M-202 INCLUSION OF DATA FROM THE SUMMIT STUDY FOR BREO ELLIPTA (FLUTICASONE FUROATE/VILANTEROL TRIFENATATE) INHALATION POWDER IN THE PACKAGE INSERT.
- M-203 PROVIDES FOR REVISIONS TO THE PACKAGE INSERT TO REFLECT RESULTS OF TWO POSTMARKETING REQUIREMENT STUDIES ROP111662 AND ROP111569
- M-204 CLINICAL INFORMATION ADDED TO THE PACKAGE INSERT REGARDING USE OF ATORVASTATIN IN CHILDREN AND ADOLESCENTS AGES 10-17 WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- M-205 INFORMATION ADDED TO THE LABELING REGARDING RANDOMIZED, MULTICENTER, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDIES ON PATIENTS WITH SEVERE RENAL IMPAIRMENT
- M-206 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM COMPLERA TO ODEFSEY
- M-207 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM ATRIPLA TO ODEFSEY
- M-208 INFORMATION ADDED TO THE LABELING TO INCLUDE RESULTS OF A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS
- M-209 INFORMATION ADDED TO THE LABELING REGARDING CABAZITAXEL AT 20 MG/M2 BASED ON THE RESULTS OF THE PROSELICA STUDY
- M-210 INFORMATION ADDED TO LABELING TO SUPPORT THE USE OF SYMBICORT TO REDUCE EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- M-211 PROVIDES FOR LABELING CHANGES REGARDING THE USE OF DAPTOMYCIN IN THE PEDIATRIC POPULATION FOR STAPHYLOCOCCUS AUREUS BACTEREMIA (SAB) BASED ON RESULTS OF A

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TRIAL IN PEDIATRIC PATIENTS 1 TO 17 YEARS OF AGE

- M-212 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND EXENATIDE EXTENDED RELEASE
- M-213 INFORMATION ADDED TO THE LABELING TO INCLUDE THE EFFICACY AND SAFETY OF CARIPRAZINE RELATIVE TO PLACEBO IN THE PREVENTION OF RELAPSE OF SYMPTOMS IN PATIENTS WITH SCHIZOPHRENIA
- M-214 INFORMATION ADDED TO THE CLINICAL TRIALS SECTION OF THE LABELING REGARDING A POSTMARKETING SAFETY AND EFFICACY STUDY EVALUATING THE RISK OF SERIOUS ASTHMA-RELATED EVENTS
- M-215 INFORMATION ADDED TO THE LABELING REGARDING THE COMPARISON OF PALIPERIDONE PALMITATE COMPARED WITH ORAL ANTIPSYCHOTIC TREATMENT IN DELAYING TIME TO TREATMENT FAILURE IN ADULTS WITH SCHIZOPHRENIA WHO HAVE BEEN INCARCERATED
- M-216 UPDATE THE PRESCRIBING INFORMATION AND PATIENT LABELING WITH FINDINGS FROM STUDY RP103-08 CONDUCTED IN TREATMENT-NAIVE NEPHROPATHIC CYSTINOSIS PATIENTS TO EXPAND THE INDICATED POPULATION TO PATIENTS 1 YEAR AND OLDER
- M-217 INCORPORATION OF THE LABELING REVISIONS PROVIDED FOR IN NDA 022253/S-039 AND NDA 022255/S-022 INTO THE LACOSAMIDE INJECTION LABELING
- M-218 ADDITIONAL INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A NEW CLINICAL TRIAL IN PATIENTS AGED 6 THROUGH 11 YEARS (TRIAL 4)
- M-219 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A NEW CLINICAL TRIAL IN PATIENTS 7 TO 14 YEARS OF AGE WITH DUCHENNE MUSCULAR DYSTROPHY
- M-220 ADDITIONAL INFORMATION ADDED TO THE LABELING FROM STUDY PC B308/13 REGARDING THE USE OF BLUE LIGHT CYSTOSCOPY WITH CYSVIEW AS AN ADJUNCT TO WHITE LIGHT CYSTOSCOPY
- M-221 DRUG FACTS LABELING CHANGES UNDER THE DIRECTIONS HEADING TO REVISE THE STATED PREPARATION TIME OF A DRY SITE FROM 120 SECONDS SCRUBBING AND 90 SECONDS DRYING TO 30 SECONDS SCRUBBING AND 30 SECONDS DRYING
- M-222 ADDITION OF DATA BASED ON THE ASSESSMENT OF SAFETY AND EFFICACY IN PEDIATRIC PATIENTS WITH MAJOR DEPRESSIVE DISORDER TO FULFILL POSTMARKETING STUDY REQUIREMENT 1229-1
- M-223 INFORMATION ADDED TO SECTION 8.1 OF THE LABELING REGARDING PREGNANT PATIENTS WHO ARE ALREADY ON A STABLE RILPIVIRINE REGIMEN PRIOR TO PREGNANCY AND WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML)
- M-224 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF EXENATIDE EXTENDED RELEASE AS ADD-ON IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON BASAL INSULIN GLARGINE WITH OR WITHOUT METFORMIN
- M-225 REVISIONS TO SECTION 8.4 OF THE PRESCRIBING INFORMATION TO INCLUDE A SAFETY AND EFFICACY STUDY IN PEDIATRIC PATIENTS AGES ≥ 6 YEARS TO < 18 YEARS WITH CHRONIC IDIOPATHIC CONSTIPATION
- M-226 CHANGES TO THE LABELING BASED ON RESULTS FROM A CONTROLLED CLINICAL TRIAL IN PATIENTS WITH LATER-ONSET SPINAL MUSCULAR ATROPHY
- M-227 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING WITH THE SUBSECTION ENTITLED DIGIT SYMBOL SUBSTITUTION TEST IN MAJOR DEPRESSIVE DISORDER
- M-228 INFORMATION ADDED TO THE PACKAGE INSERT REGARDING THE REVISION OF THE MONOTHERAPY INDICATION OF VENETOCLAX
- M-229 REVISED LABELING TO INCORPORATE THE PEDIATRIC USE OF LOTEPIREDNOL ETABONATE GEL IN PATIENTS FOR THE TREATMENT OF POSTOPERATIVE INFLAMMATION FOLLOWING OCULAR SURGERY
- M-230 REVISIONS TO THE GLECAPREVIR/PIBRENTASVIR COMBINATION PRODUCT PRESCRIBING INFORMATION TO INCLUDE SAFETY AND EFFICACY DATA FROM THE HCV/HIV-1 COINFECTION STUDY M14-730 AND FROM THE LIVER AND RENAL TRANSPLANT STUDY M13-596
- M-231 REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION (SECTION 8.3) OF THE PACKAGE INSERT WITH THE RESULTS OF CLINICAL TRIAL WV25651, CONDUCTED TO EVALUATE THE EFFECT OF VALGANCICLOVIR ON SPERMATOGENESIS AND TO FULFILL PMR 1670-3
- M-232 INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO DESCRIBE THE RESULTS FROM PEDIATRIC STUDIES
- M-233 INFORMATION ADDED TO THE LABELING TO DESCRIBE FIXED-DOSE COMBINATION OF TIOPIPIUM BROMIDE AND OLODATEROL TO INCLUDE REDUCTION OF COPD EXACERBATIONS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-234 UPDATE TO THE PRESCRIBING INFORMATION FOR VORTIOXETINE ON TREATMENT-EMERGENT SEXUAL DYSFUNCTION COMPARING VORTIOXETINE AND SSRIS
- M-235 INFORMATION ADDED TO SECTION 14 OF THE LABELING TO DESCRIBE STUDY LAP016A2307 TO FULFILL POSTMARKETING STUDY REQUIREMENT 1586-1
- M-236 INFORMATION ADDED TO THE PRESCRIBING INFORMATION TO INCLUDE EFFICACY AND SAFETY DATA FROM A STUDY IN PATIENTS WITH TREATMENT NAIVE CLL/SLL TREATED WITH IBRUTINIB IN COMBINATION WITH OBINUTUZUMAB OR CHLORAMBUCIL IN COMBINATION WITH OBINUTUZUMAB
- M-237 INFORMATION ADDED TO LABELING TO DESCRIBE A STUDY TO EVALUATE THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN ADOLESCENT SMOKERS
- M-238 INFORMATION ADDED TO THE PRESCRIBING INFORMATION TO REFLECT THAT NO DOSE ADJUSTMENT IS NEEDED FOR PATIENTS WITH AN ESTIMATED GLOMERULAR FILTRATION RATE (EGFR) OF 45 ML/MIN/1.73 M² OR GREATER AS SUPPORTED BY CLINICAL STUDY REPORT
- M-239 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A TRIAL CONDUCTED IN TREATMENT NAIVE PEDIATRIC PATIENTS, AGES 2 YEARS TO < 18 YEARS WITH TRANSFUSIONAL IRON OVERLOAD
- M-240 INFORMATION ADDED TO LABELING REGARDING A RANDOMIZED, PLACEBO-CONTROLLED CLINICAL TRIAL TO EVALUATE CARDIOVASCULAR OUTCOMES AFTER TREATMENT WITH EXENATIDE ONCE WEEKLY IN PATIENTS WITH TYPE 2 DIABETES MELLITES
- M-241 INFORMATION ADDED TO THE LABELING FOR SAFETY & EFFICACY STUDY ENTITLED, A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO CONTROLLED CLINICAL TRIAL OF DEFERASIROX IN PATIENTS WITH MYELODYSPLASTIC SYNDROMES (LOW/INT-1 RISK) & TRANSFUSIONAL IRON OVERLOAD
- M-242 INFORMATION ADDED TO THE LABELING REGARDING THE EFFICACY AND SAFETY OF INSULIN DEGLUDEC/LIRAGLUTIDE VS INSULIN GLARGINE IN PTS W/ TYPE 2 DIABETES INADEQUATELY CONTROLLED ON SGLT2I WITH OR WITHOUT ORAL ANTIDIABETIC THERAPIES
- M-243 INFORMATION ADDED TO LABELING FROM PROSPECTIVE, RANDOMIZED, OPEN-LABEL, BLIND EVALUATOR (PROBE) STUDY EVALUATING THE EFFICACY AND SAFETY OF LOW MOLECULAR WEIGHT HEPARIN/EDOXABAN VERSUS DALTEPARIN IN VENOUS THROMBOEMBOLISM ASSOCIATED WITH CANCER
- M-244 INFORMATION ADDED TO THE LABELING REGARDING EFFICACY AND SAFETY OF THE CONTINUATION OF SITAGLIPTIN COMPARED WITH THE WITHDRAWAL OF SITAGLIPTIN DURING INITIATION AND TITRATION OF INSULIN GLARGINE IN SUBJECTS WITH TYPE 2 DIABETES MELLITUS
- M-245 ADDITIONAL INFORMATION ADDED TO THE LABELING BASED ON SAFETY AND EFFICACY DATA FROM THE IMPACT TRIAL
- M-246 ADDITION OF STUDY BRf117277, A NON-RANDOMIZED, OPEN-LABEL, MULTI-CENTER, MULTI-COHORT TRIAL OF DABRAFENIB PLUS TRAMETINIB IN SUBJECTS WITH BRAF MUTATION-POSITIVE MELANOMA THAT HAS METASTASIZED TO THE BRAIN
- M-247 REVISIONS TO THE LABELING REGARDING CONTINUOUS SUBCUTANEOUS INSULIN INFUSION AS A CONDITION OF USE FOR INSULIN ASPART
- M-248 INFORMATION ADDED TO THE LABELING TO DESCRIBE A TRIAL EVALUATING A LOWER DOSE THAN THOSE APPROVED FOR PEDIATRIC PATIENTS 13 TO 17 YEARS OF AGE
- M-249 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY LVM-MD-15 TO FULFILL POSTMARKETING COMMITMENT 1943-4
- M-250 REVISIONS TO THE PEDIATRIC USE SECTION TO INCLUDE AN OPEN-LABEL CLINICAL TRIAL TO FULFILL PMR 1655-1
- M-251 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING INFLUENZA VIRUS RESISTANCE TO OSELTAMIVIR IN IMMUNOCOMPROMISED PATIENTS
- M-252 ADDITION OF INFORMATION TO CLINICAL STUDIES SECTION REGARDING CARDIOVASCULAR OUTCOME
- M-253 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY P061, A RANDOMIZED, PLACEBO-CONTROLLED, PARALLEL GROUP, MULTI-SITE, DOUBLE-BLIND STUDY TO EVALUATE SAFETY AND EFFICACY OF SUVOREXANT FOR THE TREATMENT OF INSOMNIA IN SUBJECTS WITH ALZHEIMERS DISEASE
- M-254 INFORMATION ADDED TO THE LABELING REGARDING PEDIATRIC PATIENTS AGES 7 TO 17 YEARS OF AGE WITH MAJOR DEPRESSIVE DISORDER
- M-255 INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY GS-US-320-4018 IN VIROLOGICALLY SUPPRESSED ADULTS W/ CHRONIC HEP B INFECTION WHO SWITCHED FROM TENOFOVIR DISOPROXIL FUMARATE TO TENOFOVIR ALAFENAMIDE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

M-256	INFORMATION ADDED TO THE CLINICAL STUDIES SECTION TO FULFILL A POST-MARKETING REQUIREMENT
M-257	INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE USE OF PLAQUE PSORIASIS OF THE SCALP
M-258	INFORMATION ADDED TO THE LABELING TO DESCRIBE CARMELINA TRIAL TO FULFILL POSTMARKETING COMMITMENT 1766-4
M-259	INFORMATION ADDED TO THE LABELING REGARDING SAFETY AND EFFICACY IN SUBJECTS WITH HCV SUBTYPE 3B INFECTION
M-260	INFORMATION ADDED TO THE LABELING DESCRIBING A RANDOMIZED, OPEN-LABEL STUDY THAT EXAMINED THE CONCOMITANT USE OF DIMETHYL FUMARATE AND SEVERAL NON-LIVE VACCINES IN ADULTS 27-55 YEARS OF AGE WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS
M-261	ADDITIONAL INFORMATION ADDED TO THE LABELING REGARDING THE USE IN PATIENTS ON CHRONIC HEMODIALYSIS
M-262	REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE PACKAGE INSERT TO INCLUDE THE RESULT OF STUDY P146 TO FULFILL THE REQUIREMENTS OF PMR 3003-4
M-263	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY ICL670E2419 (THETIS TRIAL) TO SUPPORT PMR 3342-2 AND 3342-3
M-264	INFORMATION ADDED TO THE LABELING DESCRIBING A PHASE 2, MULTICENTER, OPEN-LABEL STUDY TO EVALUATE THE SAFETY/EFFICACY OF SOFOSBUVIR/VELPATASVIR IN SUBJECTS WITH CHRONIC HCV INFECTION WHO HAVE RECEIVED A LIVER TRANSPLANT
M-265	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY M15-656 (VIALE-A) AND M16-043 (VIALE-C) TO SUPPORT PMR 3545-1 AND PMR 3545-2
M-266	INFORMATION ADDED TO THE LABELING TO DESCRIBE STUDY GS-US-320-4035 IN VIROLOGICALLY SUPPRESSED ADULTS W/ CHRONIC HEP B INFECTION WHO SWITCHED FROM TENOFOVIR DISOPROXIL FUMARATE TO TENOFOVIR ALFAENAMIDE
M-267	INFORMATION ADDED TO THE LABELING REGARDING THE RESULT OF STUDY LUAA21004-402
M-268	ADDITION OF INFORMATION TO THE LABEL REGARDING A CLEAR PRODUCT PRESENTATION AND 26 ML VOLUME PRODUCTS
M-269	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY 309/KEYNOTE-775 TO SUPPORT PMR 3696-1 AND 3700-1
M-270	INFORMATION ADDED TO CLINICAL PHARMACOLOGY SECTION
M-271	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY HRA2914-5016
M-272	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY E7080-G000-211 TO SUPPORT PMR 2865-1
M-273	REVISION TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY 207966 ATLAS-2M
M-274	REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY CBAF312A2130
M-275	REVISION TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY, MK-8835-004/B1521021, VERTIS CV
M-276	REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDY 1200.120, CONDUCTED TO FULFILL A PEDIATRIC WRITTEN REQUEST
M-277	UPDATES THE US PRESCRIBING INFORMATION WITH CLINICAL DATA REGARDING THE USE OF SOFOSBUVIR AND VELPATASVIR FOR THE TREATMENT OF CHRONIC HCV GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION IN PEOPLE WHO INJECT DRUGS (PWID), INCLUDING THOSE ON MEDICATION-ASSISTED TREATMENT (MAT) FOR OPIOID USE DISORDER
M-278	INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULT OF STUDY G029665
M-279	INFORMATION ADDED TO THE LABELING TO DESCRIBE THE RESULTS OF FIGARO-DKD STUDY
M-280	REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDIES E7389-G000-223 AND E7389-G000-213, CONDUCTED TO FULFILL A PEDIATRIC WRITTEN REQUEST
M-281	REVISIONS TO THE LABELING TO PROVIDE FOR THE EXPANSION OF THE USE OF STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME IN PATIENTS TAKING CLOBAZAM TO INCLUDE PEDIATRIC PATIENTS WHO ARE 6 MONTHS TO LESS THAN 2 YEARS OF AGE AND WEIGHING 7 KG OR MORE
M-282	REVISIONS TO THE LABELING TO ADD THE RESULTS OF A CLINICAL STUDY (TA-303) IN PATIENTS WITH ED FOLLOWING BILATERAL NERVE-SPARING RADICAL PROSTATECTOMY
M-283	INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULT OF STUDY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

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M-284 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM GALILEO TRIAL

M-285 REVISIONS TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULTS OF STUDY INCB 18424-269

M-286 INFORMATION ADDED TO CLINICAL PHARMACOLOGY SECTION TO INCLUDE RESULTS FROM STUDY ORGN001-102

M-287 LABELING REGARDING NEW DOSING REGIMEN IN ADULTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) AND TREATMENT OF PAH IN PEDIATRIC PATIENTS (AGES 1-17)

M-288 INFORMATION ADDED TO THE LABELING TO DESCRIBE THE RESULTS OF STUDY APL2-308

M-289 INFORMATION ADDED TO THE LABELING TO DESCRIBE THE RESULTS OF MVT-601-035

M-290 INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO INCLUDE THE RESULT OF STUDY HZA114971

M-291 REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE PACKAGE INSERT TO INCLUDE THE RESULT OF STUDY P145

M-292 REVISIONS TO THE LABELING TO DESCRIBE MODIFIED FORMULATION BASED ON RESULTS OF STUDIES EM-05-014624 AND EM-05-014815

M-293 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM CLINICAL STUDY E2006-A001-113

M-294 INFORMATION ADDED TO THE LABELING REGARDING THE RESULT OF STUDY 205860

M-295 REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDY 1218-0091, CONDUCTED TO FULFILL A PEDIATRIC WRITTEN REQUEST

M-296 REVISIONS TO THE LABELING TO INCLUDE RESULTS FROM PROTOCOL 1218- 0091

M-297 LABELING CHANGES BASED ON RESULTS FROM CLINICAL STUDY MYK-461-017

M-298 LABELING REVISIONS RELATED TO STUDY D1699CC00001

M-299 CLINICAL STUDY INFORMATION ADDED TO THE LABEL ABOUT THE TREATMENT OF MODERATE TO SEVERE GENITAL PSORIASIS

M-300 REVISIONS TO THE PEDIATRIC USE SUBSECTION OF LABELING TO INCLUDE THE RESULTS FROM CLINICAL STUDY SYR-322-309, CONDUCTED IN RESPONSE TO A PEDIATRIC WRITTEN REQUEST

M-301 CLINICAL STUDY INFORMATION ADDED TO LABEL ABOUT THE TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN PATIENTS WITH SEVERELY REDUCED RENAL FUNCTION (ESTIMATED GLOMERULAR FILTRATION RATE, EGFR < 30 ML/MIN)

M-302 INFORMATION ADDED TO LABELING REGARDING OSTEOSARCOMA

ORPHAN DRUG EXCLUSIVITY

ODE-1 TO REDUCE CHRONIC DROOLING IN PATIENTS AGED 3 - 16 WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING (E.G. CEREBRAL PALSY)

ODE-2 FOR TREATMENT OF NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE

ODE-3 TO TREAT INFANTILE SPASMS

ODE-4 TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS WHO REQUIRE THERAPEUTIC INTERVENTION BUT ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION

ODE-5 FOR SEQUENTIAL USE FOR THE TREATMENT OF CYANIDE POISONING THAT IS JUDGED TO BE LIFE-THREATENING

ODE-6 FOR THE MANAGEMENT OF POSTHERPETIC NEURALGIA

ODE-7 TO REDUCE THE RISK OF PRETERM BIRTH IN WOMEN WITH SINGLETON PREGNANCY WHO HAVE A HISTORY OF SINGLETON SPONTANEOUS PRETERM BIRTH

ODE-8 TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY

ODE-9 TREATMENT OF ASYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER IN PATIENTS WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE

ODE-10 FOR USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-11 TREATMENT OF PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- ODE-12 TREATMENT OF PERIPHERAL T-CELL LYMPHOMA (PTCL) IN PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-13 TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA WITH THE BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-14 TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA IN ADULTS 18 YEARS OF AGE AND OLDER
- ODE-15 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA APPROVED TEST
- ODE-16 TREATMENT OF PATIENTS WITH TRANSFUSIONAL IRON OVERLOAD DUE TO THALASSEMIA SYNDROMES WHEN CURRENT CHELATION THERAPY IS INADEQUATE
- ODE-17 ADJUNCTIVE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME IN PATIENTS 2 YEARS OF AGE OR OLDER
- ODE-19 TREATMENT OF PATIENTS WITH INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS, INCLUDING PRIMARY MYELOFIBROSIS, POST-POLYCYTHEMIA VERA MYELOFIBROSIS AND POST-ESSENTIAL THROMBOCYTHEMIA MYELOFIBROSIS
- ODE-20 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE A G551D MUTATION IN THE CFTR GENE.
- ODE-21 AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY
- ODE-22 FOR THE CONTROL OF HYPERGLYCEMIA SECONDARY TO HYPERCORTISOLISM IN ADULT PATIENTS WITH ENDOGENOUS CUSHING'S SYNDROME WHO HAVE TYPE 2 DIABETES MELLITUS OR GLUCOSE INTOLERANCE AND HAVE FAILED SURGERY OR ARE NOT CANDIDATES FOR SURGERY
- ODE-23 ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
- ODE-24 TREATMENT OF ADULTS WITH RENAL ANGIOMYOLIPOMA AND TUBEROUS SCLEROSIS COMPLEX (TSC) NOT REQUIRING IMMEDIATE SURGERY
- ODE-25 MANAGEMENT OF POSTHERPETIC NEURALGIA IN ADULTS.
- ODE-26 TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
- ODE-27 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
- ODE-28 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE (PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES
- ODE-29 LOCALIZATION OF LYMPH NODES DRAINING A PRIMARY TUMOR IN PATIENTS WITH MELANOMA WHEN USED WITH A HAND-HELD GAMMA COUNTER
- ODE-30 TREATMENT OF ADULT PATIENTS WITH CHRONIC, ACCELERATED OR BLAST PHASE PHILADELPHIA CHROMOSOME-POSITIVE (PH+) CHRONIC MYELOGENOUS LEUKEMIA (CML) WITH RESISTANCE, OR INTOLERANCE TO PRIOR THERAPY
- ODE-31 TREATMENT OF CORNEAL CYSTINE CRYSTAL ACCUMULATION IN PATIENTS WITH CYSTINOSIS
- ODE-32 TREATMENT OF ADULT PATIENTS WITH CHRONIC OR ACCELERATED PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE AND/OR INTOLERANCE TO TWO OR MORE TYROSINE KINASE INHIBITORS (TKI)
- ODE-33 TREATMENT OF PROGRESSIVE, METASTATIC MEDULLARY THYROID CANCER (MTC)
- ODE-34 TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-35 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) THAT IS RESISTANT OR INTOLERANT TO PRIOR TYROSINE KINASE INHIBITOR THERAPY.
- ODE-36 ADJUNCT TO A LOW-FAT DIET AND OTHER LIPID-LOWERING TREATMENTS, INCLUDING LDL APHERESIS WHERE AVAILABLE, TO REDUCE LDL-C, TC, APOLIPOPROTEIN B, & NON-HDL-C IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- ODE-37 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME (SBS) WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- ODE-38 PART OF COMBINATION THERAPY IN ADULTS (GREATER THAN OR EQUAL TO 18 YEARS) WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB)

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-39 TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS. & OLDER WITH NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT) SYNDROMES AND WITH A LIVER IRON CONCENTRATION OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT & SERUM FERRITIN GREATER THAN 300 MCG/L.
- ODE-40 TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ ALL) IN COMBINATION WITH CHEMOTHERAPY, APPROVED UNDER NDA #21588/S-037
- ODE-41 ADJUNCT TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LDL-C, APOLIPOPROTEIN B (APO B), TOTAL CHOLESTEROL (TC), AND NON-HIGH DENSITY LIPOPROTEIN-CHOLESTEROL (NON-HDL-C) IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- ODE-42 USE AS A NITROGEN-BINDING ADJUNCTIVE THERAPY FOR CHRONIC MGMT OF ADULT AND PEDIATRIC PATIENTS AT LEAST 2 YRS WITH UREA CYCLE DISORDERS THAT CANNOT BE MANAGED BY DIETARY PROTEIN RESTRICTION AND/OR AMINO ACID SUPPLEMENTATION ALONE
- ODE-43 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING LENALIDOMIDE AND BORTEZOMIB AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY.
- ODE-44 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE.
- ODE-45 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS IN ADULTS AND CHILDREN AGES 6 YEARS AND OLDER.
- ODE-46 IMPROVEMENT OF NEUROLOGICAL OUTCOME BY REDUCING THE INCIDENCE AND SEVERITY OF ISCHEMIC DEFICITS IN ADULT PATIENTS WITH SUBARACHNOID HEMORRHAGE FROM RUPTURED INTRACRANIAL BERRY ANEURYSMS REGARDLESS OF THEIR POST-ICTUS NEUROLOGICAL CONDITION
- ODE-47 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA APPROVED TEST.
- ODE-48 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA APPROVED TEST
- ODE-49 TREATMENT OF MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- ODE-50 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST.
- ODE-51 TOPICAL TREATMENT OF STAGE 1A AND 1B MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN-DIRECTED THERAPY
- ODE-52 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS AS FIRST-LINE TREATMENT, IN COMBINATION WITH GEMCITABINE.
- ODE-53 TREATMENT OF ADULTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) WHO GROUP 1, TO IMPROVE EXERCISE CAPACITY, WHO FUNCTIONAL CLASS AND TO DELAY CLINICAL WORSENING.
- ODE-54 TX OF PAH TO DELAY DISEASE PROGRESSION. DISEASE PROGRESSION INCLUDED: DEATH, INITIATION OF IV OR SC PROSTANOIDS, OR CLINICAL WORSENING OF PAH (DECREASED 6-MINUTE WALK DISTANCE, WORSENER PAH SYMPTOMS AND NEED FOR ADDITIONAL PAH TREATMENT).
- ODE-55 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-56 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DCT) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT.
- ODE-57 TRAMETINIB IN COMBO WITH DABRAFENIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-58 DABRAFENIB IN COMBO WITH TRAMETINIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-59 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER
- ODE-60 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED

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AT LEAST ONE PRIOR THERAPY

- ODE-61 TREATMENT OF NEUROGENIC SYMPTOMATIC ORTHOSTATIC HYPOTENSION IN PATIENTS WITH PRIMARY AUTONOMIC FAILURE, DOPAMINE-BETA-HYDROXYLASE DEFICIENCY, AND NONDIABETIC AUTONOMIC NEUROPATHY
- ODE-62 TREATMENT OF PROLIFERATING INFANTILE HEMANGIOMA REQUIRING SYSTEMIC THERAPY.
- ODE-63 TREATMENT OF VISCERAL LEISHMANIASIS DUE TO LEISHMANIA DONOVANI; CUTANEOUS LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS, LEISHMANIA GUYANENSIS, AND LEISHMANIA PANAMENSIS; AND MUCOSAL LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS.
- ODE-64 SELECTIVE HEPATIC INTRA-ARTERIAL USE FOR IMAGING TUMORS IN ADULTS WITH KNOWN HEPATOCELLULAR CARCINOMA (HCC)
- ODE-65 TREATMENT OF PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS PART OF A COMBINATION REGIMEN.
- ODE-66 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB.
- ODE-67 GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY
- ODE-68 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA
- ODE-69 TREATMENT OF MALIGNANT HYPERTHERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTHERMIA IN PATIENTS AT HIGH RISK
- ODE-70 RELAPSED CLL, IN COMBO. WITH RITUXIMAB, IN PATIENTS FOR WHOM RITUXIMAB ALONE WOULD BE CONSIDERED APPROPRIATE THERAPY DUE TO OTHER CO-MORBIDITIES; AND RELAPSED SLL IN PATIENTS WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES
- ODE-71 RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA (FL) IN PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-72 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-73 LONG-TERM TREATMENT OF ADULT PATIENTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS (EMS), INTERMEDIATE METABOLIZERS (IMS), OR POOR METABOLIZERS (PMS) AS DETECTED BY AN FDA-CLEARED TEST.
- ODE-74 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY
- ODE-75 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY.
- ODE-76 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE NOT RECEIVED AT LEAST 1 PRIOR THERAPY
- ODE-77 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- ODE-78 TREATMENT OF HYPERCALCEMIA IN ADULT PATIENTS WITH PRIMARY HYPERPARATHYROIDISM FOR WHOM PARATHYROIDECTOMY WOULD BE INDICATED ON THE BASIS OF SERUM CALCIUM LEVELS, BUT WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY.
- ODE-79 TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA
- ODE-80 TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S
- ODE-81 TREATMENT OF PATIENTS WITH ACROMEGALY WHO HAVE HAD AN INADEQUATE RESPONSE TO SURGERY AND/OR FOR WHOM SURGERY IS NOT AN OPTION
- ODE-82 TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL- OR MODERATELY-DIFFERENTIATED LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS TO IMPROVE PROGRESSION-FREE SURVIVAL
- ODE-83 USE OF AS MONOTHERAPY FOR PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA MUTATED (AS DETECTED BY AN FDA-APPROVED TEST) ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY
- ODE-84 TREATMENT OF MOTOR FLUCTUATIONS IN PATIENTS WITH ADVANCED PARKINSON'S DISEASE
- ODE-85 AS A REPLACEMENT SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY (CRRT) AND IN CASE OF DRUG POISONING WHEN CRRT IS USED TO REMOVE DIALYZABLE SUBSTANCES
- ODE-86 TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA
- ODE-87 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE,

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RADIOACTIVE IODINE REFRACTORY DIFFERENTIATED THYROID CANCER

- ODE-88 FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE TREATMENT)
- ODE-89 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR REGIMENS, INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT
- ODE-90 TREATMENT OF INVASIVE MUCORMYCOSIS IN PATIENTS 18 YEARS OF AGE AND OLDER
- ODE-91 TREATMENT OF BILE ACID SYNTHESIS DISORDERS DUE TO SINGLE ENZYME DEFECTS
- ODE-92 TREATMENT OF LYMPHANGIOLEIOMYOMATOSIS (LAM)
- ODE-93 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR F508DEL MUTATION IN THE CFTR GENE
- ODE-94 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY TRANSPLANT PATIENTS CONVERTED FROM TACROLIMUS IMMEDIATE-RELEASE FORMULATIONS IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-95 FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-96 TREATMENT OF PRIMARY HYPERKALEMIC PERIODIC PARALYSIS, PRIMARY HYPOKALEMIC PERIOD PARALYSIS, AND RELATED VARIANTS
- ODE-97 TO EXPAND THE INDICATION TO PEDIATRIC PATIENTS 2-6 YEARS OF AGE WITH NEPHROPATHIC CYSTINOSIS
- ODE-98 TREATMENT OF HEREDITARY OROTIC ACIDURIA
- ODE-99 FOR USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, FOR THE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED FOLLOWING GEMCITABINE-BASED THERAPY
- ODE-100 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA OR LEIOMYOSARCOMA WHO RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-101 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATION, IN COMBINATION WITH VEMURAFENIB. COTELLIC IS NOT INDICATED FOR TREATMENT OF PATIENTS WITH WILD-TYPE BRAF MELANOMA
- ODE-102 FOR TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), AS DETECTED BY AN FDA-APPROVED TEST, WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY
- ODE-103 USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-104 EMERGENCY TX OF PTS FOLLOWING A FU OR CAPECITABINE OD, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING TOXICITY AFFECTING THE CARDIAC SYSTEM OR CNS, AND/OR EARLY-ONSET, UNUSUALLY SEVERE AR W/IN 96 HRS FOLLOWING THE END OF FU OR CAPECITABINE ADMIN.
- ODE-105 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-106 FOR USE OF UPTRAVI (SELEXIPAG) TABLETS, 200, 400, 600, 800, 1000, 1200, 1400, AND 1600 MCG FOR TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH, WHO GROUP I) TO REDUCE THE RISKS OF DISEASE PROGRESSION AND HOSPITALIZATION FOR PAH
- ODE-107 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-108 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL-DIFFERENTIATED, NON-FUNCTIONAL, NEUROENDOCRINE TUMORS (NET) OF GASTROINTESTINAL (GI) OR LUNG ORIGIN, (EXCLUDING PANCREATIC) WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- ODE-109 INDICATED FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITHOUT 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE THERAPY)
- ODE-110 FOR HIGH-DOSE CONDITIONING TREATMENT PRIOR TO HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- ODE-111 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS ARE ROS-1 POSITIVE.

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- ODE-112 FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH HEPATIC VENO-OCCLUSIVE DISEASE (VOD), ALSO KNOWN AS SINUSOIDAL OBSTRUCTION SYNDROME (SOS), WITH RENAL OR PULMONARY DYSFUNCTION FOLLOWING HEMATOPOIETIC STEM CELL TRANSPLANTATION (HSCT).
- ODE-113 FOR TREATMENT OF PEDIATRIC AND ADULT PATIENTS WITH ACQUIRED METHEMOGLOBINEMIA.
- ODE-114 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION, AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-115 TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSING AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-116 TREATMENT OF PROGRESSIVE KERATOCONUS
- ODE-117 FOR TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-118 AN ADJUNCT TO DIET TO REDUCE LDL-C, TOTAL-C, NONHDL-C AND APOB IN CHILDREN AND ADOLESCENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS (E.G., LDL APHERESIS)
- ODE-119 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN COMBINATION WITH URSODEOXYCHOLIC ACID (UDCA) IN ADULTS WITH AN INADEQUATE RESPONSE TO UDCA, OR AS MONOTHERAPY IN ADULTS UNABLE TO TOLERATE UDCA
- ODE-120 FOR USE AFTER RADIOLABELING WITH GA 68, WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT AND PEDIATRIC PATIENTS.
- ODE-121 TREATMENT OF CORNEAL ECTASIA FOLLOWING REFRACTIVE SURGERY
- ODE-122 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- ODE-123 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6-11 YEAR OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
- ODE-124 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE FORMULATIONS, WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE, IN ADULTS WITH THE FOLLOWING SEIZURE TYPES: PARTIAL WITH COMPLEX SYMPTOMOLOGY, GENERALIZED CLONIC-TONIC, AND MIXED
- ODE-125 INDICATED IN PEDIATRIC PATIENTS 10 YEARS AND OLDER FOR THE PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGES 3 AND 4 AND CKD STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS
- ODE-126 AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES
- ODE-127 TREATMENT OF SPINAL MUSCULAR ATROPHY IN PEDIATRIC AND ADULT PATIENTS
- ODE-128 TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY
- ODE-129 INDICATED FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED IN CONJUNCTION WITH HIGH-DOSE BUSULFAN & CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENIC HEMATOPOIETIC PROGENITOR CELL TRANSPLANTATION FOR PEDIATRIC PATIENTS WITH CLASS 3 BETA-THALASSEMIA
- ODE-130 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS 5 YEARS OF AGE AND OLDER
- ODE-131 TREATMENT OF MULTIPLE MYELOMA (MM), AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)
- ODE-132 TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY
- ODE-133 INDICATED FOR MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-134 TREATMENT OF CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
- ODE-135 TREATMENT OF CHRONIC HCV GENOTYPE 2 OR 3 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-136 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1, 4, 5, OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS

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- ODE-137 TREATMENT OF OLIGOARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PERSISTENT OLIGOARTHRITIS, PSORIATIC JUVENILE IDIOPATHIC ARTHRITIS, ENTHESITIS-RELATED ARTHRITIS, OR UNDIFFERENTIATED ARTHRITIS) & POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS IN CHILDREN 0-16 YRS
- ODE-138 TREATMENT OF PEDIATRIC PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS A COMPONENT OF A COMBINATION CHEMOTHERAPY MAINTENANCE REGIMEN
- ODE-139 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC OR LIVER CANCER) WHO HAVE BEEN PREVIOUSLY TREATED WITH THE DRUG SORAFENIB.
- ODE-140 TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)
- ODE-141 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) THAT IS FLT3 MUTATION-POSITIVE AS DETECTED BY AN FDA APPROVED TEST, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION
- ODE-142 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-143 TO DECREASE THE RECURRENCE OF PNEUMOTHORAX IN ADULTS
- ODE-144 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)
- ODE-145 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- ODE-146 OPTICAL IMAGING AGENT INDICATED IN PATIENTS WITH GLIOMA (SUSPECTED WORLD HEALTH ORGANIZATION GRADES III OR IV ON PREOPERATIVE IMAGING) AS AN ADJUNCT FOR THE VISUALIZATION OF MALIGNANT TISSUE DURING SURGERY
- ODE-147 DABRAFENIB IN COMBINATION WITH TRAMETINIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-148 TRAMETINIB IN COMBINATION WITH DABRAFENIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-149 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY
- ODE-150 TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS 5 YEARS OF AGE AND OLDER.
- ODE-151 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-152 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD)
- ODE-153 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- ODE-154 FOR USE IN CHILDREN AGES 2 TO 12 YEARS OLD WITH CHAGAS DISEASE
- ODE-155 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-156 TREATMENT OF ADULTS WITH CARCINOID SYNDROME; WHEN USED, IT REDUCES THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY
- ODE-157 FOR USE AS A NITROGEN-BINDING AGENT FOR CHRONIC MANAGEMENT OF PEDIATRIC PATIENTS ≥ 2 MONTHS AND < 2 YEARS OF AGE WITH UREA CYCLE DISORDERS (UCDS) WHO CANNOT BE MANAGED BY DIETARY PROTEIN RESTRICTION AND/OR AMINO ACID SUPPLEMENTATION ALONE
- ODE-158 TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION
- ODE-159 FOR TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK) POSITIVE, METASTATIC NON-SMALL-CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST, EXCLUDING PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-160 FOR TREATMENT OF SCURVY IN ADULT AND PEDIATRIC PATIENTS AGE 5 MONTHS AND OLDER FOR WHOM ORAL ADMINISTRATION IS NOT POSSIBLE, INSUFFICIENT OR CONTRAINDICATED
- ODE-161 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) IN PEDIATRIC PATIENTS AGED 3 YRS AND OLDER WITH IDIOPATHIC OR CONGENITAL PAH TO IMPROVE PULMONARY VASCULAR RESISTANCE (PVR), WHICH IS EXPECTED TO RESULT IN AN

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IMPROVEMENT IN EXERCISE ABILITY

- ODE-162 TREATMENT OF NEPHROPATHIC CYSTINOSIS IN PEDIATRIC PATIENTS 1 YEAR OF AGE TO LESS THAN 2 YEARS OF AGE
- ODE-163 TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+ CML)
- ODE-164 TREATMENT OF PEDIATRIC PATIENTS WITH PHILADELPHIA CHROMOSOME-POSITIVE (PH+) CHRONIC MYELOID LEUKEMIA (CML) IN CHRONIC PHASE
- ODE-165 PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE IN ADULT CMV-SEROPOSITIVE RECIPIENTS [R+] OF AN ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANT (HSCT)
- ODE-166 TREATMENT OF SOMATOSTATIN RECEPTOR-POSITIVE GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS) INCLUDING FOREGUT, MIDGUT, AND HINDGUT NEUROENDOCRINE TUMORS IN ADULTS
- ODE-167 ARSENIC TRIOXIDE FOR USE IN COMBINATION WITH TRETINOIN FOR TREATMENT OF ADULTS WITH NEWLY-DIAGNOSED LOW-RISK ACUTE PROMYELOCYTIC LEUKEMIA (APL) WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- ODE-168 FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-169 FOR THE ADJUNCTIVE TREATMENT OF ADULT AND PEDIATRIC PATIENTS AGED 2 YEARS AND OLDER WITH TUBEROUS SCLEROSIS COMPLEX (TSC)-ASSOCIATED PARTIAL-ONSET SEIZURES
- ODE-170 FOR THE DIAGNOSIS OF ADULT GROWTH HORMONE DEFICIENCY (AGHD)
- ODE-171 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML) IN CHRONIC PHASE
- ODE-172 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH CHRONIC PHASE PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA WITH RESISTANCE OR INTOLERANCE TO PRIOR TYROSINE-KINASE INHIBITOR THERAPY
- ODE-173 TREATMENT OF PATIENTS WITH CYSTIC FIBROSIS AGED 12 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR GENE RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- ODE-174 FOR THE TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA (ITP) WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- ODE-175 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-176 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-177 TO REDUCE THE FREQUENCY OF PAINFUL CRISES AND TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS IN PEDIATRIC PATIENTS, 2 YEARS OF AGE AND OLDER, WITH SICKLE CELL ANEMIA WITH RECURRENT MODERATE TO SEVERE PAINFUL CRISIS
- ODE-178 INDICATED TO SLOW KIDNEY FUNCTION DECLINE IN ADULTS AT RISK OF RAPIDLY PROGRESSING AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)
- ODE-179 TREATMENT OF PATIENTS WITH CLL AND TREATMENT OF PATIENTS WITH INDOLENT B-CELL NHL THAT HAS PROGRESSED DURING OR WITHIN SIX MONTHS OF TREATMENT WITH RITUXIMAB OR A RITUXIMAB CONTAINING REGIMEN
- ODE-180 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-181 TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY
- ODE-182 TRAMETINIB IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION

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- ODE-183 TRAMETINIB AND DABRAFENIB IN COMBINATION, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- ODE-184 INDICATED IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF PEDIATRIC PATIENTS WITH HIV-1 INFECTION
- ODE-185 INDICATED FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-186 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, OR S549R
- ODE-187 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE
- ODE-188 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGES 2 TO LESS THAN 6 YEARS WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: G551D, G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, S549R, AND R117H
- ODE-189 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 2 YEARS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: 711+3A-G, E831X, 2789+5G-A, 3272-26A-G, AND 3849+10KBC-T
- ODE-190 TX OF CF IN PTS 2 YRS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: E56K, P67L, R74W, D110E, D110H, R117C, E193K, L206W, R347H, R352Q, A455E, D579G, S945L, S977F, F1052V, K1060T, A1067T, G1069R, R1070Q, R1070W, F1074L, D1152H, D1270N
- ODE-191 TO DECREASE THE RECURRENCE OF MALIGNANT PLEURAL EFFUSIONS IN SYMPTOMATIC PATIENTS FOLLOWING MAXIMAL DRAINAGE OF THE PLEURAL EFFUSION
- ODE-192 INDICATED TO INDUCE CONTROLLED CARDIAC SEPTAL INFRACTION TO IMPROVE EXERCISE CAPACITY IN ADULTS WITH SYMPTOMATIC HYPERTROPHIC OBSTRUCTIVE CARDIOMYOPATHY WHO ARE NOT CANDIDATES FOR SURGICAL MYECTOMY
- ODE-193 INDICATED FOR THE TREATMENT OF ONCHOCERCIASIS DUE TO ONCHOCERCA VOLVULUS IN PATIENTS AGED 12 YEARS AND OLDER
- ODE-194 ENCORAFENIB IS INDICATED IN COMBINATION WITH BINIMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH A BRAF V600E OR V600K MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-195 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 2 THROUGH 5 YEARS OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
- ODE-196 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH UNRESECTABLE HEPATOCELLULAR CARCINOMA (HCC)
- ODE-197 INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- ODE-198 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME (DS) IN PATIENTS 2 YEARS OF AGE AND OLDER TAKING CLOBAZAM
- ODE-199 THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 12 MONTHS AND OLDER WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-200 INDICATED FOR THE TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 13 KG
- ODE-201 INDICATED FOR THE RADICAL CURE (PREVENTION OF RELAPSE) OF PLASMODIUM VIVAX MALARIA IN PATIENTS AGED 16 YEARS AND OLDER WHO ARE RECEIVING APPROPRIATE ANTIMALARIAL THERAPY FOR ACUTE P. VIVAX INFECTION
- ODE-202 INDICATED AS A SOURCE OF CALORIES AND FATTY ACIDS IN PEDIATRIC PATIENTS WITH PARENTERAL NUTRITION-ASSOCIATED CHOLESTASIS (PNAC)
- ODE-203 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-204 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH IOBENGUANE SCAN POSITIVE, UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC PHEOCHROMOCYTOMA OR PARAGANGLIOMA WHO REQUIRE SYSTEMIC ANTICANCER THERAPY
- ODE-205 INDICATED FOR THE TREATMENT OF ADULTS WITH A CONFIRMED DIAGNOSIS OF FABRY DISEASE AND AN AMENABLE GALACTOSIDASE ALPHA GENE (GLA) VARIANT BASED ON IN VITRO ASSAY DATA

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- ODE-206 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-207 TREATMENT OF STATUS EPILEPTICUS IN ADULTS
- ODE-208 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) AFTER AT LEAST TWO PRIOR THERAPIES
- ODE-209 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) AFTER AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-210 INDICATED IN COMBINATION WITH STANDARD IMMUNOSUPPRESSIVE THERAPY FOR THE FIRST-LINE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS AND OLDER WITH SEVERE APLASTIC ANEMIA
- ODE-211 INDICATED IN COMBO WITH AZACITIDINE, OR DECITABINE, OR LOW-DOSE CYTARABINE FOR THE TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-212 INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- ODE-213 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR (EGFR) EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-214 TX OF MAC LUNG DISEASE IN ADULTS WITH LIMITED OR NO ALTERNATIVE TX OPTIONS AS PART OF A COMBO ANTIBACTERIAL DRUG REGIMEN WHO DO NOT ACHIEVE NEGATIVE SPUTUM CULTURES AFTER A MINIMUM OF 6 CONSECUTIVE MONTHS OF A MULTIDRUG BACKGROUND REGIMEN THERAPY
- ODE-215 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC RECEPTOR TYROSINE KINASE (NTRK) GENE FUSION WITHOUT A KNOWN ACQUIRED RESISTANCE MUTATION
- ODE-216 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) OR DRAVET SYNDROME (DS) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-217 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON CRIZOTINIB AND AT LEAST ONE OTHER ALK INHIBITOR FOR METASTATIC DISEASE
- ODE-218 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON ALECTINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-219 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON CERITINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-220 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY
- ODE-221 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT HAVE NO SATISFACTORY ALTERNATIVE TREATMENTS OR THAT HAVE PROGRESSED FOLLOWING TREATMENT
- ODE-222 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WHO HAVE RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A FMS-LIKE TYROSINE KINASE 3 (FLT3) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-223 TREATMENT OF LAMBERT-EATON MYASTHENIC SYNDROME (LEMS)
- ODE-224 INDICATED, IN COMBINATION WITH LOW-DOSE CYTARABINE, FOR THE TREATMENT OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULT PATIENTS WHO ARE ≥ 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-225 INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN COMBINATION WITH CHEMOTHERAPY
- ODE-226 MAINTENANCE TREATMENT OF ADULTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR

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- PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- ODE-227 INDICATED FOR TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
- ODE-228 INDICATED FOR THE TREATMENT OF FASCIOLIASIS IN PATIENTS 6 YEARS OF AGE AND OLDER
- ODE-229 TREATMENT OF ADULT PATIENTS WITH METASTATIC GASTRIC OR GASTROESOPHAGEAL JUNCTION ADENOCARCINOMA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
- ODE-230 FIRST-LINE TREATMENT OF METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS HAVE NON-RESISTANT EPIDERMAL GROWTH FACTOR (EGFR) MUTATIONS OTHER THAN EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-231 INDICATED FOR THE TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PEDIATRIC PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY
- ODE-232 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1,2,3,4,5 OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS (CHILD-PUGH A)
- ODE-233 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG WITH HCV GENOTYPE 1 INFECTION, WHO PREVIOUSLY HAVE BEEN TREATED WITH A REGIMEN CONTAINING AN HCV NS5A INHIBITOR OR AN NS3/4A PROTEASE INHIBITOR (PI), BUT NOT BOTH
- ODE-234 INDICATED FOR THE TREATMENT OF STABLE SYMPTOMATIC HEART FAILURE DUE TO DILATED CARDIOMYOPATHY (DCM) IN PEDIATRIC PATIENTS AGED 6 MONTHS AND OLDER, WHO ARE IN SINUS RHYTHM WITH AN ELEVATED HEART RATE
- ODE-235 INDICATED FOR THE TREATMENT OF ACUTE HERPETIC KERATITIS (DENDRITIC ULCERS) IN PATIENTS WITH HERPES SIMPLEX (HSV-1 AND HSV-2) VIRUS
- ODE-236 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 MONTHS TO LESS THAN 12 MONTHS WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-237 TREATMENT OF THE CARDIOMYOPATHY OF WILD TYPE OR HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM) IN ADULTS TO REDUCE CARDIOVASCULAR MORTALITY AND CARDIOVASCULAR-RELATED HOSPITALIZATION
- ODE-238 TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (GVHD) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- ODE-239 TREATMENT OF PREVIOUSLY UNTREATED ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-240 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH SHORT BOWEL SYNDROME (SBS) WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- ODE-241 INDICATED IN COMBINATION WITH A RITUXIMAB PRODUCT FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED FOLLICULAR LYMPHOMA (FL)
- ODE-242 TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 MUTATION AS DETECTED BY AN FDA-APPROVED TEST IN ADULT PTS WHO ARE >=75 YRS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-243 ACUTE TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E., SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
- ODE-244 TREATMENT OF LAMBERT-EATON MYASTHENIC SYNDROME (LEMS) IN PATIENTS 6 TO LESS THAN 17 YEARS OF AGE
- ODE-245 INDICATED IN COMBINATION WITH A RITUXIMAB PRODUCT FOR THE TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA (MZL)
- ODE-246 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- ODE-247 TX OF PTS W/ CYSTIC FIBROSIS (CF) AGE 6 TO <12 YRS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR W/ AT LEAST 1 MUTATION IN CF TRANSMEMBRANE CONDUCTANCE REGULATORY GENE RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
- ODE-248 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCETS DISEASE

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- ODE-249 TREATMENT OF PATIENTS WITH CYSTIC FIBROSIS (CF) AGE 6 YEARS TO LESS THAN 12 YEARS WHO ARE HOMOZYGOUS FOR F508DEL MUTATION OR WHO HAVE AT LEAST ONE MUTATION IN THE CF TRANSMEMBRANE CONDUCTANCE REGULATOR GENE THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- ODE-250 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH SYMPTOMATIC TENOSYNOVIAL GIANT CELL TUMOR (TGCT) ASSOCIATED WITH SEVERE MORBIDITY OR FUNCTIONAL LIMITATIONS AND NOT AMENABLE TO IMPROVEMENT WITH SURGERY
- ODE-251 INDICATED AS PART OF COMBINATION THERAPY IN THE TREATMENT OF PEDIATRIC PATIENTS (12 TO LESS THAN 18 YEARS OF AGE AND WEIGHING AT LEAST 30 KG) WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS
- ODE-252 TREATMENT OF DUCHENE MUSCULAR DYSTROPHY IN PATIENTS 2 YEARS OF AGE TO LESS THAN 5 YEARS OF AGE
- ODE-253 INDICATED AS PART OF A COMBINATION REGIMEN WITH BEDAQUILINE AND LINEZOLID FOR THE TREATMENT OF ADULTS WITH PULMONARY EXTENSIVELY DRUG RESISTANT (XDR) OR TREATMENT-INTOLERANT OR NONRESPONSIVE MULTIDRUG-RESISTANT (MDR) TUBERCULOSIS (TB)
- ODE-254 INDICATED TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY
- ODE-255 INDICATED FOR THE TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS (EDS) IN ADULT PATIENTS WITH NARCOLEPSY
- ODE-256 FOR HIV-1 INFECTION IN PEDIATRIC PTS AT LEAST 25 KG W/ NO ANTIRETROVIRAL (ARV) TX HX OR TO REPLACE CURRENT ARV REGIMEN FOR VIROLOGICALLY-SUPPRESSED ON STABLE ARV W/ NO HX TX FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED W/ RESISTANCE TO BIC, FTC, OR TAF
- ODE-257 IN COMBO W/ DEXAMETHASONE FOR ADULTS W/ RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO RECEIVED AT LEAST 4 PRIOR THERAPIES AND REFRACTORY TO AT LEAST 2 PROTEASOME INHIBITORS, AT LEAST 2 IMMUNOMODULATORY AGENTS, AND AN ANTI-CD38 MONOCLONAL ANTIBODY
- ODE-258 FOR THE TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 2 OR 3 INFECTION IN PEDIATRIC PATIENTS BETWEEN 3 YEARS OF AGE AND 12 YEARS OF AGE OR WEIGHING 35 KG WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-259 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY (POST-POLYCYTHEMIA VERA OR POST-ESSENTIAL THROMBOCYTHEMIA) MYELOFIBROSIS (MF)
- ODE-260 INDICATED TO INCREASE SYSTEMIC EXPOSURE OF ATAZANAVIR IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS IN THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 35 KG
- ODE-261 INDICATED TO SLOW THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD)
- ODE-262 TREATMENT OF PEDIATRIC PATIENTS BETWEEN 3 YEARS OF AGE AND 12 YEARS OF AGE OR WEIGHING 35 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 4, 5, OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS
- ODE-263 TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH CHRONIC HCV GENOTYPE 1 INFECTION WITH DECOMPENSATED CIRRHOSIS, FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-264 TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH CHRONIC HCV GENOTYPE 1 OR 4 INFECTION WHO ARE LIVER TRANSPLANT RECIPIENTS WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS, FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-265 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ROS1-POSITIVE
- ODE-266 ADULT & PED >=12YRS OLD W/ SOLID TUMORS THAT HAVE NTRK W/O KNOWN ACQUIRED RESISTANCE MUTATION, ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY & HAVE EITHER PROGRESSED FOLLOWING TX OR HAVE NO SATISFACTORY ALTERNATIVE TX
- ODE-267 INDICATED IN COMBINATION WITH HIGH FLUID INTAKE, ALKALI, AND DIET MODIFICATION, FOR THE PREVENTION OF CYSTINE STONE FORMATION IN PEDIATRIC PATIENTS 20KG TO 9 YEARS OF AGE W/SEVERE HOMOZYGOUS CYSTINURIA, WHO ARE NOT RESPONSIVE TO THESE MEASURES ALONE
- ODE-268 INDICATED FOR TREATMENT OF PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE

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ODE-269	PROPHYLAXIS OF ORGAN REJECTION IN PEDIATRIC PATIENTS RECEIVING ALLOGENEIC KIDNEY TRANSPLANT, LIVER TRANSPLANTS, AND HEART TRANSPLANT, IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
ODE-270	INDICATED TO INCREASE PAIN FREE LIGHT EXPOSURE IN ADULT PATIENTS WITH A HISTORY OF PHOTOTOXIC REACTIONS FROM ERYTHROPOIETIC PROTOPORPHYRIA (EPP)
ODE-271	INDICATED IN COMBINATION WITH OTHER ANTI-MYELOMA PRODUCTS FOR THE TREATMENT OF ADULTS WITH MULTIPLE MYELOMA (MM)
ODE-272	INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO DELAY DISEASE PROGRESSION
ODE-273	INDICATED FOR THE TREATMENT OF ADULTS WITH ACUTE HEPATIC PORPHYRIA (AHP)
ODE-274	INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
ODE-275	INDICATED FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE
ODE-276	INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
ODE-277	TX OF ADULTS W/ ADV OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & CANCER ASSOCIATED W/ HRD+ STATUS DEFINED BY A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION
ODE-278	TX OF ADULTS W/ ADV OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER TREATED W/ >=3 PRIOR CHEMO REGIMENS & CANCER ASSOCIATED W/ HRD+ STATUS DEFINED BY GENOMIC INSTABILITY & PROGRESSED >6 MONTHS AFTER RESPONSE TO THE LAST PLATINUM-BASED CHEMOTHERAPY
ODE-279	INDICATED FOR THE ACUTE TX OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E. SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 6 YEARS OF AGE AND OLDER
ODE-280	INDICATED FOR THE TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
ODE-281	INDICATED FOR THE TREATMENT OF SICKLE CELL DISEASE (SCD) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
ODE-282	INDICATED TO SELECTIVELY STAIN THE INTERNAL LIMITING MEMBRANE (ILM)
ODE-283	MAINTENANCE TX OF ADULTS W/ DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DZ HAS NOT PROGRESSED ON >=16WKS OF 1ST LINE PLATINUM BASED CHEMO REGIMEN. SELECT PTS FOR THERAPY BASED ON APPROVED COMPANION DIAGNOSTIC
ODE-284	IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS, FOR THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 35KG
ODE-285	IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS OTHER THAN PROTEASE INHIBITORS THAT REQUIRE A CYP3A INHIBITOR, FOR THE TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 25KG AND LESS THAN 35KG
ODE-286	INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
ODE-287	TREATMENT OF ADULTS WITH NEWLY DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
ODE-288	INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH NEUROFIBROMATOSIS TYPE 1 (NF1) WHO HAVE SYMPTOMATIC, INOPERABLE PLEXIFORM NEUROFIBROMAS (PN)
ODE-289	INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH LOW-GRADE UPPER TRACT UROTHELIAL CANCER (LG-UTUC)
ODE-290	INDICATED FOR THE INITIAL TREATMENT OF SEVERE MALARIA IN ADULT AND PEDIATRIC PATIENTS TO ALWAYS BE FOLLOWED BY A COMPLETE TREATMENT COURSE OF AN APPROPRIATE ORAL ANTIMALARIAL REGIMEN
ODE-291	INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE A MUTATION THAT LEADS TO MESENCHYMAL-EPIHELIAL TRANSITION (MET) EXON 14 SKIPPING AS DETECTED BY AN FDA-APPROVED TEST
ODE-292	INDICATED FOR THE TREATMENT OF ADULTS WITH PREVIOUSLY TREATED, UNRESECTABLE

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LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT AS DETECTED BY AN FDA-APPROVED TEST

- ODE-293 TX OF PED PTS 6 YRS OF AGE & OLDER OR WEIGHING AT LEAST 17 KG WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION: WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS; OR WITH DECOMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-294 PROPHYLAXIS OF ORGAN REJECTION IN PEDIATRIC PATIENTS RECEIVING ALLOGENEIC KIDNEY OR HEART TRANSPLANTS, IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-295 INDICATED FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH ADVANCED EPITHELIAL OVARIAN CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY
- ODE-296 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH AIDS-RELATED KAPOSI SARCOMA (KS) AFTER FAILURE OF HIGHLY ACTIVE ANTIRETROVIRAL THERAPY (HAART)
- ODE-297 FOR THE TREATMENT OF KAPOSI SARCOMA (KS) IN ADULT PATIENTS WHO ARE HIV-NEGATIVE
- ODE-298 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE RECEIVED PRIOR TREATMENT WITH 3 OR MORE KINASE INHIBITORS, INCLUDING IMATINIB
- ODE-299 INDICATED FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS AGED 16 YEARS AND OLDER WITH METASTATIC OR LOCALLY ADVANCED EPITHELIOID SARCOMA NOT ELIGIBLE FOR COMPLETE RESECTION
- ODE-300 FOR THE TREATMENT OF ADULT PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA-APPROVED TEST, NOT INCLUDING PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-301 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- ODE-302 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- ODE-303 ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- ODE-304 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC SMALL CELL LUNG CANCER (SCLC) WITH DISEASE PROGRESSION ON OR AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-305 TREATMENT OF INVASIVE ASPERGILLOSIS
- ODE-306 W/ BEVACIZUMAB FOR MAINT TX OF ADULTS W/ ADV EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CA IN COMPLETE OR PARTIAL RESPONSE TO 1ST LINE PT BASED CHEMO & WHOSE CA IS ASSOC W/ HOMOLOGOUS RECOMB DEF + STATUS DEFINED BY GENOMIC INSTABILITY
- ODE-307 INDICATED AS PART OF COMBINATION THERAPY IN THE TREATMENT OF PEDIATRIC PATIENTS 5 YEARS AND OLDER TO LESS THAN 12 YEARS OF AGE AND WEIGHING AT LEAST 15 KG WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB)
- ODE-308 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO IMPROVE EXERCISE CAPACITY
- ODE-309 INDICATED FOR USE IN COMBINATION WITH TRASTUZUMAB AND CAPECITABINE FOR TREATMENT OF ADULT PATIENTS WITH METASTATIC HER2-POSITIVE BREAST CANCER AND BRAIN METASTASES, WHO HAVE RECEIVED ONE OR MORE PRIOR ANTI-HER2-BASED REGIMENS IN THE METASTATIC SETTING
- ODE-310 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- ODE-311 INDICATED AS A SOURCE OF CALORIES AND FATTY ACIDS FOR THE TREATMENT OF PEDIATRIC AND ADULT PATIENTS WITH MOLECULARLY CONFIRMED LONG-CHAIN FATTY ACID OXIDATION DISORDERS (LC-FAOD)
- ODE-312 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-313 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION WITHOUT A KNOWN ACQUIRED RESISTANCE MUTATION, ARE METASTATIC OR

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WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY, AND HAVE EITHER PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY

- ODE-314 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA WHOSE TUMORS ARE POSITIVE FOR AN EZH2 MUTATION AS DETECTED BY AN FDA-APPROVED TEST AND WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES, AND FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- ODE-315 FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-316 TREATMENT OF ADULT PATIENTS WITH MYELODYSPLASTIC SYNDROMES (MDS), INCLUDING PREVIOUSLY TREATED AND UNTREATED, DE NOVO AND SECONDARY MDS WITH THE FOLLOWING FRENCH-AMERICAN-BRITISH SUBTYPES (REFRACTORY ANEMIA, REFRACTORY ANEMIA WITH RINGED SIDEROBLASTS, REFRACTORY ANEMIA WITH EXCESS BLASTS, AND CHRONIC MYELOMONOCYTIC LEUKEMIA [CMML]) AND INTERMEDIATE-1, INTERMEDIATE-2, AND HIGH-RISK INTERNATIONAL PROGNOSTIC SCORING SYSTEM GROUPS.
- ODE-317 FOR USE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT PATIENTS
- ODE-318 TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST
- ODE-319 INDICATED IN PEDIATRIC PATIENTS (BIRTH TO LESS THAN 18 YEARS OF AGE AND WEIGHING AT LEAST 2.5 KG) FOR THE TREATMENT OF CHAGAS DISEASE (AMERICAN TRYPANOSOMIASIS) CAUSED BY TRYPANOSOMA CRUZI
- ODE-320 INDICATED FOR CONTINUED TREATMENT OF ADULT PATIENTS WITH ACUTE MYELOID LEUKEMIA WHO ACHIEVED FIRST COMPLETE REMISSION (CR) OR COMPLETE REMISSION WITH INCOMPLETE BLOOD COUNT RECOVERY (CRI) FOLLOWING INTENSIVE INDUCTION CHEMOTHERAPY AND ARE NOT ABLE TO COMPLETE INTENSIVE CURATIVE THERAPY
- ODE-321 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 4 MONTHS TO LESS THAN 6 MONTHS WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-322 TREATMENT OF NARCOLEPSY
- ODE-323 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE A MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA
- ODE-324 TREATMENT OF HUTCHINSON-GILFORD PROGERIA SYNDROME (HGPS) AND PROGEROID LAMINOPATHIES
- ODE-325 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) HARBORING MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING ALTERATIONS
- ODE-326 TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) OR DRAVET SYNDROME (DS) IN PATIENTS BETWEEN 1 AND 2 YEARS OF AGE
- ODE-327 ADD-ON MAINTENANCE THERAPY TO IMPROVE PULMONARY FUNCTION IN ADULT PATIENTS 18 YEARS OF AGE AND OLDER WITH CYSTIC FIBROSIS AND WHO HAVE PASSED THE BRONCHITOL TOLERANCE TEST
- ODE-328 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ALK-POSITIVE
- ODE-329 FOR THE TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PEDIATRIC PATIENTS 3 TO 15 YEARS OF AGE
- ODE-330 THE TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME (SMS) IN PATIENTS 16 YEARS OF AGE AND OLDER
- ODE-331 TREATMENT OF CATAPLEXY IN ADULT PATIENTS WITH NACROLEPSY
- ODE-332 TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX (TSC) IN PATIENTS 1 YEAR OF AGE AND OLDER
- ODE-333 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-334 TREATMENT OF SPINAL MUSCULAR ATROPHY (SMA) IN PATIENTS 2 MONTHS OF AGE AND OLDER
- ODE-335 FOR TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE AT LEAST ONE OF THE ADDITIONAL MUTATIONS IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT HAVE BEEN IDENTIFIED AS RESPONSIVE TO

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TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND IDENTIFIED IN THE APPROVAL ON DECEMBER 21, 2020

- ODE-336 INDICATED FOR CHRONIC WEIGHT MANAGEMENT IN ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH OBESITY DUE TO PROOPIOMELANOCORTIN (POMC), PROPROTEIN CONVERTASE SUBTILISIN/KEXIN TYPE 1 (PCSK1), OR LEPTIN RECEPTOR (LEPR) DEFICIENCY CONFIRMED BY GENETIC TESTING DEMONSTRATING VARIANTS IN POMC, PCSK1, OR LEPR GENES THAT ARE INTERPRETED AS PATHOGENIC, LIKELY PATHOGENIC, OR OF UNCERTAIN SIGNIFICANCE (VUS)
- ODE-337 FOR ADJUVANT THERAPY AFTER TUMOR RESECTION IN ADULT PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-338 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 4 MONTHS AND OLDER WHO HAVE ONE OF THE ADDITIONAL MUTATIONS IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE THAT HAVE BEEN IDENTIFIED AS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON IN VITRO DATA AND IDENTIFIED IN THE APPROVAL ON DECEMBER 21, 2020
- ODE-339 TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1) TO LOWER URINARY OXALATE LEVELS IN PEDIATRIC AND ADULT PATIENTS
- ODE-340 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- ODE-341 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- ODE-342 INDICATED TO REDUCE THE RISK OF MORTALITY IN PATIENTS WITH MOLYBDENUM COFACTOR DEFICIENCY (MOCD) TYPE A
- ODE-343 FOR TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED REGIMEN
- ODE-344 FOR TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) WHO HAVE RECEIVED AT LEAST THREE PRIOR LINES OF SYSTEMIC THERAPY
- ODE-345 IN PEDIATRIC AND ADULT PATIENTS AS ADJUNCTIVE THERAPY TO STANDARD OF CARE FOR THE TREATMENT OF ACUTE HYPERAMMONEMIA DUE TO PROPIONIC ACIDEMIA (PA) OR METHYLMALONIC ACIDEMIA (MMA)
- ODE-346 FOR THE TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY, EXCLUDING ADULT PATIENTS COVERED BY XPOVIO'S PREVIOUS INDICATION FOR MULTIPLE MYELOMA APPROVED ON JULY 3, 2019
- ODE-347 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING
- ODE-348 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST FOUR PRIOR LINES OF THERAPY AND WHOSE DISEASE IS REFRACTORY TO AT LEAST ONE PROTEASOME INHIBITOR, ONE IMMUNOMODULATORY AGENT, AND ONE CD-38 DIRECTED MONOCLONAL ANTIBODY
- ODE-349 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST, EXCLUDING PATIENTS WHOSE DISEASE HAS PROGRESSED ON CRIZOTINIB AND AT LEAST ONE OTHER ALK INHIBITOR FOR METASTATIC DISEASE; OR ALECTINIB OR CERITINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-350 TREATMENT OF NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PEDIATRIC PATIENTS AGES 1 YEAR AND OLDER
- ODE-351 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH)
- ODE-352 TREATMENT OF ADULT PATIENTS WITH KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), AS DETERMINED BY AN FDA-APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR SYSTEMIC THERAPY
- ODE-353 FOR TREATMENT OF ADULTS WITH PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT AS DETECTED BY AN FDA-APPROVED TEST
- ODE-354 TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULT AND PEDIATRIC PATIENTS, INCLUDING NEONATES

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- ODE-355 FOR THE TREATMENT OF INVASIVE ASPERGILLOSIS IN ADULTS AND PEDIATRIC PATIENTS 13 YEARS OF AGE AND OLDER
- ODE-356 FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED SYSTEMIC MASTOCYTOSIS (ADVSM). ADVSM INCLUDES PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), AND MAST CELL LEUKEMIA (MCL)
- ODE-357 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 6 THROUGH 11 YEARS OLD WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA
- ODE-358 FOR THE TREATMENT OF VERNAL KERATOCONJUNCTIVITIS (VKC) IN CHILDREN AND ADULTS
- ODE-359 FOR THE TREATMENT OF BOTH THE FIRST-STAGE (HEMOLYMPHATIC) AND SECOND-STAGE (MENINGOENCEPHALITIC) HUMAN AFRICAN TRYPANOSOMIASIS (HAT) DUE TO TRYPANOSOMA BRUCEI GAMBIENSE IN PATIENTS 6 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 20 KG
- ODE-360 FOR PROPHYLAXIS OF ORGAN REJECTION IN ADULT AND PEDIATRIC PATIENTS RECEIVING ALLOGENEIC LUNG TRANSPLANT
- ODE-361 INDICATED FOR THE TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY
- ODE-362 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CHRONIC GVHD) AFTER FAILURE OF AT LEAST TWO PRIOR LINES OF SYSTEMIC THERAPY
- ODE-363 TREATMENT OF PRURITUS IN PATIENTS 3 MONTHS OF AGE AND OLDER WITH PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- ODE-364 TREATMENT OF ADULT PATIENTS WITH VON HIPPEL-LINDAU (VHL) DISEASE WHO REQUIRE THERAPY FOR ASSOCIATED RENAL CELL CARCINOMA (RCC), CENTRAL NERVOUS SYSTEM (CNS) HEMANGIOBLASTOMAS, OR PANCREATIC NEUROENDOCRINE TUMORS (PNET), NOT REQUIRING IMMEDIATE SURGERY
- ODE-366 INDICATED FOR THE TREATMENT OF ADULTS WITH UNRESECTABLE OR METASTATIC GIST HARBORING A PLATELET-DERIVED GROWTH FACTOR RECEPTOR ALPHA (PDGFRA) EXON 18 MUTATION, INCLUDING PDGFRA D842V MUTATIONS
- ODE-367 PEDIATRIC PATIENTS AGED 6 MONTHS AND OLDER FOR THE TREATMENT OF C. DIFFICILE-ASSOCIATED DIARRHEA (CDAD)
- ODE-368 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH AN ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-369 THE TREATMENT OF IDIOPATHIC HYPERSOMNIA (IH) IN ADULTS
- ODE-370 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
- ODE-371 TREATMENT OF ADULT PATIENTS WITH WALDENSTRM'S MACROGLOBULINEMIA (WM)
- ODE-372 FOR TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WEIGHING LESS THAN 45 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5 OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS (CHILD-PUGH A); AND TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE WEIGHING LESS THAN 45 KG WITH HCV GENOTYPE 1 INFECTION, WHO PREVIOUSLY HAVE BEEN TREATED WITH A REGIMEN CONTAINING AN HCV NS5A INHIBITOR OR AN NS3/4A PROTEASE INHIBITOR (PI), BUT NOT BOTH
- ODE-373 TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER
- ODE-374 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 20 INSERTION MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, WHOSE DISEASE HAS PROGRESSED ON OR AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-375 THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH LOCALLY ADVANCED OR METASTATIC DIFFERENTIATED THYROID CANCER (DTC) THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY OR INELIGIBLE
- ODE-376 FOR TREATMENT OF PEDIATRIC PATIENTS 3 YEARS OF AGE TO LESS THAN 6 YEARS OF AGE WEIGHING LESS THAN 17 KG WITH CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION: WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS; OR WITH

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DECOMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN

- ODE-377 AS AN ADJUNCTIVE TREATMENT OF ADULT PATIENTS WITH SEVERE ACTIVE ANTI-NEUTROPHIL CYTOPLASMIC AUTOANTIBODY (ANCA)-ASSOCIATED VASCULITIS (GRANULOMATOSIS WITH POLYANGIITIS [GPA] AND MICROSCOPIC POLYANGIITIS [MPA])
- ODE-378 A COMPLETE REGIMEN FOR THE TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTION IN PEDIATRIC PATIENTS WEIGHING 14 KG TO LESS THAN 25 KG WHO HAVE NO ANTIRETROVIRAL TREATMENT HISTORY OR TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY-SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES PER ML) ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO THE INDIVIDUAL COMPONENTS OF BIKTARVY
- ODE-379 FOR TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS WITH ALAGILLE SYNDROME (ALGS) 1 YEAR OF AGE AND OLDER
- ODE-380 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH ACCELERATED PHASE PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) WITH RESISTANCE OR INTOLERANCE TO PRIOR TYROSINE-KINASE INHIBITOR (TKI) THERAPY
- ODE-381 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP) WITH THE T315I MUTATION
- ODE-382 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML) IN CHRONIC PHASE (CP), PREVIOUSLY TREATED WITH TWO OR MORE TYROSINE KINASE INHIBITORS (TKIS)
- ODE-383 FOR USE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR THE LOCALIZATION OF KNOWN SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT AND PEDIATRIC PATIENTS
- ODE-384 TO REDUCE THE RISK OF OTOTOXICITY ASSOCIATED WITH CISPLATIN IN PEDIATRIC PATIENTS 1 MONTH OF AGE AND OLDER WITH LOCALIZED, NON-METASTATIC SOLID TUMORS
- ODE-385 FOR TREATMENT OF ENDOGENOUS HYPERCORTISOLEMIA IN ADULT PATIENTS WITH CUSHING'S SYNDROME FOR WHOM SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-386 FOR THE TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA)
- ODE-387 TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS WITH ACHONDROPLASIA WHO ARE 5 YEARS OF AGE AND OLDER WITH OPEN EPIPHYSES
- ODE-388 FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS (12 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 35 KG) WITH POST-TRANSPLANT CYTOMEGALOVIRUS (CMV) INFECTION/DISEASE THAT IS REFRACTORY TO TREATMENT (WITH OR WITHOUT GENOTYPIC RESISTANCE) WITH GANCICLOVIR, VALGANCICLOVIR, CIDOFOVIR OR FOSCARNET
- ODE-389 TO REDUCE PROTEINURIA IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) AT RISK OF RAPID DISEASE PROGRESSION, GENERALLY A URINE PROTEIN-TO-CREATININE RATIO (UPCR) > OR = 1.5 G/G
- ODE-390 AS AN ADJUNCT FOR INTRAOPERATIVE IDENTIFICATION OF MALIGNANT LESIONS IN ADULT PATIENTS WITH OVARIAN CANCER
- ODE-391 TREATMENT OF FACIAL ANGIOFIBROMA ASSOCIATED WITH TUBEROUS SCLEROSIS IN ADULTS AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER
- ODE-392 TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY
- ODE-393 TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-394 FOR TREATMENT OF SICKLE CELL DISEASE (SCD) IN PEDIATRIC PATIENTS 4 YEARS OF AGE TO LESS THAN 12 YEARS OF AGE
- ODE-395 TREATMENT OF SEIZURES ASSOCIATED WITH CYCLIN-DEPENDENT KINASE-LIKE 5 (CDKL5) DEFICIENCY DISORDER (CDD) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-396 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH SEVERE MANIFESTATIONS OF PIK3CA-RELATED OVERGROWTH SPECTRUM (PROS) WHO REQUIRE SYSTEMIC THERAPY
- ODE-397 TREATMENT OF ADULTS WITH INTERMEDIATE OR HIGH-RISK PRIMARY OR SECONDARY (POST-POLYCYTHEMIA VERA OR POST-ESSENTIAL THROMBOCYTHEMIA) MYELOFIBROSIS (MF) WITH A PLATELET COUNT BELOW $50 \times 10^9/L$
- ODE-398 TREATMENT OF ADULTS WITH SYMPTOMATIC NEW YORK HEART ASSOCIATION (NYHA) CLASS II-III OBSTRUCTIVE HYPERTROPHIC CARDIOMYOPATHY (HCM) TO IMPROVE FUNCTIONAL CAPACITY AND SYMPTOMS

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ODE-399	TREATMENT OF PEDIATRIC PATIENTS AGED ONE MONTH AND OLDER WITH NEWLY DIAGNOSED JUVENILE MYELOMONOCYTIC LEUKEMIA (JMML)
ODE-400	TREATMENT OF SPINAL MUSCULAR ATROPHY (SMA) IN PEDIATRIC PATIENTS BETWEEN BIRTH AND 2 MONTHS OF AGE
ODE-401	TREATMENT OF ADULT PATIENTS WITH STABLE WILSON'S DISEASE WHO ARE DE-COPPERED AND TOLERANT TO PENICILLAMINE
ODE-402	FOR CHRONIC WEIGHT MANAGEMENT IN ADULT AND PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WITH MONOGENIC OR SYNDROMIC OBESITY DUE TO BARDET-BIEDL SYNDROME (BBS)
ODE-403	TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME (DS) IN PATIENTS TAKING CLOBAZAM WHO ARE 6 MONTHS TO LESS THAN 2 YEARS OF AGE AND WEIGHING 7 KG OR MORE
ODE-404	TREATMENT OF ADULTS WITH RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS (MLNS) WITH FIBROBLAST GROWTH FACTOR RECEPTOR 1 (FGFR1) REARRANGEMENT
ODE-405	TREATMENT OF PEDIATRIC PATIENTS AGE 1 YEAR AND OLDER WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
ODE-406	TO IMPROVE KIDNEY FUNCTION IN ADULTS WITH HEPATORENAL SYNDROME WITH RAPID REDUCTION IN KIDNEY FUNCTION
ODE-407	TREATMENT OF ADULT AND PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH UNRESECTABLE, RECURRENT, OR REFRACTORY INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT) THAT IS ALK-POSITIVE
ODE-408	TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 1 YEAR TO LESS THAN 2 YEARS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
ODE-409	TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC SOLID TUMORS WITH A RET GENE FUSION THAT HAVE PROGRESSED ON OR FOLLOWING PRIOR SYSTEMIC TREATMENT OR WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
ODE-410	TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY TREATED, UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC INTRAHEPATIC CHOLANGIOCARCINOMA HARBORING FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) GENE FUSIONS OR OTHER REARRANGEMENTS
ODE-411	TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS
ODE-412	TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED NON-SMALL CELL LUNG CANCER (NSCLC) WITH A REARRANGED DURING TRANSFECTION (RET) GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST
ODE-413	TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
ODE-414	TREATMENT OF NEONATAL SEIZURES IN TERM AND PRETERM INFANTS
ODE-415	TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1) TO LOWER PLASMA OXALATE LEVELS IN PEDIATRIC AND ADULT PATIENTS
ODE-416	TREATMENT OF ADULT PATIENTS WITH HISTIOCYTIC NEOPLASMS
ODE-417	TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN ADULT PATIENTS WITH THALASSEMIA SYNDROMES EXCLUDING ADULT PATIENTS COVERED BY THE INDICATION FOR THALASSEMIA SYNDROMES APPROVED ON OCTOBER 14, 2011
ODE-418	TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH THALASSEMIA SYNDROMES
ODE-419	TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN ADULT AND PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER WITH SICKLE CELL DISEASE OR OTHER ANEMIAS
ODE-420	TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN PEDIATRIC PATIENTS 8 YEARS OF AGE AND OLDER WITH THALASSEMIA SYNDROMES
ODE-421	TREATMENT OF TRANSFUSIONAL IRON OVERLOAD IN ADULT AND PEDIATRIC PATIENTS 8 YEARS OF AGE AND OLDER WITH SICKLE CELL DISEASE OR OTHER ANEMIAS
ODE-422	TREATMENT OF ADULT PATIENTS WITH RAS WILD-TYPE, HER2-POSITIVE UNRESECTABLE OR METASTATIC COLORECTAL CANCER THAT HAS PROGRESSED FOLLOWING TREATMENT WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-BASED CHEMOTHERAPY
ODE-423	FOR PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT RECIPIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE [D+/R-])
ODE-424	TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MANTLE CELL LYMPHOMA (MCL) AFTER AT LEAST TWO LINES OF SYSTEMIC THERAPY, INCLUDING A BTK INHIBITOR
ODE-425	TREATMENT OF RETT SYNDROME IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND

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OLDER

- ODE-426 FOR USE IN PATIENTS 18 YEARS OF AGE OR OLDER WHO HAVE LIMITED OR NO ALTERNATIVE OPTIONS FOR THE TREATMENT OF CANDIDEMIA AND INVASIVE CANDIDIASIS
- ODE-427 TREATMENT OF FRIEDREICH'S ATAXIA IN ADULTS AND ADOLESCENTS AGED 16 YEARS AND OLDER
- ODE-428 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA (LGG) WITH A BRAF V600E MUTATION WHO REQUIRE SYSTEMIC THERAPY
- ODE-429 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 3 MONTHS OF AGE TO LESS THAN 1 YEAR OF AGE WITH ALAGILLE SYNDROME (ALGS)
- ODE-430 TREATMENT OF ACTIVATED PHOSPHOINOSITIDE 3-KINASE DELTA (PI3K DELTA) SYNDROME (APDS) IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- ODE-431 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN ADULTS WITH NARCOLEPSY
- ODE-432 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS WHO HAVE A MUTATION IN THE SUPEROXIDE DISMUTASE 1 (SOD1) GENE
- ODE-433 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGED 2 YEARS TO LESS THAN 6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA
- ODE-434 TREATMENT OF ADULT PATIENTS WITH INDOLENT SYSTEMIC MASTOCYTOSIS (ISM)
- ODE-435 THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS 1 MONTH TO LESS THAN 4 MONTHS OF AGE WHO HAVE AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-436 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS 12 MONTHS OF AGE AND OLDER WITH ALAGILLE SYNDROME (ALGS)
- ODE-437 FOR USE IN COMBINATION WITH STANDARD INDUCTION AND CONSOLIDATION, AND AS MAINTENANCE THERAPY FOLLOWING CONSOLIDATION CHEMOTHERAPY, FOR THE TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) THAT IS FLT3 INTERNAL TANDEM DUPLICATION (ITD)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- ODE-438 AS A LIVER-DIRECTED TREATMENT FOR ADULT PATIENTS WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES AFFECTING LESS THAN 50% OF THE LIVER AND NO EXTRAHEPATIC DISEASE OR EXTRAHEPATIC DISEASE LIMITED TO THE BONE, LYMPH NODES, SUBCUTANEOUS TISSUES, OR LUNG THAT IS AMENABLE TO RESECTION OR RADIATION
- ODE-439 FOR THE REDUCTION IN VOLUME OF NEW HETEROTOPIC OSSIFICATION IN ADULTS AND PEDIATRIC PATIENTS AGED 8 YEARS AND OLDER FOR FEMALES AND 10 YEARS AND OLDER FOR MALES WITH FIBRODYSPLASIA OSSIFICANS PROGRESSIVA (FOP)
- ODE-440 FOR TREATMENT OF ADULTS WITH CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
- ODE-441 TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS (MF), INCLUDING PRIMARY MF OR SECONDARY MF [POST-POLYCYTHEMIA VERA (PV) AND POST-ESSENTIAL THROMBOCYTHEMIA (ET)], IN ADULTS WITH ANEMIA
- ODE-442 TO MOBILIZE HEMATOPOIETIC STEM CELLS TO THE PERIPHERAL BLOOD FOR COLLECTION AND SUBSEQUENT AUTOLOGOUS TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- ODE-443 TO LOWER URINARY OXALATE LEVELS IN CHILDREN 9 YEARS OF AGE AND OLDER AND ADULTS WITH PRIMARY HYPEROXALURIA TYPE 1 (PH1) AND RELATIVELY PRESERVED KIDNEY FUNCTION, E.G., EGFR GREATER THAN OR EQUAL TO 30 ML/MIN/1.73 M²
- ODE-444 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+ CML), NEWLY-DIAGNOSED OR RESISTANT OR INTOLERANT TO PRIOR THERAPY
- ODE-445 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-446 TREATMENT OF GENERALIZED MYASTHENIA GRAVIS (GMG) IN ADULT PATIENTS WHO ARE ANTI-ACETYLCHOLINE RECEPTOR (ACHR) ANTIBODY POSITIVE
- ODE-447 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES (MDS) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-448 TREATMENT OF PEDIATRIC PATIENTS OLDER THAN 1 MONTH UP TO 12 YEARS OF AGE WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST WITHOUT A KNOWN ACQUIRED RESISTANCE MUTATION, ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN

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SEVERE MORBIDITY, AND HAVE PROGRESSED FOLLOWING TREATMENT OR HAVE NO SATISFACTORY ALTERNATIVE THERAPY

ODE-449 TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS LESS THAN 5 YEARS OF AGE WITH ACHONDROPLASIA WITH OPEN EPIPHYSES

ODE-450 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS 2 YEARS OF AGE AND OLDER

ODE-451 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA (CLL/SLL) WHO HAVE RECEIVED AT LEAST TWO PRIOR LINES OF THERAPY, INCLUDING A BTK INHIBITOR AND A BCL-2 INHIBITOR

ODE-452 FOR ADULT PATIENTS WITH PROGRESSING DESMOID TUMORS WHO REQUIRE SYSTEMIC TREATMENT

ODE-453 TREATMENT OF INVASIVE MUCORMYCOSIS IN PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER

ODE-454 TREATMENT OF INVASIVE MUCORMYCOSIS IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WHO WEIGH 16 KG AND GREATER

ODE-455 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ROS1-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) WITH ADENOCARCINOMA HISTOLOGY

ODE-456 TREATMENT OF ADULTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH)

ODE-457 TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS UNDER 12 YEARS OF AGE WEIGHING AT LEAST 14 KG TO LESS THAN 25 KG

ODE-458 TREATMENT OF INVASIVE ASPERGILLOSIS IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER WHO WEIGH 16 KILOGRAMS (KG) AND GREATER

ODE-459 TREATMENT OF INVASIVE ASPERGILLOSIS IN PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER

PATENT USE

U-1 PREVENTION OF PREGNANCY

U-2 TREATMENT OR PROPHYLAXIS OF ANGINA PECTORIS AND ARRHYTHMIA

U-3 TREATMENT OF HYPERTENSION

U-4 PROVIDING PREVENTION AND TREATMENT OF EMESIS AND NAUSEA IN MAMMALS

U-5 METHOD OF PRODUCING BRONCHODILATION

U-6 METHOD OF PRODUCING SYMPATHOMIMETIC EFFECTS

U-7 INCREASING CARDIAC CONTRACTILITY

U-8 ACUTE MYOCARDIAL INFARCTION

U-9 CONTROL OF EMESIS ASSOCIATED WITH ANY CANCER CHEMOTHERAPY AGENT

U-10 DIAGNOSTIC METHOD FOR DISTINGUISHING BETWEEN HYPOTHALAMIC MALFUNCTIONS OR LESIONS IN HUMANS

U-11 TREATMENT OR PROPHYLAXIS OF CARDIAC DISORDERS

U-12 METHOD OF TREATING [A] HUMAN SUFFERING FROM DEPRESSION

U-13 A METHOD FOR TREATING ANXIETY IN A HUMAN SUBJECT IN NEED OF SUCH TREATMENT

U-14 ADJUNCTIVE THERAPY FOR THE PREVENTION AND TREATMENT OF HYPERAMMONEMIA IN THE CHRONIC MANAGEMENT OF PATIENTS WITH UREA CYCLE ENZYMOPATHIES

U-15 METHOD OF LOWERING INTRAOCULAR PRESSURE

U-16 USE IN LUNG SCANNING PROCEDURES

U-17 TREATMENT OF VENTRICULAR AND SUPRAVENTRICULAR ARRHYTHMIAS

U-18 METHOD FOR INHIBITING GASTRIC SECRETION IN MAMMALS

U-19 TREATMENT OF INFLAMMATION

U-20 A PROCESS FOR TREATING A PATIENT SUFFERING FROM PARKINSON'S SYNDROME AND IN NEED OF TREATMENT

U-21 TREATMENT OF HUMANS SUFFERING UNDESIRE UROTOXIC SIDE EFFECTS CAUSED BY CYTOSTATICALLY ACTIVE ALKYLATING AGENTS

U-22 METHOD OF COMBATTING PATHOLOGICALLY REDUCED CEREBRAL FUNCTIONS AND PERFORMANCE WEAKNESSES, CEREBRAL INSUFFICIENCY AND DISORDERS IN CEREBRAL CIRCULATION AND

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

METABOLISM IN WARM-BLOODED ANIMALS

- U-23 METHOD FOR TREATING PROSTATIC CARCINOMA COMPRISING ADMINISTERING FLUTAMIDE
- U-24 METHOD FOR TREATING PROSTATE ADENOCARCINOMA COMPRISING ADMINISTERING AN ANTIANDROGEN INCLUDING FLUTAMIDE AND AN LHRH AGONIST
- U-25 REDUCING CHOLESTEROL IN CHOLELITHIASIS PATIENTS
- U-26 REDUCING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-27 DISSOLVING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-28 CEREBRAL, CORONARY, PERIPHERAL, VISCERAL AND RENAL ARTERIOGRAPHY, AORTOGRAPHY AND LEFT VENTRICULOGRAPHY
- U-29 CT IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY
- U-30 CEREBRAL ANGIOGRAPHY, AND VENOGRAPHY
- U-31 INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY
- U-32 PALLIATIVE TREATMENT OF PATIENTS WITH OVARIAN CARCINOMA RECURRENT AFTER PRIOR CHEMOTHERAPY, INCLUDING PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED WITH CISPLATIN
- U-33 TREATING VIRAL INFECTIONS IN A MAMMAL
- U-34 TREATING VIRAL INFECTIONS IN A WARM-BLOODED ANIMAL
- U-35 TREATING CYTOMEGALOVIRUS IN A HUMAN WITH AN INJECTABLE COMPOSITION
- U-36 METHODS OF TREATING BACTERIAL ILLNESSES
- U-37 METHOD OF TREATING GASTROINTESTINAL DISEASE
- U-38 TREATMENT OF PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA
- U-39 ANGINA PECTORIS
- U-40 METHOD OF TREATMENT OF BURNS
- U-41 METHOD OF TREATING CARDIAC ARRHYTHMIAS
- U-42 ADJUVANT TREATMENT IN COMBINATION WITH FLUOROURACIL AFTER SURGICAL RESECTION IN PATIENTS WITH DUKES' STAGE C COLON CANCER
- U-43 MANAGEMENT OF CHRONIC PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA
- U-44 RELIEF OF NAUSEA AND VOMITING
- U-45 TREATMENT OF INFLAMMATION AND ANALGESIA
- U-46 TREATMENT OF PANIC DISORDER
- U-47 STIMULATION OF THE RELEASE OF GROWTH HORMONE
- U-48 ANALGESIA
- U-49 SYMPTOMATIC CANCER-RELATED HYPERCALCEMIA
- U-50 USE IN TREATING INFLAMMATORY DERMATOSES
- U-51 BLOOD POOL IMAGING, INCLUDING CARDIAC FIRST PASS AND GATED EQUILIBRIUM IMAGING AND FOR DETECTION OF SITES OF GASTROINTESTINAL BLEEDING
- U-52 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER SIX MONTHS OF AGE) WITH ADVANCED HIV INFECTION
- U-53 HYPERCALCEMIA OF MALIGNANCY
- U-54 REVERSAL AGENT OR ANTAGONIST OF NONDEPOLARIZING NEUROMUSCULAR BLOCKING AGENTS
- U-55 TREATMENT OF PAIN
- U-56 AID TO SMOKING CESSATION
- U-57 OPHTHALMIC USE OF NORFLOXACIN
- U-58 METHOD OF TREATING INFLAMMATORY INTESTINAL DISEASES
- U-59 METHOD OF TREATING HYPERCHOLESTEROLEMIA
- U-60 NASAL ADMINISTRATION OF BUTORPHANOL
- U-61 CEREBRAL AND PERIPHERAL ARTERIOGRAPHY AND CT IMAGING OF THE HEAD
- U-62 CORONARY ARTERIOGRAPHY, LEFT VENTRICULOGRAPHY, CT IMAGING OF THE BODY, INTRAVENOUS EXCRETORY UROGRAPHY, INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY AND VENOGRAPHY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-63 ISOPRENALINE ANTAGONISM ON THE HEART RATE OR BLOOD PRESSURE

U-64 TREATMENT OF VIRAL INFECTIONS

U-65 METHOD OF TREATMENT OF A PATIENT INFECTED WITH HIV

U-66 TRIPHASIC REGIMEN

U-67 METHOD OF INDUCING ANESTHESIA IN A WARM BLOODED ANIMAL

U-68 TREATMENT OF ACTINIC KERATOSIS

U-69 TREATMENT OF PNEUMOCYSTIS CARINII INFECTIONS

U-70 TREATMENT OF TRANSIENT INSOMNIA

U-71 METHOD OF TREATMENT OF HEART FAILURE

U-72 TREATMENT OF MIGRAINE

U-73 METHOD OF TREATING DISEASES OR INFECTIONS CAUSED BY MYCETES

U-74 METHOD OF PROVIDING HYPNOTIC EFFECT

U-75 RELIEF OF OCULAR ITCHING DUE TO SEASONAL ALLERGIC CONJUNCTIVITIS

U-76 USE TO IMAGE A SUBJECT WITH A MAGNETIC RESONANCE IMAGING SYSTEM

U-77 TREATMENT OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

U-78 ULCERATIVE COLITIS

U-79 SYMPTOMATIC TREATMENT OF PATIENTS WITH NOCTURNAL HEARTBURN DUE TO GERD

U-80 METHOD OF TREATING OCULAR BACTERIAL INFECTIONS

U-81 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS

U-82 TREATMENT FOR DEMENTIA IN PATIENTS WITH ALZHEIMER'S DISEASE

U-83 TREATMENT OF SEIZURES

U-84 A METHOD OF BLOCKING THE UPTAKE OF MONOAMINES BY BRAIN NEURONS IN ANIMALS

U-85 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-86 METHOD OF TREATING CERTAIN FORMS OF EPILEPSY

U-87 METHOD FOR NONINVASIVE ADMINISTRATION OF SEDATIVES, ANALGESICS, AND ANESTHETICS

U-88 TREATMENT OF MODERATE PLAQUE PSORIASIS

U-89 TREATMENT OR PROPHYLAXIS OF EMESIS

U-90 TREATMENT OF PSYCHOTIC DISORDERS

U-91 ALTERNATIVE THERAPY TO TRIMETHOPRIM-SULFAMETHOXAZOLE FOR TREATMENT OF MODERATE-TO-SEVERE PNEUMOCYSTIS CARINII PNEUMONIA IN IMMUNOCOMPROMISED AND AIDS PATIENTS

U-92 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN DEPENDENT DIABETES MELLITUS AND RETINOPATHY

U-93 USE AS AN ANTIHISTAMINE/DECONGESTANT

U-94 TREATMENT-ADULTS W/ ADVANCED HIV, INTOLERANT OF APPROVED THERAPIES, INTOLERANT OF APPROVED THERAPIES W/PROVEN BENEFIT OR HAVE EXPERIENCED CLINICAL/IMMUNOLOGICAL DETERIORATION WHILE RECEIVING..OR FOR WHOM SUCH THERAPIES-CONTRAINDICATED

U-95 SHORT TERM MANAGEMENT OF MODERATE PRURITIS IN ADULTS WITH ATOPIC DERMATITIS AND LICHEN SIMPLEX CHRONICUS

U-96 METHOD OF TREATING VARICELLA ZOSTER (SHINGLES) INFECTIONS

U-97 A METHOD OF TREATING A PATIENT IN NEED OF MEMORY ENHANCEMENT

U-98 A METHOD OF INDUCING REGRESSION OF LEUKEMIA CELL GROWTH IN A MAMMAL

U-99 METHOD OF PROVIDING POTASSIUM TO A SUBJECT IN NEED OF POTASSIUM

U-100 METHOD OF TREATING OCULAR INFLAMMATION

U-101 ADJUNCT TO CONVENTIONAL CT OR MRI IMAGING IN THE LOCALIZATION OF STROKE IN PATIENTS IN WHOM STROKE HAS ALREADY BEEN DIAGNOSED

U-102 METHOD OF HORMONALLY TREATING MENOPAUSAL OR POST-MENOPAUSAL DISORDERS IN WOMEN

U-103 TREATMENT OF OCULAR HYPERTENSION

U-104 TREATMENT OF AQUEOUS HUMOR FORMATION AND INTRAOCULAR PRESSURE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-105 EMESIS

U-106 TREATMENT OF EPILEPSY

U-107 TREATMENT OF HYPERTENSION AND ANGINA PECTORIS

U-108 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER, GASTROESOPHAGEAL REFLUX DISEASE (GERD), SEVERE EROSIIVE ESOPHAGITIS, POORLY RESPONSIVE SYMPTOMATIC GERD AND PATHOLOGICAL HYPERSECRETORY CONDITIONS AND MAINTENANCE HEALING OF EROSIIVE ESOPHAGITIS

U-109 ADJUNCT DIET IN THE TX OF ELEVATED TOTAL CHOLESTEROL AND LDL-C LEVELS IN PTS W/PRIMARY HYPERCHOLESTEROLEMIA WHOSE RESPONSE TO DIETARY RESTRICTION OF SAT FAT AND CHOLESTEROL AND OTHER NONPHARMACOLOGICAL MEASURES HAS NOT BEEN ADEQUATE

U-110 USE AS A RETRIEVABLE PESSARY

U-111 DIABETES

U-112 CONTRACEPTION

U-113 METHOD OF CONDUCTING RADIOLOGICAL EXAMINATION OF A PATIENT BY ADMINISTERING TO SAID PATIENT A RADIOPAQUE AMOUNT OF IOPROMIDE

U-114 USE FOR INHIBITING BONE RESORPTION

U-115 USE OF VASODILATORS TO EFFECT AND ENHANCE AN ERECTION (AND THUS TREAT ERECTILE DYSFUNCTION), BY INJECTION INTO THE PENIS

U-116 METHOD OF MYOCARDIAL IMAGING

U-117 TREATMENT OF OCULAR ALLERGIC RESPONSE IN HUMAN EYES

U-118 METHOD OF LOWERING BLOOD SUGAR LEVEL

U-119 TREATMENT OF NASAL HYPERSECRETION

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- U-210 METHOD OF TREATING CONGESTIVE HEART FAILURE
- U-211 USE IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
- U-212 METHOD OF TREATMENT OF PARKINSON'S DISEASE
- U-213 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS AND TREATING HYPERCHOLESTEROLEMIA AND METHOD FOR TREATING HYPERLIPIDEMIA
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PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-215 TREATMENT OF EPILEPSY TWICE DAILY. TREATING A PATIENT BY ADMINISTERING CARBAMAZEPINE IN A DOSAGE FORM CAPABLE OF MAINTAINING BLOOD CONCENTRATION FROM 4-12MCG/ML OVER 12 HOURS

U-216 TREATMENT OF ADENOCARCINOMA, INCLUDING STAGE B2-C BY ADMINISTERING AN AGONIST OF LH-RH AND FLUTAMIDE

U-217 METHOD OF PRODUCING ANESTHESIA

U-218 METHOD FOR LIMITING THE POTENTIAL FOR MICROBIAL GROWTH IN THE DRUG PRODUCT

U-219 TREATMENT OF PARKINSON'S DISEASE

U-220 METHOD OF DIAGNOSIS

U-221 SELECTIVE VASODILATION BY CONTINUOUS ADENOSINE INFUSION

U-222 METHOD OF TREATING PAGET'S DISEASE USING ACTONEL

U-223 TREATMENT OF BACTERIAL CONJUNCTIVITIS CAUSED BY SUSCEPTIBLE STRAINS OF MICROORGANISMS

U-224 CONTROLLING INTRAOCULAR PRESSURE

U-225 METHOD FOR DELIVERY

U-226 METHOD OF ENHANCING THE DISSOLUTION PROFILE OF A PHARMACEUTICAL FROM A SOLID DOSAGE FORM CONTAINING THE PHARMACEUTICAL AND SIMETHICONE

U-227 NASAL ADMINISTRATION

U-228 ASTHMA

U-229 CARDIAC INSUFFICIENCY (CONGESTIVE HEART FAILURE)

U-230 PREVENTION OF ACUTE CARDIAC ISCHEMIC EVENTS

U-231 USE IN PARKINSON'S DISEASE

U-232 METHOD OF TREATING MIGRAINE

U-233 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE

U-234 METHOD OF USING RIBAVIRIN TO TREAT VIRAL INFECTIONS IN MAMMALS

U-235 METHOD OF MODULATING TH1 AND TH2 RESPONSE IN ACTIVATED T CELLS OF A HUMAN COMPRISING ADMINISTERING RIBAVIRIN TO THE T CELLS IN A DOSAGE WHICH PROMOTES THE TH1 RESPONSE AND SUPPRESSES THE TH2 RESPONSE

U-236 TREATING MALE PATTERN BALDNESS WITH 0.05 TO 3.0MG/DAY

U-237 METHOD OF PERFORMING NMR IMAGING WITH A PATIENT COMPRISING ADMINISTERING TO THE PATIENT AN EFFECTIVE AMOUNT OF CONTRAST AGENT DISCLOSED IN THE CLAIMS

U-238 IMAGING A BODY TISSUE AND SUBJECTING TO NMR TOMOGRAPHY, ADMINISTERING AN AMOUNT OF PHARMACEUTICAL AGENT FOR AFFECTING THE RELAXATION TIMES OF ATOMS IN BODY TISSUES UNDERGOING NMR DIAGNOSIS, WHEREBY THE IMAGE CONTRAST IS ENHANCED....

U-239 TREATING OR CONTROLLING OCULAR INFLAMMATION WHICH COMPRISES TOPICALLY ADMINISTERING TO AFFECTED EYE A COMPOSITION COMPRISING AN NSAID, A POLYMERIC QUATERNARY AMMONIUM COMPOUND AND BORIC ACID

U-240 TREATMENT OF ACUTE MIGRAINE ATTACKS

U-241 FOR SHORT-TERM TREATMENT ACTIVE DUODENAL ULCER, MAINTENANCE THERAPY FOR DUODENAL ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING OF ACTIVE ULCER, SHORT-TERM TREATMENT ACTIVE BENIGN GASTRIC ULCER & GERD, PATHOLOGICAL HYPERSECRETORY CONDITIONS

U-242 USE OF FOLLITROPIN ALPHA ALONE IN IN-VITRO FERTILIZATION

U-243 TOPICAL ADMINISTRATION

U-244 PLATELET AGGREGATION INHIBITORS

U-245 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS

U-246 PHOSPHATE BINDING

U-247 TREATMENT OF RHEUMATOID ARTHRITIS

U-248 TREATMENT OF HIV

U-249 METHOD OF TREATING ALLERGIC OR NON-ALLERGIC RHINITIS IN PATIENTS BY ADMINISTERING AEROSOLIZED PARTICLES OF MOMETASONE FUROATE

U-250 TREATMENT OF HEPATITIS B INFECTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-251 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS IN THE TREATMENT OF TYPE II DIABETES

U-252 METHOD OF TREATING A HUMAN SUBJECT HAVING GAUCHER'S DISEASE

U-253 ORAL TRANSMUCOSAL USE

U-254 USE OF AGGRASTAT IN COMBINATION WITH HEPARIN

U-255 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY

U-256 TREATMENT OF HIV INFECTION IN COMBINATION WITH ONE OR MORE ADDITIONAL HIV ANTIVIRAL AGENTS

U-257 TREATMENT OF HIV INFECTION

U-258 TREATMENT OF NEURODEGENERATIVE DISEASES

U-259 TREATMENT OF ANDROGENIC ALOPECIA BY ORAL ADMINISTRATION DRUG SUBSTANCE

U-260 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA AND OCULAR HYPERTENSION WHO ARE INTOLERANT OF OTHER IOP LOWERING MEDICATIONS OR INSUFFICIENTLY RESPONSIVE TO ANOTHER IOP LOWERING MEDICATION

U-261 TREATING BENIGN PROSTATIC HYPERPLASIA WITH A GENUS OF COMPOUNDS, INCLUDING FINASTERIDE

U-262 TREATING BENIGN PROSTATIC HYPERTROPHY WITH FINASTERIDE

U-263 METHOD OF TREATING A MALIGNANT CONDITION THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING LEUKEMIA OR LYMPHOMA IN A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVENOUS ADMINISTRATION OF BUSULFAN

U-264 METHOD OF TREATING A MALIGNANT DISEASE THROUGH PARENTERAL ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN

U-265 USE AS LAXATIVE

U-266 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS; RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS; MANAGEMENT OF ACUTE PAIN IN ADULTS; TREATMENT OF PRIMARY DYSMENORRHEA; ACUTE TREATMENT OF MIGRAINE ATTACKS IN ADULTS

U-267 PREVENTING HEARTBURN EPISODES FOLLOWING INGESTION OF HEARTBURN-INDUCING FOOD/BEVERAGE, COMPRISING ADMIN TO PT, 30 MIN PRIOR TO CONSUMPTION BY THE PT THE FOOD/BEVERAGE, A COMPOSITION COMPRISING 10MG FAMOTIDINE

U-268 ACROMEGALY

U-269 EXCESS GH-SECRETION OR GASTRO-INTESTINAL DISORDERS

U-270 METHOD OF IMPROVING THE TIME FOR ADMINISTRATION OR THE TIME BETWEEN CHANGES OF GIVING SETS FOR THE DRUG PRODUCT

U-271 METHOD OF TREATING TUMORS

U-272 METHOD OF TREATING CARCINOMA

U-273 CUTANEOUS T-CELL LYMPHOMA

U-274 ZANAMIVIR FOR INHALATION

U-275 METHOD OF USE OF THE DRUG SUBSTANCE

U-276 METHOD OF USE OF LEVOBUPIVACAINE

U-277 NEUROLOGICAL AND OTHER DISORDERS (TREATMENT OF EPILEPSY, BID ORAL DOSING)

U-278 METHOD OF USE OF THE INDICATION OF THE DRUG PRODUCT

U-279 METHOD OF USE OF THE APPROVED PRODUCT

U-280 TREATING PRECIPITATED ACUTE URINARY RETENTION WITH FINASTERIDE

U-281 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS

U-282 METHOD OF TREATING BACTERIAL INFECTIONS

U-283 METHOD FOR TREATING MENOPAUSAL SYMPTOMS IN A POSTMENOPAUSAL FEMALE

U-284 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE, AND VULVAR AND VAGINAL ATROPHY) AND OSTEOPOROSIS

U-285 DEPRESSION AND SOCIAL ANXIETY DISORDER/SOCIAL PHOBIA

U-286 DEPRESSION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-287 TREATMENT OR PREVENTION OF OSTEOPOROSIS

U-288 THERAPY OF INFLUENZA

U-289 TREATMENT OF NON-HYPERKERATOTIC ACTINIC KERATOSES OF FACE AND SCALP

U-290 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS)

U-291 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH CYCLOSPORIN

U-292 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH AZATHIOPRINE

U-293 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH A CORTICOSTEROID

U-294 TREATMENT OF HYPERPIGMENTARY DISORDERS

U-295 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-296 TREATING MIGRAINE PAIN AND ONE OR MORE OF A CLUSTER OF SYMPTOMS CHARACTERISTIC OF A MIGRAINE ATTACK SYMPTOMS BEING SELECTED FROM PHOTOPHOBIA, PHONOPHOBIA NAUSEA AND FUNCTIONAL DISABILITY

U-297 PREVENTION OR TREATMENT OF REVERSIBLE VASOCONSTRICTION BY THE INHALATION OF NITRIC OXIDE WITH AN OXYGEN CONTAINING GAS

U-298 METHOD OF COMBATING BACTERIA IN A PATIENT

U-299 TREATMENT OF ADENOMATOUS POLYPS

U-300 INDICATED FOR THE REDUCTION OF ELEVATED TOTAL AND LDL CHOLESTEROL LEVELS IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA

U-301 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS AND BIGUANIDES IN THE TREATMENT OF TYPE II DIABETES

U-302 TO REDUCE THE RISK OF STROKE IN PATIENTS WHO HAVE HAD TRANSIENT ISCHEMIA OF THE BRAIN OR COMPLETED ISCHEMIC STROKE DUE TO THROMBOSIS

U-303 METHOD OF USE PATENT-PRODUCT APPROVED FOR TREATMENT OF OSTEOPOROSIS, PAGET'S DISEASE, PREVENTION AND TREATMENT OF GLUCOCORTICOID INDUCED OSTEOPOROSIS

U-304 A METHOD OF TREATMENT OF A CONDITION INVOLVING AN ANTIBODY ANTIGEN REACTION

U-305 METHODS FOR USING THE DRUG PRODUCT

U-306 TREATMENT OF POST-MENOPAUSAL UROGENITAL SYMPTOMS ASSOCIATED WITH ESTROGEN DEFICIENCY

U-307 CLAIMS AN OLANZAPINE POLYMORPH USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATION OF THIS NDA

U-308 CLAIMS A SOLID ORAL FORMULATION INCLUDING TABLETS AND GRANULES OF OLANZAPINE USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATIONS OF THIS NDA

U-309 TREATING SJOEGREN SYNDROME

U-310 TREATMENT OF XEROSTOMIA

U-311 HORMONE REPLACEMENT

U-312 PANIC DISORDER, OBSESSIVE-COMPULSIVE DISORDER, POSTTRAUMATIC STRESS DISORDER

U-313 TREATMENT OF CONGESTIVE HEART FAILURE

U-314 METHOD FOR TREATING HYPERPARATHYROIDISM WHICH COMPRISES SUPPRESSING PARATHYROID ACTIVITY

U-315 METHOD FOR ADMINISTERING DRUG TO GASTROINTESTINAL TRACT

U-316 METHOD OF TREATING A SUBJECT SUFFERING FROM PROSTATE CANCER

U-317 METHOD OF USING TROGLITAZONE TO TREAT PATIENTS HAVING INSULIN RESISTANCE

U-318 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE

U-319 TREATMENT OF MICROBIAL INFECTIONS

U-320 INHIBITING OR ELIMINATING ACUTE MYELOID LEUKEMIA

U-321 REDUCTION OF ELEVATED IPTH LEVELS IN THE MGT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS UNDERGONG CHRONIC RENAL DIALYSIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-322 TREATMENT OF ALZHEIMER'S DEMENTIA

U-323 USE AS A BILE ACID SEQUESTRANT

U-324 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN, SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS OR ACUTE MANIA EMPLOYING OLANZAPINE

U-325 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS, INCLUDING "BIPOLAR DISORDER NOS" EMPLOYING OLANZAPINE

U-326 METHOD OF TREATING SCHIZOPHRENIA AND BIPOLAR DISORDER

U-327 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED PSYCHOTIC CONDITIONS EMPLOYING OLANZAPINE

U-328 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS INCLUDING "A PSYCHOTIC CONDITION" EMPLOYING AN OLANZAPINE POLYMORPH

U-329 USE OF AVANDIA AS MONOTHERAPY, IN COMBINATION WITH METFORMIN, AND IN COMBINATION WITH SULFONYLUREAS TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS

U-330 TREATMENT OF NAUSEA AND VOMITING

U-331 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-332 TREATMENT OR PREVENTION OF BRONCHOSPASM

U-333 METHOD OF TREATING OCULAR HYPERTENSION

U-334 TREATMENT OF EXCESSIVE FEMALE FACIAL HAIR

U-335 USE OF PRAVASTATIN SODIUM FOR SECONDARY PREVENTION OF CORONARY EVENTS IN MEN AND WOMEN WHO HAVE HAD A MYOCARDIAL INFARCTION AND HAVE NORMAL CHOLESTEROL LEVELS

U-336 DIAGNOSTIC RADIOIMAGING

U-337 USE OF CARDIOLITE/MIRALUMA KIT FOR THE PREPARATION OF TC99M SESTAMIBI

U-338 METHODS FOR TREATING DISTURBANCES OF MOOD, DISTURBANCES OF APPETITE, DEPRESSED MOOD, OR CARBOHYDRATE CRAVING ALL ASSOCIATED WITH PREMENSTRUAL SYNDROME

U-339 PREVENTION OF CARDIO-TOXICITY CAUSED BY THE ADMINISTRATION OF DOXORUBICIN

U-340 THE LONG TERM TREATMENT OF GROWTH FAILURE DUE TO LACK OF ADEQUATE ENDOGENOUS GROWTH HORMONE SECRETION IN CHILDREN

U-341 METHOD FOR ENHANCING THE TREATMENT OF ... LATE LUTEAL PHASE DYSPHORIC DISORDER

U-342 METHOD FOR TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER

U-343 REDUCTION OF INTESTINAL GAS, CRAMPING AND ANORECTAL IRRITATION

U-344 METHOD FOR INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH ANOTHER HIV PROTEASE INHIBITOR

U-345 RITONAVIR AND ANOTHER HIV PROTEASE INHIBITOR FOR CONCOMITANT ADMINISTRATION FOR THE TREATMENT OF AN HIV INFECTION

U-346 METHOD FOR INHIBITING CYTOCHROME P450 MONOOXYGENASE WITH RITONAVIR AND A METHOD FOR IMPROVING THE PHARMACOKINETICS OF A DRUG THAT IS METABOLIZED BY CYTOCHROME P450 MONOOXYGENASE BY ADMIN THE DRUG AND RITONAVIR

U-347 METHOD OF USE IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS

U-348 METHOD OF USE FOR INHIBITING HIV INFECTION

U-349 METHOD OF USE WHICH IS SUBJECT OF THE APPLICATION

U-350 PREPARATION OF A PHARMACEUTICAL COMPOSITION FOR CONCOMITANT ADMIN WITH A REVERSE TRANSCRIPTASE INHIBITOR

U-351 INHIBITING PROTEASE WITH LOPINAVIR AND INHIBITING AN HIV INFECTION WITH LOPINAVIR

U-352 INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH A REVERSE TRANSCRIPTASE INHIBITOR

U-353 PREVENTION AND TREATMENT OF OSTEOPOROSIS

U-354 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-355 METHOD OF ASSISTING PERSON TO QUIT SMOKING...TRANSDERMALLY ADMIN NICOTINE VIA..PATCH ADHERED TO SKIN AT DOSING RATE APPROX SAME AS ABSORBED FROM SMOKING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-356 DELIVERING A MEDICINAL AEROSOL FORMULATION USING CFC-FREE PROPELLANT 134A.

U-357 USE OF THE DRUG PRODUCT IN PHOTODYNAMIC THERAPEUTIC PROTOCOLS FOR THE TREATMENT OF AGE-RELATED MACULAR DEGENERATION AND RELATED CONDITIONS INVOLVING UNWANTED NEOVASCULATURE IN THE EYE

U-358 DEPRESSION, OBSESSIVE COMPULSIVE DISORDER, PANIC DISORDER AND SOCIAL ANXIETY DISORDER

U-359 METHOD OF USE OF VISICOL

U-360 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS INCLUDING MENTAL DISORDERS EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011

U-361 MANAGEMENT OF ANXIETY DISORDERS AND THE SHORT-TERM RELIEF OF THE SYMPTOMS OF ANXIETY

U-362 USE OF APPROVED FORMULATIONS TO TREAT ALL APPROVED DISEASE INDICATIONS

U-363 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS THAT RELATE TO THE USE OF A PSYCHOACTIVE SUBSTANCE EMPLOYING OLANZAPINE AS PER THE INDICATION THE SUBJECT MATTER OF SUPPLEMENT 011

U-364 TREATING A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ANY NUMBER OF LISTED CONDITIONS INCLUDING PSYCHOSIS, EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011

U-365 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN OUR EXTENDED, CONTROLLED RELEASE FORMULATION

U-366 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN A DELAYED RELEASE FORMULATION

U-367 TREATMENT OF CARDIOVASCULAR DISORDERS

U-368 HEARTBURN

U-369 METHOD OF CONTROLLING AND LOWERING INTRAOCULAR PRESSURE

U-370 INTRAVAGINAL TREATMENT OF VAGINAL INFECTIONS WITH BUFFERED METRONIDAZOLE COMPOSITIONS

U-371 APPROVAL FOR MARKETING ONLY UNDER A SPECIAL RESTRICTION PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)

U-372 METHOD FOR ADMINISTERING A BENEFICIAL DRUG TO THE GI TRACT OF AN ANIMAL, WHICH METHOD COMPRISES ADMITTING AN OSMOTIC DEVICE ORALLY INTO THE ANIMAL...

U-373 GENERAL USE CLAIM SUBMITTED FOR 12 NEXIUM PATIENTS STATING "PERTINENT TO THE CAPSULE FORMULATION FOR NEXIUM AND ITS INDICATIONS FOR THE TREATMENT OF GERD AND ERADICATION OF H.PYLORI TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

U-374 KIT ADAPTED AND DESIGNED TO PROVIDE BOTH DATA ON THE CURRENT REPRODUCTIVE STATUS OF A PATIENT AND CONTRACEPTION FOR THOSE WHO ARE NOT PREGNANT, BUT RECENTLY ENGAGED IN UNPROTECTED SEX

U-375 METHOD OF USING RIBAVIRIN FOR TREATING A DISEASE RESPONSIVE TO RIBAVIRIN, E.G. HEPATITIS C

U-376 TREATMENT OF INFLUENZA

U-377 METHOD OF TREATING PT WITH CHRONIC HEPATITIS C HAVING HCV GENOTYPE 1 AND VIRAL LOAD GREATER THAN 2 MILLION COPIES/ML TO ERADICATE DETECTABLE HCV-RNA BY ADMIN COMBINATION OF RIBAVIRIN AND INTERFERON ALFA-2B FOR A LEAST 24 WEEKS

U-378 METHOD FOR TREATING INCONTINENCE

U-379 METHOD OF TREATING ONYCHOMYCOSIS

U-380 COMBINATIONS OF TAXOL (PACLITAXEL) AND CISPLATIN WHICH ARE SUITABLE FOR THE TREATMENT OF OVARIAN AND NON-SMALL CELL LUNG CARCINOMAS

U-381 TREATMENT OF HYPERPHOSPHATEMIA

U-382 METHOD OF STABILIZING PROSTAGLANDIN

U-383 METHOD FOR TREATING GLAUCOMA AND OCULAR HYPERTENSION

U-384 TREATMENT OF CMV RETINITIS

U-385 TREATMENT OF PEPTIC ULCERS

U-386 TREATMENT OF PATIENTS SUFFERING FROM A LATE ASTHMATIC REACTION OR LATE PHASE ASTHMA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-387 TREATMENT OF PATIENTS WITH RESPIRATORY DISORDERS

U-388 SMOKING CESSATION AID APPLIED TO THE SKIN

U-389 SMOKING CESSATION AID APPLIED TO THE SKIN ON WAKING AND REMOVED PRIOR TO SLEEP AFTER ABOUT 16 HOURS

U-390 METHOD OF USING THE DRUG TO TREAT NEUROIMMUNOLOGIC DISEASES (INCLUDING MULTIPLE SCLEROSIS)

U-391 USE OF CASODEX IN COMBINATION WITH LHRH AGONISTS FOR THE TREATMENT OF PROSTATE CANCER

U-392 TREATMENT OF PATIENTS FOR INFLAMMATION

U-393 MANAGEMENT OF INCONTINENCE, MGT OF HORMONE REPLACEMENT THERAPY, TREATMENT OF INVOLUNTARY INCONTINENCE, MGT OVERACTIVE BLADDER AND INCREASING COMPLIANCE IN SUCH PT

U-394 METHOD OF USE OF ALPHAGAN

U-395 METHOD OF USE OF ALPHAGAN P

U-396 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION

U-397 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION WITHOUT AN INCREASE IN NAUSEA

U-398 TREATMENT OF GENERALIZED ANXIETY DISORDER

U-399 IN-THE-EYE USE OF CHLORINE DIOXIDE CONTAINING COMPOSITIONS

U-400 USE OF RIBAVIRIN TO INCREASE TYPE 1 CYTOKINE RESPONSE AND SUPPRESS TYPE 2 CYTOKINE RESPONSE TO LYMPHOCYTES, INCLUDING METHODS THAT TAKE ADVANTAGE OF SUCH MODULATION TO TREAT INFECTIONS AND INFESTATIONS

U-401 USE OF LOPINAVIR IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS FOR TREATING HIV INFECTION AND IN COMBO WITH OTHER HIV PROTEASE INHIBITORS

U-402 TREATMENT OF ACTINIC KERATOSES

U-403 ANTI-ALLERGIC FOR VARIOUS ALLERGIC DISEASES

U-404 TREATMENT OF ALLERGIC CONJUNCTIVITIS

U-405 FOR WOMEN WITH SEVERE DIARRHEA-PREDOMINANT IRRITABLE BOWEL SYNDROME (IBS)

U-406 METHOD OF USE OF ATOVAQUONE AND PROGUANIL

U-407 METHOD OF TREATING OTOPATHY

U-408 FOR INDUCING OVULATION IN CONJUNCTION WITH A GONADOTROPIN RELEASING FACTOR ANTAGONIST AND RECRUITING OOCYTES FOR IN-VITRO FERTILIZATION

U-409 METHOD OF TREATING INFLAMMATION USING DRUG SUBSTANCE

U-410 METHOD OF REDUCING AMOUNT OF RESPECTIVE ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (INCLUDING PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-411 METHOD OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN PREPARATION

U-412 TREATMENT OF TYPE 2 DIABETES

U-413 USE OF THE ACTIVE INGREDIENT FOR INHIBITING THE BIOSYNTHESIS OF CHOLESTEROL AND TREATMENT OF ATHEROSCLEROSIS

U-414 A METHOD OF TREATING GLYCOMETABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE

U-415 A METHOD FOR REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS

U-416 A METHOD FOR REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS

U-417 COMBINATION USE OF AD-4833 WITH A BIGUANIDE

U-418 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

U-419 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE

U-420 METHOD OF TREATMENT OF TYPE II DIABETES

U-421 USE FOR SEDATION

U-422 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER AND ATTENTION DEFICIT HYPERACTIVITY DISORDER

U-423 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER, ATTENTION DEFICIT HYPERACTIVITY DISORDER, OR AIDS RELATED DEMENTIA

U-424 FOR ONCE DAILY, BOLUS ADMINISTRATION TO A PATIENT IN ORDER TO ENGENDER TREATMENT FOR A NERVOUS DISORDER FOR SUBSTANTIALLY AN ENTIRE DAY ON A CHRONIC BASIS

U-425 METHOD OF REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMIN TO A DIABETIC BY ADMIN A CHEMICAL COMPOUND HAVING FORMULA (INCL PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-426 PREVENTION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION

U-427 METHOD OF TREATING ALLERGIC REACTIONS IN MAMMALS

U-428 METHOD OF TREATING ALLERGY IN A MAMMAL USING THIS ACTIVE METABOLITE

U-429 METHOD OF USING DESLORATADINE TO TREAT ALLERGIC RHINITIS

U-430 METHOD OF TREATING A DIABETIC BY ADMINISTERING AN INSULIN SENSITIZER IN COMBINATION WITH AN INSULIN SECRETION ENHANCER, AND A DRUG PRODUCT COMPRISING AN INSULIN SENSITIZER AND AN INSULIN SECRETION ENHANCER

U-431 POSTTRAUMATIC STRESS DISORDER

U-432 REDUCTION OF ATHEROSCLEROTIC EVENTS (MYOCARDIAL INFARCTION, STROKE, AND VASCULAR DEATH) IN PATIENTS WITH ATHEROSCLEROSIS DOCUMENTED BY RECENT STROKE, RECENT MYOCARDIAL INFARCTION OR ESTABLISHED PERIPHERAL ARTERIAL DISEASE

U-433 USE OF LEVOCARNITINE IN PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS

U-434 CONTROLLED SYMPTOMS OF DIARRHEA, BLOATING PRESSURE AND CRAMPS, COMMONLY REFERRED TO AS GAS

U-435 A TITRATION DOSING REGIMEN FOR THE TREATMENT OF PAIN USING AN INITIAL DOSE OF ABOUT 25MG

U-436 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS

U-437 METHOD OF USE EQUAL TO PROCESS OF PREPARATION

U-438 TREATMENT/PREVENTION OF NEURODEGENERATIVE DISEASE

U-439 TREATMENT OF OBESITY

U-440 METHOD FOR TRANSDERMAL ADMINISTRATION OF A DRUG THROUGH NON-SCROTAL SKIN USING A TRANSDERMAL DRUG DELIVERY DEVICE CONTAINING THE DRUG AND HAVING AN ADHESIVE SURFACE

U-441 METHOD OF TREATING MS BY ADMINISTERING COPAXONE

U-442 METHOD FOR DELIVERING A DRUG TO A PATIENT IN NEED OF THE DRUG, WHILE AVOIDING THE OCCURRENCE OF AN ADVERSE SIDE EFFECT KNOWN OR SUSPECTED OF BEING CAUSED BY SAID DRUG

U-443 MANAGEMENT OF MODERATE TO SEVERE PAIN WHEN A CONTINUOUS, AROUND-THE-CLOCK ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME

U-444 TREATMENT OF MIGRAINE

U-445 USE AS AN ANTIMYCOTIC AGENT

U-446 TOPICAL TREATMENT OF OCULAR HYPERTENSION AND GLAUCOMA

U-447 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-448 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-449 USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER WHERE THE DOSE OF LEUCOVORIN IS AT LEAST 200MG PER SQUARE METER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-450 INTERMEDIATE REL NICOTINIC ACID FORMULATIONS HAVING UNIQUE URINARY METAB PROFILES RESULTING FROM ABSORPTION PROFILES OF NICOTINIC ACID FROM THE INTERMEDIATE NICOTINIC ACID FORMULATIONS, SUITABLE FOR TX HYPERLIPIDEMIA FOLLOWING QD DOSING

U-451 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER

U-452 USE OF LANSOPRAZOLE FOR COMBATING DISEASES CAUSED BY THE GENUS CAMPYLOBACTER (C.PYLORI=H.PYLORI)

U-453 TREATMENT OF PLATELET ASSOCIATED ISCHEMIC DISORDERS

U-454 METHOD OF TX A PT SUSPECTED OF HAVING HEPATITIS C BY ADMIN, IN COMBINATION, A CONJUGATE COMPRISING PEG 12000 & INTERFERON ALFA-2B IN AN AMT OF FROM 0.5MCG/KG TO 2MCG/KG, ONCE WEEKLY, AND RIBAVIRIN

U-455 TREATMENT OF PULMONARY HYPERTENSION WITH UT-15

U-456 METHOD OF DECREASING THE PRODUCTION OF A-BETA USING A COMPOSITION WHICH DECREASES BLOOD CHOLESTEROL IN PATIENTS AT RISK OF OR EXHIBITING SYMPTOMS OF ALZHEIMER'S DISEASE

U-457 METHOD OF TREATING A VAGINAL FUNGAL INFECTION IN A FEMALE HUMAN

U-458 METHOD OF USE OF IMAGENT

U-459 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER

U-460 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING SERTRALINE

U-461 METHOD OF TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER (PMDD) USING SERTRALINE

U-462 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND ADULT RHEUMATOID ARTHRITIS AND TREATMENT OF PRIMARY DYSMENORRHEA

U-463 VENOGRAPHY

U-464 PERIPHERAL ARTERIOGRAPHY

U-465 CT IMAGING OF THE HEAD

U-466 TREATMENT OF IRRITABLE BOWEL SYNDROME

U-467 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR FOR TREATING HYPERTENSION

U-468 METHOD OF USING FEXOFENADINE HCL IN TREATING ALLERGIC RHINITIS

U-469 TREATMENT OF GASTROESOPHAGEAL REFLEX DISEASE (GERD) AND ERADICATION OF H.PYLORI TO REDUCE RISK OF DUODENAL ULCER RECURRENCE

U-470 THERAPY IN CHRONIC HEPATITIS B VIRUS INFECTION

U-471 METHOD OF TREATING A PATIENT SUFFERING FROM DIABETES MELLITUS

U-472 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING METHYLPHENIDATE BI-MODAL RELEASE PROFILE EXTENDED-RELEASE CAPSULES

U-473 TO REDUCE PLASMA CHOLESTEROL LEVELS IN A MAMMAL

U-474 TO REDUCE PLASMA CHOLESTEROL LEVELS BY ADMIN EZETIMIBE IN COMBO WITH CHOLESTEROL BIOSYNTHESIS INHIB SELECTED FROM GROUP CONSISTING OF HMG COA REDUCTASE INHIBITORS INCL SIMVASTATIN

U-475 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY

U-476 METHOD OF TREATING ANDROGEN RESPONSIVE/MEDIATED CONDITION IN MAMMAL BY ADMIN A SAFE, EFFECTIVE AMOUNT OF DUTASTERIDE OR PHARMACEUTICALLY ACCEPTABLE DERIVATIVE THEREOF..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY

U-477 METHOD OF INHIBITING 5 ALPHA TESTOSTERONE REDUCTASE ENZYME WITH DUTASTERIDE OR ITS DERIVATIVE AND TREATING ANDROGEN RESPONSIVE/MEDIATED DISEASE INCLUDING BENIGN PROSTATIC HYPERPLASIA

U-478 METHOD OF TREATING HEPATITIS C VIRAL INFECTION BY CONTINUOUS PARENTERAL ADMIN INTERFERON ALPHA 2-10 MILLION IU WEEKLY, SUBCUTANEOUSLY, INJECTION OF POLYMER-INTERFERON ALPHA CONJUGATE-POLYMER IS PEG-INTERFERON IS ALPHA 2B

U-479 METHOD OF USING PEG-INTRON/REBETOL COMBINATION THERAPY AND INTRON/REBETOL COMBINATION THERAPY

U-480 CONTRAST AGENT FOR MRI

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-481 DISUBSTITUTED ACETYLENES BEARING HETEROAROMATIC AND HETEROBICYCLIC GROUPS HAVING RETINOID-LIKE ACTIVITY
- U-482 METHOD OF IN VITRO FERTILIZATION THERAPY INCLUDING MEANS FOR INDUCING OVULATION....
- U-483 METHOD FOR THE ADMINISTRATION OF DRUGS USING THAT COMPOUND
- U-484 METHOD OF TREATING A SKIN DISEASE WITH A CORTICOSTEROID-CONTAINING PHARMACEUTICAL COMPOSITION
- U-485 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-486 EXTERNAL PREPARATION FOR APPLICATION TO THE SKIN CONTAINING LIDOCAINE-DRUG RETAINING LAYER PLACED ON SUPPORT AND COMPRISES ADHESIVE GEL BASE 1-10% BY WEIGHT OF LIDOCAINE
- U-487 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-488 METHOD FOR REDUCING THE PAIN ASSOCIATED WITH HERPES-ZOSTER AND POST-HERPETIC NEURALGIA
- U-489 EXPECTORANT
- U-490 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE
- U-491 METHOD OF DELIVERING A DRUG TO THE LUNG
- U-492 METHOD FOR THE TREATMENT OF SKIN, SUFFERING FROM A CONDITION SELECTED FROM A GROUP CONSISTING OF NONACNE INFLAMMATORY DERMATOSES... COMPRISING APPLYING TO AFFECTED AREA. A THERAPEUTICALLY EFFECTIVE AMT AZELAIC ACID
- U-493 TREATMENT OF TYPE 2 DIABETES MELLITUS
- U-494 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER
- U-495 PERITONEAL DIALYSIS SOLUTION
- U-496 METHOD FOR TREATING CHRONIC RENAL FAILURE
- U-497 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND RHEUMATOID ARTHRITIS
- U-498 INTRA-ARTERIAL AND INTRAVENOUS USES OF ULTRAVIST
- U-499 METHOD OF USING REBETOL CAPSULES IN COMBINATION WITH A CONJUGATE COMPRISING POLYETHYLENE GLYCOL (PEG) AND AN ALPHA INTERFERON, INCLUDING, FOR EXAMPLE, PEG-INTRON POWDER FOR INJECTION
- U-500 USE AS AN ANTIHYPERTENSIVE AGENT
- U-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN ADULTS
- U-502 PITYRIASIS VERSICOLOR
- U-503 GENERATOR MUST BE USED WITH INFUSION SYSTEM SPECIFICALLY LABELED FOR USE WITH GENERATOR
- U-504 TINEA PEDIS, TINEA CRURIS, TINEA CORPORIS
- U-505 ULTRASOUND CONTRAST AGENT
- U-506 PHARM PRODUCT CONTAINER 1ST CHAMBER IS DISPOSED AQUEOUS DILUENT SOL 2ND CHAMBER PHARM ACTIVE AGENT COMPRISING ACETYLCHOLINE, BUFFER IN 1ST CHAM IS SUFFICIENT TO BUFFER PH OF MIXED SOL RESULTING MIXTURE OF AQUEOUS DILUENT SOL & PHARM ACTIVE..
- U-507 ACROMEGALY IN PATIENTS W/INADEQUATE RESPONSE TO SURGERY AND/OR RADIATION THERAPY AND/OR MEDICAL THERAPIES, OR FOR WHOM THESE THERAPIES ARE NOT APPROPRIATE
- U-508 METHOD OF RELEASING 17-BETA OESTRADIOL PRECURSOR IN A SUBSTANTIALLY ZERO ORDER PATTERN FOR AT LEAST THREE WEEKS
- U-509 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-510 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (STAGE IA AND IB) WHO HAVE REFRACTORY OR PERSISTENT DISEASE AFTER OTHER THERAPIES OR WHO HAVE NOT TOLERATED OTHER THERAPIES
- U-511 USE OF QUINOLONE COMPOUNDS AGAINST ANAEROBIC PATHOGENIC BACTERIA
- U-512 USE OF QUINOLONE COMPOUNDS AGAINST ATYPICAL UPPER RESPIRATORY PATHOGENIC BACTERIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-513 METHODS OF USE OF ANTIMICROBIAL COMPOUNDS AGAINST PATHOGENIC AMYCOPLASMA BACTERIA

U-514 PREVENTION OF OVULATION IN A WOMAN

U-515 TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON THE LAST THERAPY

U-516 METHOD OF TREATING A PSYCHOTIC DISEASE

U-517 STABLE GEL FORMULATION FOR TOPICAL TREATMENT OF SKIN CONDITIONS

U-518 OBSESSIVE COMPULSIVE DISORDER

U-519 POST OPERATIVE NAUSEA AND VOMITING

U-520 PREMENOPAUSAL OSTEOPOROSIS

U-521 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTRON A (INTERFERON ALPHA-2 B RECOMBINANT) INJECTION TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-522 TREATMENT OF CMV RETINITIS BY INTRAVITREAL ADMIN OF A PHOSPHOROTHIOATE OLIGONUCLEOTIDE CAPABLE OF HYBRIDIZING WITH CMV MRNA

U-523 METHOD OF TREATING INFECTION BY CRYPTOSPORIDIUM PARVUM IN AN IMMUNOCOMPROMISED MAMMAL

U-524 METHOD OF TREATING DIARRHEA

U-525 METHOD OF TREATING PARASITIC INFECTIONS

U-526 METHOD OF PROVIDING CONTROLLED RELEASE OF A TREATING AGENT USING A CONTROLLED RELEASE COMPOSITION

U-527 METHOD OF DELIVERING AN ACTIVE INGREDIENT USING A PROGRESSIVE HYDRATION BIOADHESIVE

U-528 PREVENTION OF CHEMOTHERAPY-INDUCED NAUSEA AND VOMITING

U-529 ONCE DAILY TREATMENT OF ASTHMA WITH NEBULIZED BUDESONIDE

U-530 TREATMENT OF HERPES ZOSTER, TREATMENT OF GENITAL HERPES, TREATMENT OF COLD SORES, SUPPRESSION OF GENITAL HERPES IN IMMUNOCOMPETENT AND HIV-INFECTED INDIVIDUALS, REDUCTION OF RISK OF HETEROSEXUAL TRANSMISSION OF GENITAL HERPES

U-531 TREATMENT OF PATIENTS WITH ESSENTIAL HYPERTENSION. MAY BE USED ALONE OR GIVEN WITH OTHER CLASSES OF ANTIHYPERTENSIVES, ESPECIALLY THIAZIDE DERIVATIVES

U-532 TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD IN PATIENTS REQUIRING MORE THAN ONE BRONCHO DILATOR

U-533 ERECTILE DYSFUNCTION

U-534 HUMALOG IS AN INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS FOR THE CONTROL OF HYPERGLYCEMIA

U-535 TREATMENT OF SOCIAL ANXIETY DISORDER

U-536 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING

U-537 TREATMENT OF CONDITIONS RELATED TO HYPERALDOSTERONISM SUCH AS HYPERTENSION AND CARDIAC INSUFFICIENCY, WITH EPLERENONE

U-538 FIRST LINE TREATMENT OF SEVERE HYPERTENSION, IN PATIENTS WITH HYPERTENSION SEVERE ENOUGH THAT THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY IN THESE PATIENTS

U-539 TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE

U-540 TREATMENT OF FUNGAL INFECTIONS

U-541 METHOD OF TREATMENT OF ADULTS INFECTED WITH HIV-1

U-542 METHOD OF TREATING PATIENT WITH TYPE 2 DIABETES BY ONCE DAILY ADMINISTRATION

U-543 TREATMENT OF SCHIZOPHRENIA

U-544 TREATMENT OF OVERACTIVE BLADDER. TREATMENT OF URINARY INCONTINENCE.

U-545 METHOD FOR THE PREVENTION AND/OR TREATMENT OF THROMBOTIC EPISODES, SUCH AS MYOCARDIAL INFARCTION, IN A HUMAN PATIENT AND METHOD FOR THE PREVENTION OF VENOUS THROMBOSIS IN A POSTOPERATIVE HUMAN PATIENT

U-546 USE OF REPAGLINIDE IN COMBINATION WITH METFORMIN TO LOWER BLOOD GLUCOSE

U-547 MAINTENANCE MONOTHERAPY FOR BIPOLAR DISORDER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-548 A METHOD OF REDUCING FLUSH IN AN INDIVIDUAL BEING TREATED FOR A LIPIDEMIC DISORDER AND EFFECTIVELY TREATING THE LIPIDEMIC DISORDER
- U-549 USE IN THE TREATMENT OF MEN WITH ADVANCED SYMPTOMATIC PROSTATE CANCER
- U-550 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA
- U-551 METHOD FOR REDUCING TOXICITY OF ALIMTA TREATED PATIENTS BY ADMINISTERING FOLIC ACID
- U-552 TREATMENT OF HYPERTENSION AND HYPERLIPIDEMIA WITH A SINGLE COMPOSITION
- U-553 MANAGEMENT OF PAIN AND DISCOMFORT ASSOCIATED WITH PERIDONTAL SCALING AND ROOT PLANNING PROCEDURES BY APPLICATION OF AN EUTECTIC MIXTURE OF LOCAL ANESTHETICS TO PERIDONTAL POCKETS
- U-554 TREATING HIV INFECTION WITH INDINAVIR SULFATE IN COMBINATION WITH ANTIRETROVIRAL AGENTS
- U-555 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
- U-556 USE AS ADJUNCT DIAGNOSTIC FOR SERUM THYROGLOBULIN (TG) TESTING
- U-557 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-558 INDICATED FOR THE RELIEF OF BRONCHOSPASM IN PATIENTS 2-12 YEARS OF AGE WITH ASTHMA (REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE)
- U-559 METHOD OF DECREASING OR REDUCING PARATHYROID HORMONE LEVEL; METHOD OF MODULATING PARATHYROID HORMONE SECRETION;METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF REDUCING SERUM IONIZED CALCIUM LEVEL
- U-560 METHOD OF DECREASING PARATHYROID HORMONE LEVEL;METHOD OF TREATING HYPERPARATHYROIDISM
- U-561 COSOPT IS INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION WHO ARE INSUFFICIENTLY RESPONSIVE TO BETA BLOCKERS
- U-562 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
- U-563 MARINOL IS INDICATED FOR, INTER ALIA, ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS
- U-564 TREATMENT OF HIV IN CONCOMITANT THERAPY
- U-565 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS, AND CHRONIC URTICARIA
- U-566 FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-567 METHOD OF TREATING INFERTILITY
- U-568 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION
- U-569 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THEREAFTER AN OVULATORY INDUCING AMOUNT OF HCG IS ADMINISTERED
- U-570 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THE DAILY AMOUNT OF FSH IS ABOUT 5-10 IU/KG
- U-571 TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA AND BIPOLAR I MANIA
- U-572 INTENSIVE CARE UNIT SEDATION
- U-573 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-574 PROPHYLAXIS AND TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS AND TREATMENT OF THE NASAL SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-575 LOTEMAX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TREATMENT OF STEROID RESPONSIVE CONDITIONS OF THE PALPEBRAL BULBAR CONJUNCTIVA, CORNEA AND ANTERIOR SEGMENT OF THE GLOBE.
- U-576 ALREX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TEMPORARY RELIEF OF THE SIGNS AND SYMPTOMS OF SEASONAL ALLERGIC CONJUNCTIVITIS.
- U-577 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA WITH FINASTERIDE IN COMBINATION WITH DOXAZOSIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-578 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA, ACUTE EXACERBATION OF CHRONIC BRONCHITIS, AND ACUTE BACTERIAL SINUSITIS CAUSED BY SUSCEPTIBLE STRAINS OF DESIGNATED MICROORGANISMS IN PATIENTS 18 YEARS AND OLDER.

U-579 TREATMENT OF EPILEPSY AND/OR MIGRAINE.

U-580 TREATMENT OF DISORDERS OF THE SEROTONERGIC SYSTEM SUCH AS DEPRESSION AND ANXIETY-RELATED DISORDERS

U-581 METHOD OF TREATING A CONDITION CAPABLE OF TREATMENT BY INHALATION, E.G. ASTHMA, COMPRISING ADMINISTRATION OF A FORMULATION CLAIMED IN US PATENT NO. 6743413

U-582 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6253762

U-583 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING TO A PATIENT BY INHALATION, A METERED AEROSOL DOSE OF A DRUG FORMULATION FROM THE METERED DOSE INHALER SYSTEM CLAIMED IN US 6546928

U-584 SINGLE-DOSE ADMINISTRATION BY THE EPIDURAL ROUTE, AT THE LUMBAR LEVEL, FOR THE TREATMENT OF PAIN FOLLOWING MAJOR SURGERY

U-585 TO PROMOTE WEIGHT GAIN AFTER WEIGHT LOSS IN CERTAIN TYPES OF PATIENTS

U-586 AN INTERMEDIATE RELEASE NICOTINIC ACID FORMULATION SUITABLE FOR ORAL ADMINISTRATION ONCE-A-DAY AS A SINGLE DOSE FOR TREATING HYPERLIPIDEMIA WITHOUT CAUSING DRUG-INDUCED HEPATOTOXICITY OR ELEVATIONS IN URIC ACID OR GLUCOSE OR BOTH

U-587 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR (AND OPTIONALLY A DIURETIC) FOR TREATING CONGESTIVE HEART FAILURE AND HYPERTENSION

U-588 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER; TREATMENT OF HEARTBURN AND OTHER SYMPTOMS ASSOCIATED WITH GERD; SHORT-TERM TREATMENT OF EROSIIVE ESOPHAGITIS; MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS

U-589 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN U.S. PATENT NO. 6131966

U-590 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING TO A PATIENT BY ORAL OR NASAL INHALATION A DRUG FORMULATION BY USING THE METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6532955

U-591 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING A DOSAGE FORM WHICH PROVIDES ONCE-DAILY ORAL ADMINISTRATION OF A PHENIDATE DRUG

U-592 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)

U-593 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)

U-594 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS

U-595 35 MG ORALLY ONCE A WEEK FOR PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN; 35 MG ORALLY ONCE A WEEK FOR TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

U-596 TREATMENT OF HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY

U-597 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT HIGH RISK FOR FRACTURE

U-598 PROPHYLACTIC TREATMENT OF MIGRAINE

U-599 METHOD FOR TREATING ALLERGIC CONJUNCTIVITIS

U-600 A METHOD OF TREATING A PATIENT IN NEED OF OPHTHALMIC ANTIMICROBIAL THERAPY WITH LEVOFLOXACIN

U-601 TREATMENT OF BIPOLAR DISORDER

U-602 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS, RHEUMATOID ARTHRITIS IN ADULTS, AND/OR PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS, ACUTE PAIN IN ADULTS; PRIMARY DYSMENORRHEA; AND/OR ACUTE MIGRAINE ATTACKS IN ADULTS

U-603 METHOD OF TREATING INFECTIONS COMPRISING ORALLY ADMINISTERING AN EFFECTIVE AMOUNT OF THE FDA APPROVED ORAL SUSPENSION

U-604 METHOD OF LOWERING BLOOD GLUCOSE BY ONCE DAILY ADMINISTRATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-605 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD); ALTHOUGH THE MECHANISM OF THE ANTIDEPRESSANT ACTION OF DULOXETINE IN HUMANS IS UNKNOWN, IT IS BELIEVED TO BE RELATED TO ITS POTENTIATION OF SERATONERGIC AND NORADRENERGIC ACTIVITY IN THE CNS
- U-606 USE OF IRINOTECAN IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER
- U-607 CANCIDAS IS INDICATED FOR EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS.
- U-608 USE OF QUINOLONE COMPOUNDS AGAINST PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-609 USE OF QUINOLONE COMPOUNDS AGAINST QUINOLONE-RESISTANT PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-610 ATROVENT HFA (IPRATROPIUM BROMIDE HFA) INHALATION AEROSOL IS INDICATED AS A BRONCHODILATOR FOR MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA.
- U-611 METHOD OF USING DESLORATADINE TO TREAT SEASONAL AND PERENNIAL ALLERGIC RHINITIS, PRURITIS, AND CHRONIC IDIOPATHIC URTICARIA IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-612 TREATMENT OF SEASONAL ALLERGY SYMPTOMS WITH NASAL CONGESTION IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-613 REDUCTION OF SERUM PHOSPHATE
- U-614 TREATMENT OF SEXUAL DYSFUNCTION
- U-615 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TOTAL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)
- U-616 MANAGEMENT OF PERSISTENT, MODERATE TO SEVERE PAIN IN PATIENTS REQUIRING CONTINUOUS, AROUND-THE-CLOCK ANALGESIA WITH A HIGH POTENCY OPIOID FOR AN EXTENDED PERIOD OF TIME GENERALLY WEEKS TO MONTHS OR LONGER
- U-617 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-618 USE OF ROSUVASTATIN CALCIUM TO REDUCE ELEVATED TOTAL-C, LDL-C, APOB, NONHDL-C OR TG LEVELS; TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA; AND TO SLOW THE PROGRESSION OF ATHEROSCLEROSIS.
- U-619 TREATMENT OF MALIGNANT NEOPLASM
- U-620 TREATMENT OF INSOMNIA
- U-621 METHOD OF TREATING CANCER
- U-622 TREATMENT OF VEGF MEDIATED OCULAR DISEASE.
- U-623 SHORT TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- U-624 REDUCTION OF RISK OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS
- U-625 ALLERGIC RHINITIS OR NASAL POLYPS
- U-626 CLOLAR IS INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 TO 21 YEARS OLD WITH RELAPSED OR REFRACTORY ACUTE LYMPHOBLASTIC LEUKEMIA AFTER AT LEAST TWO PRIOR REGIMENS
- U-627 TREATMENT OF PATIENTS USING EXTENDED-RELEASE CARBAMAZEPINE
- U-628 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA, AND IN COMBINATION WITH METFORMIN AND A SULFONYLUREA TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-629 METHOD OF INDUCING A HYPNOTIC OR SEDATIVE EFFECT IN A HUMAN BY ADMINISTERING ESZOPICLONE
- U-630 TREATING URINARY INCONTINENCE BY ADMINISTERING AN EXTENDED-RELEASE FORM OF DARIFENACIN
- U-631 TREATING A DISEASE OF ALTERED MOTILITY OR TONE OF SMOOTH MUSCLE BY ADMINISTERING A MUSCARINIC RECEPTOR ANTAGONIZING AMOUNT OF DARIFENACIN
- U-632 METHOD OF TREATMENT OF CANCER BY ADMINISTERING PARTICLES OF PACLITAXEL THAT HAVE A PROTEIN COATING
- U-633 METHOD FOR TREATMENT OF TUMORS BY ADMINISTERING PACLITAXEL AT A DOSE IN THE RANGE OF ABOUT 30MG/METER SQUARE TO ABOUT 100MG/METER SQUARE IN A PHARMACEUTICALLY ACCEPTABLE FORMULATION THAT DOES NOT CONTAIN CREMOPHOR

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-634 METHOD FOR DELIVERY OF A BIOLOGIC (INCLUDING ANTINEOPLASTIC AGENTS) BY ADMINISTERING TO A PATIENT AN EFFECTIVE AMOUNT OF A BIOLOGIC AS A SOLID OR LIQUID WITH A POLYMERIC BIOCOMPATIBLE MATERIAL

U-635 TREATMENT OF GERD, MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND RISK REDUCTION OF NSAID ASSOCIATED GASTRIC ULCERS

U-636 TREATMENT OR PREVENTION OF BRONCHOSPASM OR ASTHMATIC SYMPTOMS

U-637 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST

U-638 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST, INCLUDING WITH INSULIN

U-639 TREATMENT OF A MAMMAL HAVING A NEED OF OR REDUCED ABILITY TO PRODUCE INSULIN WITH AN INSULIN AND AN AMYLIN SUCH AS PRAMLINTIDE

U-640 USE OF AN AMYLIN AGONIST TO REDUCE GASTRIC MOTILITY AND TREAT POST PRANDIAL HYPERGLYCEMIA

U-641 USE OF AN AMYLIN AGONIST HAVING SPECIFIED BINDING ACTIVITY TO REDUCE GASTRIC MOTILITY, INCLUDING USE THROUGH PARENTERAL ADMINISTRATION

U-642 TREATMENT AND PREVENTION OF OSTEOPOROSIS

U-643 THE SHORT TERM TREATMENT (UP TO 10 DAYS) IN PTS HAVING GASTROESOPHAGEAL REFLUX DISEASE (GERD) AS AN ALTERNATIVE TO ORAL THERAPY IN PTS WHEN THERAPY WITH NEXIUM CAPSULES IS NOT POSSIBLE OR APPROPRIATE

U-644 TREATMENT OF SEASONAL ALLERGIC RHINITIS

U-645 TREATMENT OF ASTHMA

U-646 METHOD OF TREATING OTITIS

U-647 TREATMENT OF OSTEOPOROSIS IN POST MENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

U-648 THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN

U-649 A METHOD FOR TREATING A TUMOR DISEASE

U-650 TREATMENT OF ESOPHAGEAL CANDIDIASIS AND PROPHYLAXIS OF CANDIDA INFECTIONS IN HSCT PATIENTS

U-651 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL)

U-652 TREATMENT OF CARDIAC ARRHYTHMIA

U-653 STIMULATING INSULIN RELEASE BY ADMINISTERING EXENATIDE

U-654 LOWERING PLASMA GLUCAGON IN A SUBJECT IN NEED THEREOF, INCLUDING ONE WITH TYPE 2 DIABETES, BY ADMINISTERING AN EXENDIN OR ANALOG, SUCH AS EXENDIN-4

U-655 TREATMENT OF MILD TO MODERATE ACTIVE CHROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON AND THE MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR ASCENDING COLON FOR UP TO 3 MONTHS

U-656 REDUCING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4

U-657 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

U-658 TREATMENT OF ADVANCED HORMONE-DEPENDENT BREAST CANCER

U-659 TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) AFTER FAILURE OF AT LEAST ONE PRIOR CHEMOTHERAPY REGIMEN

U-660 TREATMENT OF HYPERTENSION AND TREATMENT OF HEART FAILURE

U-661 TREATMENT OF SEIZURE DISORDER

U-662 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

U-663 THE TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS

U-664 TREATMENT OF CONDITIONS FOR WHICH AN ALDOSTERONE RECEPTOR BLOCKER IS INDICATED, SUCH AS HYPERTENSION, HEART FAILURE, AND POST-MYOCARDIAL INFARCTION

U-665 METHOD OF USING THE DRUG SUBSTANCE/DRUG PRODUCT FOR ULTRASOUND IMAGING

U-666 METHOD OF TREATING ADHD

U-667 MANAGEMENT OF INCONTINENCE; METHOD FOR TREATING INCONTINENCE

U-668 LEVEMIR IS A LONG-ACTING BASAL INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- OF PATIENTS WITH DIABETES MELLITUS
- U-669 INDICATION OF TYPE II DIABETES
- U-670 TREATMENT OF HIV-1 INFECTION BY THE CO-ADMINISTRATION OF TIPRANAVIR AND RITONAVIR.
- U-671 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 3 AND 4
- U-672 TREATMENT OF INFLAMMATION OR AN INFLAMMATION-ASSOCIATED DISORDER
- U-673 METHOD OF TREATMENT WITH ONCE-DAILY DOSES OF 625MG/5ML
- U-674 METHOD OF TREATING INSOMNIA CHARACTERIZED BY DIFFICULTY WITH SLEEP ONSET
- U-675 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA; RELIEF OF SYMPTOMS OF ALLERGIC RHINITIS
- U-676 METHOD OF TREATING ATTENTION DEFICIT DISORDER USING ORAL ADMINISTRATION OF A BI-MODAL OR PULSATILE RELEASE COMPOSITION
- U-677 A METHOD OF TREATING DISEASE AMENABLE TO TREATMENT WITH A PHENIDATE DRUG BY ONCE DAILY ORAL ADMINISTRATION OF AN EXTENDED RELEASE DOSAGE FORM
- U-678 METHOD OF TREATING ATTENTION DEFICIT DISORDER AND/OR ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-679 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN
- U-680 A METHOD OF TREATING DYSLIPIDEMIA AND DYSLIPOPTEINEMIA USING A DOSAGE FORM THAT CAN PROVIDE AN EFFECTIVE AMOUNT OF FENOFIBRATE TO A PATIENT IN A FASTED STATE WHICH IS AT LEAST 90% OF THE AUC AMOUNT PROVIDED BY THE DOSAGE FORM
- U-681 TREATMENT OF PRIMARY IGF-1 DEFICIENCY
- U-682 NON-BENZODIAZEPINE HYPNOTIC AGENT INDICATED FOR TREATMENT OF INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE
- U-683 PREVENTION OR TREATMENT OF ISCHEMIC HEART DISEASE
- U-684 TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-685 EXPECTORANT AND COUGH SUPPRESSANT
- U-686 EXPECTORANT AND NASAL DECONGESTANT
- U-687 REDUCING FOOD INTAKE IN A SUBJECT WITH TYPE 2 DIABETES BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4
- U-688 TREATMENT OF HIV-INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS
- U-689 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-690 TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-691 USE AS A MONOTHERAPY, IN COMBINATION WITH A SULFONYLUREA, METFORMIN OR INSULIN OR IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-692 USE OF VALSARTAN TO REDUCE CARDIOVASCULAR MORTALITY IN CLINICALLY STABLE PATIENTS WITH LEFT VENTRICULAR FAILURE OR LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- U-693 THE RECOMMENDED INITIAL DOSE OF EQUETRO IS 400MG/DAY GIVEN IN DIVIDED DOSES, TWICE DAILY. THE DOSE SHOULD BE ADJUSTED IN 200MG DAILY INCREMENTS TO ACHIEVE OPTIMAL CLINICAL RESPONSE.
- U-694 LENALIDOMIDE IS AN ANALOGUE OF THALIDOMIDE. THALIDOMIDE IS A KNOWN HUMAN TERATOGEN THAT CAUSES SEVERE LIFE-THREATENING HUMAN BIRTH DEFECTS. IF LENALIDOMIDE IS TAKEN DURING PREGNANCY, IT MAY CAUSE BIRTH DEFECTS OR DEATH TO AN UNBORN BABY.
- U-695 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA AND T-CELL LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-696 TREATMENT OF PATIENTS WITH T-CELL LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-697 A METHOD OF USING RINFABATE RECOMBINANT (RHIGFBP-3) WITH MECASERMIN RECOMBINANT (RHIGF-1) TO PROMOTE LINEAR GROWTH IN THE TREATMENT OF PRIMARY IGF-1 DEFICIENCY

U-698 METHOD OF USING ANTAGONIST OF ARGININE VASOPRESSIN (AVA) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH EUVOLEMIC HYPONATREMIA

U-699 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-700 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

U-701 TREATMENT OF HYPERCHOLESTEROLEMIA AND/OR HYPERTRIGLYCERIDEMIA

U-702 TOPICAL AEROSOL HAIR REGROWTH TREATMENT

U-703 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMOR AND RENAL CELL CARCINOMA WITH SUNITINIB

U-704 METHOD OF ADMINISTERING INSULIN VIA INHALATION

U-705 TREATING CHRONIC ANGINA BY ADMINISTERING AN EXTENDED RELEASE FORM OF RANOLAZINE

U-706 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA

U-707 ALLERGIC RHINITIS

U-708 TREATMENT OF CHRONIC NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE

U-709 METHOD OF COMBATING BACTERIA IN A PATIENT

U-710 A METHOD OF TREATING RESPIRATORY DISORDERS, E.G., ASTHMA, WHICH COMPRISES ADMINISTRATION BY INHALATION OF AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT NO. 5658549

U-711 ACUTE AND LONGER-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER

U-712 A METHOD OF USING A NICOTINIC ACID FORMULATION TO REDUCE ELEVATED TC, LDL-C AND TG LEVELS, AND RAISE HDL-C LEVELS IN PATIENTS WITH HYPERLIPIDEMIA

U-713 TREATMENT OF MILD TO MODERATE DEMENTIA OF THE ALZHEIMER'S TYPE

U-714 TOPICAL TREATMENT OF INTERDIGITAL TINEA PEDIS AND TINEA CORPORIS DUE TO TRICHOPHYTON RUBRUM, TRICHOPHYTON MENTAGROPHYTES OR EPIDERMOPHYTON FLOCCOSUM

U-715 FOR CLEANSING THE BOWEL IN PREPARATION FOR COLONOSCOPY, IN ADULTS 18 YEARS OF AGE OR OLDER

U-716 THE TREATMENT OR PREVENTION OF BRONCHOSPASM IN ADULTS AND CHILDREN 4 YEARS OF AGE AND OLDER WITH REVERSIBLE OBSTRUCTIVE AIRWAYS DISEASE AND THE PREVENTION OF EXERCISED-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER

U-717 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT

U-718 TREATMENT OF FUNGAL INFECTIONS

U-719 TREATMENT OF PSYCHOSIS

U-720 TREATMENT OF NEUROLEPTIC DISEASES

U-721 TREATMENT OF INFLUENZA

U-722 PROPHYLAXIS OF INFLUENZA

U-723 PROPHYLACTIC TREATMENT OF MIGRAINE

U-724 METHOD OF TREATING SEIZURES

U-725 ALLERGIC RHINITIS AND URTICARIA

U-726 ALLERGIC RHINITIS

U-727 FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)

U-728 METHOD FOR TREATING BACTERIAL INFECTION

U-729 TREATMENT OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER, H. PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

U-730 USE AS A NASAL SPRAY FOR TREATMENT OF THE SYMPTOMS OF SEASONAL ALLERGIC RHINITIS AND VASOMOTOR RHINITIS

U-731 USE IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

U-732 ACUTE TREATMENT OF THE CUTANEOUS MANIFESTATIONS OF MODERATE TO SEVERE ERYTHEMA NODOSUM LEPROSUM (ENL)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-733 MAINTENANCE THERAPY FOR PREVENTION AND SUPPRESSION OF THE CUTANEOUS MANIFESTATIONS OF ENL RECURRENCE

U-734 FIRST LINE THERAPY FOR TYPE 2 DIABETES MELLITUS

U-735 METHOD OF TREATING CHRONIC IRON OVERLOAD

U-736 METHOD FOR IONTOPHORETIC TRANSDERMAL DELIVERY OF FENTANYL HYDROCHLORIDE

U-737 DISINFECTION OF PATIENT SKIN PRIOR TO AN INVASIVE PROCEDURE

U-738 INDICATED FOR THE LONG-TERM, TWICE-DAILY MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE OR OLDER

U-739 METHOD FOR TREATING CONSTIPATION BY OPENING CIC CHANNELS IN A MAMMALIAN SUBJECT

U-740 FOR THE TREATMENT OF PATIENTS WITH PRIMARY BILIARY CIRRHOSIS

U-741 COMBINATION THERAPY WITH CISPLATIN FOR THE TREATMENT OF LATE STAGE CERVICAL CANCER

U-742 TWICE DAILY TOPICAL TREATMENT OF MODERATE TO SEVERE PLAQUE PSORIASIS.

U-743 ONCE A DAY TOPICAL TREATMENT OF THE INFLAMMATORY LESIONS OF ROSACEA

U-744 TREATMENT OF HIV INFECTION IN ANTIRETROVIRAL TREATMENT-EXPERIENCED ADULT PATIENTS

U-745 TREATMENT OR PREVENTION OF EMESIS

U-746 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT

U-747 PREVENTION OR TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING

U-748 A METHOD FOR THE TREATMENT OF A PROTEIN TYROSINE KINASE-ASSOCIATED DISORDER

U-749 METHOD OF CONTRACEPTION

U-750 TREATMENT OF HIV-1 INFECTION IN ADULTS

U-751 ONCE DAILY DOSING OF BUDESONIDE VIA NEBULIZER FOR THE TREATMENT OF ASTHMA

U-752 SUNSCREEN

U-753 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES

U-754 USE FOR THE LONG-TERM MAINTENANCE TREATMENT OF ASTHMA

U-755 TREATMENT OF ANOREXIA, CACHEXIA, OR AN UNEXPLAINED, SIGNIFICANT WEIGHT LOSS IN PATIENTS WITH A DIAGNOSIS OF ACQUIRED IMMUNODEFICIENCY SYNDROME (AIDS)

U-756 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

U-757 USE AS A BILE ACID SEQUESTRANT FOR LOWERING CHOLESTEROL

U-758 TREATMENT OF SYMPTOMS OF PREMENSTRUAL DYSPHORIC DISORDER

U-759 METHOD OF USE OF ADMINISTERING LEVOTHYROXINE

U-760 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS AND TREATMENT OF OROPHARYNGEAL CANDIDIASIS

U-761 TREATMENT OF SCHIZOPHRENIA INCLUDING MAINTAINING STABILITY IN PATIENTS WITH SCHIZOPHRENIA

U-762 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

U-763 ADMINISTRATION OF ARIPIRAZOLE BY INJECTION

U-764 TREATMENT OF SCHIZOPHRENIA

U-765 METHOD OF TREATING ALLERGIC CONJUNCTIVITIS

U-766 TREATMENT OF SEIZURES

U-767 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER

U-768 A METHOD OF REDUCING THE CAPACITY OF EXTENDED RELEASE NICOTINIC ACID TO PROVOKE A FLUSHING REACTION BY PRETREATING AN INDIVIDUAL WITH A FLUSH INHIBITING AGENT PRIOR TO THE ADMINISTRATION OF THE EXTENDED RELEASE NICOTINIC ACID

U-769 REVLIMID (LENALIDOMIDE) IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-770 LONG-TERM TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS

U-771 METHOD FOR THE TREATMENT OF DIABETES MELLITUS, SUCH AS TYPE 1 DIABETES MELLITUS OR TYPE 2 DIABETES MELLITUS, IN A HUMAN PATIENT

U-772 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN CHILDREN 2 TO 11 YEARS AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 MONTHS TO 11 YEARS

U-773 PATHOLOGICAL HYPERSECRETORY CONDITIONS

U-774 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR

U-775 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND/OR A SULFONYLUREA

U-776 TREATMENT OF CUTANEOUS MANIFESTATION IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL) WHO HAVE PROGRESSIVE, PERSISTENT OR RECURRENT DISEASE ON OR FOLLOWING TWO SYSTEMIC THERAPIES.

U-777 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE

U-778 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-779 A METHOD FOR TREATMENT OF A CANCER, WHEREIN THE CANCER IS CHRONIC MYELOGENOUS LEUKEMIA

U-780 A METHOD FOR THE TREATMENT OF CANCER

U-781 FOR TREATMENT OF ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE NAIVE TO PHARMACOLOGIC THERAPY

U-782 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS WITH EVIDENCE OF VIRAL REPLICATION AND EITHER EVIDENCE OF PERSISTENT ELEVATIONS IN SERUM AMINOTRANSFERASES (ALT OR AST) OR HISTOLOGICALLY ACTIVE DISEASE

U-783 DESONATE GEL IS INDICATED FOR THE TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE AND OLDER

U-784 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)

U-785 USE AS REPLACEMENT SOLUTION, HEMOFILTRATION SOLUTION OR HEMODIAFILTRATION SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY

U-786 PRODUCT IS APPROVED FOR THE TOPICAL TREATMENT OF TINEA PEDIS

U-787 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND PEDIATRIC PATIENTS SIX YEARS OF AGE OR OLDER, INCLUDING PATIENTS REQUIRING ORAL CORTICOSTEROID THERAPY FOR ASTHMA

U-788 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING PAROXETINE

U-789 TREATMENT OF KNOWN OR SUSPECTED CYANIDE POISONING

U-790 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT RISK FOR FRACTURE. FORTEO CAN BE USED BY PEOPLE WHO HAVE HAD A FRACTURE RELATED TO OSTEOPOROSIS

U-791 GLEEVEC IS ALSO INDICATED FOR THE TREATMENT OF PATIENTS WITH KIT (CD117) POSITIVE UNRESECTABLE AND/OR METASTATIC MALIGNANT GASTROINTESTINAL STROMAL TUMORS (GIST)

U-792 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS

U-793 FOR THE LONG TERM TREATMENT, TWICE DAILY (MORNING AND EVENING) MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA

U-794 CLOSURE OF A CLINICALLY SIGNIFICANT PATENT DUCTUS ARTERIOSUS IN PREMATURE INFANTS WEIGHING BETWEEN 500 AND 1500G, WHO ARE NO MORE THAN 32 WEEKS GESTATIONAL AGE WHEN USUAL MEDICAL MANAGEMENT IS INEFFECTIVE

U-795 METHOD FOR INHIBITING NOREPINEPHRINE UPTAKE

U-796 METHOD OF TREATING DEPRESSION

U-797 METHOD OF TREATING ANXIETY

U-798 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN BY ONCE-MONTHLY ORAL ADMINISTRATION OF IBANDRONATE SODIUM MONOHYDRATE EQUIVALENT TO 150MG OF IBANDRONIC ACID

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-799 METHOD FOR INHIBITING SEROTONIN UPTAKE

U-800 TREATMENT OF PATIENTS WITH ADVANCED OR METASTATIC BREAST CANCER WHOSE TUMORS OVEREXPRESS HER2 AND WHO HAVE RECEIVED PRIOR THERAPY INCLUDING ANTHRACYCLINE, A TAXANE AND TRASTUZUMAB

U-801 METHOD OF TREATING CANCER

U-802 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR

U-803 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN

U-804 TREATMENT OF ACTINIC KERATOSES BY PHOTODYNAMIC THERAPY

U-805 TREATMENT OF IMPETIGO DUE TO STAPHYLOCOCCUS AUREUS OR STREPTOCOCCUS PYOGENES

U-806 INTRATHECAL TREATMENT OF LYMPHOMATOUS MENINGITIS

U-807 PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION

U-808 THE TREATMENT OF THE SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN PATIENTS 2 YEARS OF AGE AND OLDER

U-809 TREATMENT OF CHRONIC IDIOPATHIC URTICARIA

U-810 METHOD OF TREATMENT TO ALLEVIATE INFLAMMATION OF THE EYE

U-811 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS AND TREATMENT OF THE UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA

U-812 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS

U-813 MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-814 TREATMENT OF SCHIZOPHRENIA

U-815 TREATS COLD SORES/FEVER BLISTERS ON THE FACE OR LIPS. SHORTENS HEALING TIME AND DURATION OF SYMPTOMS: TINGLING, PAIN, BURNING AND/OR ITCHING

U-816 DEPRESSION, PANIC DISORDER, PREMENSTRUAL DISORDERS AND SOCIAL ANXIETY DISORDER

U-817 NASAL ADMINISTRATION OF CYANOCOBALAMIN

U-818 TOPICAL TREATMENT OF ACNE VULGARIS

U-819 MANAGEMENT OF FIBROMYALGIA

U-820 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE SLEEPINESS ASSOCIATED WITH NARCOLEPSY, OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME, AND SHIFT WORK SLEEP DISORDER

U-821 METHOD OF INHIBITING ENTHOTHELIN RECEPTORS BY ADMINISTERING AMBRISENTAN TO A PATIENT TO TREAT PULMONARY ARTERIAL HYPERTENSION.

U-822 USE IN LIPID MANAGEMENT

U-823 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS AND FOR THE TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 TO 11 YEARS OF AGE

U-824 METHOD OF TREATING PATIENTS INFECTED WITH CCR5-TROPIC HIV-1

U-825 USE FOR PREVENTION OF BREAST CANCER

U-826 RELIEF OF MODERATE TO SEVERE PAIN

U-827 USE FOR TREATMENT OF DIABETES, PARTICULARLY TYPE 2 DIABETES

U-828 PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION

U-829 TREATMENT OF EXTRAVASATION RESULTING FROM IV ANTHRACYCLINE CHEMOTHERAPY

U-830 TREATMENT OF RELAPSED SMALL CELL LUNG CANCER

U-831 METHOD OF ADMINISTERING LANREOTIDE ACETATE

U-832 ZINGO IS INDICATED FOR THE USE ON INTACT SKIN TO PROVIDE LOCAL ANALGESIA PRIOR TO VENIPUNCTURE OR INTRAVENOUS CANNULATION.

U-833 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.25% BY WEIGHT OF ROPIVACAINE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-834 INVIRASE IN COMBINATION WITH RITONAVIR AND OTHER ANTIRETROVIRAL AGENTS IS INDICATED FOR THE TREATMENT OF HIV INFECTION
- U-835 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF ATOPIC DERMATITIS IN PATIENTS ONE YEAR OF AGE OR OLDER
- U-836 A METHOD FOR THE TREATMENT OF LEUKEMIAS
- U-837 GASTROINTESTINAL LAVAGE INDICATED FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY IN ADULTS
- U-838 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.5% BY WEIGHT OF ROPIVACAINE
- U-839 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- U-840 TREATMENT FOR TYPE 2 DIABETES MELLITUS
- U-841 INDICATED FOR THE LONG-TERM, MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE AND OLDER
- U-842 INDICATED FOR THE TREATMENT OF ATTENTION-DEFICIT/HYPERACTIVITY DISORDER (ADHD)
- U-843 METHOD FOR ADMINISTRATION OF TESTOSTERONE
- U-844 PREFEST IS INDICATED IN WOMEN WHO HAVE A UTERUS FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE; TREATMENT OF VULVAR AND VAGINAL ATROPHY; PREVENTION OF OSTEOPOROSIS
- U-845 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESSSES
- U-846 USE FOR DELINEATION (VISUALIZATION) DURING A VITRECTOMY SURGICAL PROCEDURE
- U-847 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)
- U-848 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-849 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP. DOSE IS ONE DROP OF COMBIGAN IN THE AFFECTED EYE TWICE DAILY
- U-850 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT
- U-851 TREATMENT OF TYPE 2 DIABETES MELLITUS
- U-852 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- U-853 TREATMENT OR PREVENTION OF EMESIS
- U-854 PREVENTION OF CMV DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)
- U-855 METHOD TO INDUCE NATRIURESIS, DIURESIS AND/OR VASODILATION
- U-856 SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN
- U-857 INHIBITION OF TRANSPLANT REJECTION
- U-858 PEDIATRIC USE AGED 1-11 YEARS, GERD AND EROSIVE ESOPHAGITIS
- U-859 EROSIVE ESOPHAGITIS, HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME, MAINTENANCE OF HEALING OF EROSIVE ESOPHAGITIS AND REDUCTION OF SYMPTOMS IN PATIENTS WITH GERD
- U-860 FOR THE APPROVED USES AND CONDITIONS OF USE, INCLUDING DEPRESSION
- U-861 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER
- U-862 ADJUNCT TO DIET TO REDUCE ELEVATED TOTAL-C, LDL-C, NON-HDL-C, APO B, TG, AND LP(A) LEVELS AND TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA, MIXED DYSLIPIDEMIA, AND HYPERTRIGLYCERIDEMIA
- U-863 TAKING ASPIRIN OR NON-STEROIDAL ANTI-INFLAMMATORY MEDICATIONS APPROXIMATELY 30 MINUTES BEFORE DOSING CAN MINIMIZE FLUSHING, A COMMON SIDE EFFECT OF NIACIN THERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-864 PEDIATRIC USE AGES 1-2 YEARS, GERD AND EROSIIVE ESOPHAGITIS

U-865 TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND A HIGH RISK FOR BONE FRACTURE BY REDUCING THE RISK OF VERTEBRAL AND NONVERTEBRAL BONE FRACTURE

U-866 THE LABEL REFERENCES THE EFFECTS OF THE ACTIVE INGREDIENT OF REVLIMID UPON CYTOKINES

U-867 TREATMENT OF MIGRAINE

U-868 METHOD OF USING ANTAGONIST OF ARGININE VASOPRESSIN (AVA) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH HYPERVOLEMIC HYPONATREMIA

U-869 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART

U-870 METHOD OF PRODUCING CORONARY VASODILATION WITHOUT PERIPHERAL VASODILATION

U-871 METHOD OF REDUCING RISK OF MYOCARDIAL INFARCTION, STROKE AND DEATH

U-872 TWICE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA. TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS

U-873 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME BY OPENING CHLORIDE CHANNELS (CIC)

U-874 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME

U-875 FIRST-LINE TREATMENT OF LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER, IN COMBINATION WITH GEMCITABINE

U-876 TREATMENT OF MIGRAINE WITH OR WITHOUT AURA

U-877 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PEPTIC ULCER

U-878 A METHOD FOR BINDING A PERIPHERAL OPIOID RECEPTOR

U-879 A METHOD OF TREATING OR PREVENTING ILEUS

U-880 ENDOMETRIN IS A PROGESTERONE INDICATED TO SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN

U-881 TREATMENT OF NON-SMALL CELL LUNG CANCER

U-882 MANAGEMENT OF FIBROMYALGIA (FM)

U-883 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR WITH SUNITINIB

U-884 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA

U-885 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY

U-886 ADMINISTERING DESLORATADINE TO TREAT THE SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS, SEASONAL ALLERGIC RHINITIS, OR CHRONIC IDIOPATHIC URTICARIA

U-887 TREATMENT AND PREVENTION OF OSTEOPOROSIS

U-888 FEMALE HORMONE REPLACEMENT THERAPY FOR POSTMENOPAUSAL WOMEN

U-889 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE)

U-890 REDUCTION OF SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE

U-891 USE AS AN INTRAOCULAR IRRIGATING SOLUTION DURING SURGICAL PROCEDURES INVOLVING PERFUSION OF THE EYE

U-892 TREATMENT OF CUTANEOUS MANIFESTATIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL)

U-893 CLEVIPREX IS A DIHYDROPYRIDINE CALCIUM CHANNEL BLOCKER INDICATED FOR THE REDUCTION OF BLOOD PRESSURE WHEN ORAL THERAPY IS NOT FEASIBLE OR NOT DESIRABLE

U-894 TREATMENT OF COLD SORES IN PEDIATRIC PATIENTS TWELVE YEARS OF AGE AND OLDER

U-895 TREATMENT OF HIV INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS

U-896 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN TWO YEARS OF AGE AND OLDER

U-897 METHOD OF TREATING TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES IN A ONCE-A-DAY AMOXICILLIN PRODUCT

U-898 USE OF GLUTAMINE TOGETHER WITH GROWTH HORMONE FOR THE TREATMENT OF PATIENTS WITH

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

SHORT BOWEL SYNDROME

- U-899 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA
- U-900 INTEGRASE INHIBITION FOR THE TREATMENT OF HIV INFECTION
- U-901 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
- U-902 USE IN THE TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
- U-903 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN ADULT PATIENTS
- U-904 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE
- U-905 TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY, ASSOCIATED WITH MENOPAUSE
- U-906 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY, LIVER AND HEART ALLOGENIC TRANSPLANTS; TREATMENT OF PATIENTS WITH SEVERE ACTIVE, RHEUMATOID ARTHRITIS; TREATMENT OF ADULT, NONIMMUNOCOMPROMISED PATIENTS WITH SEVERE, RECALCITRANT, PLAQUE PSORIASIS
- U-907 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS IN SUBJECTS 18 YEARS OF AGE AND OLDER
- U-908 PROPHYLAXIS OF ORGAN REJECTION IN PATIENTS RECEIVING ALLOGENEIC RENAL TRANSPLANTS
- U-909 TREATMENT OF CYSTIC FIBROSIS PATIENTS WITH PSEUDOMONAS AERUGINOSA
- U-910 TREATMENT OF METASTATIC CARCINOMA OF THE OVARY AFTER FAILURE OF INITIAL OR SUBSEQUENT CHEMOTHERAPY
- U-911 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL TREATMENT IS TEMPORARILY NOT FEASIBLE
- U-912 SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES
- U-913 TREATMENT OF OVERACTIVE BLADDER WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND FREQUENCY
- U-914 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER
- U-915 TREATMENT OF MUSCULOSKELETAL CONDITIONS
- U-916 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 12 YEARS OR OLDER
- U-917 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS
- U-918 TO TREAT OR PREVENT INFECTIONS CAUSED BY SUSCEPTIBLE BACTERIA USING DELAYED-RELEASE TABLETS CONSISTING OF DOXYCYCLINE HYCLATE COATED PELLETS IN A TABLET
- U-919 FOR THE TREATMENT OF DERMATITIS
- U-920 STEROID-RESPONSIVE INFLAMMATORY OCULAR CONDITIONS FOR WHICH A CORTICOSTEROID IS INDICATED AND WHERE SUPERFICIAL BACTERIAL OCULAR INFECTION OR A RISK OF BACTERIAL OCULAR INFECTION EXISTS
- U-921 TREATMENT OF ACNE VULGARIS
- U-922 FOR THE TREATMENT OF FUNGAL INFECTIONS
- U-923 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION
- U-924 TREATMENT OF MILD TO MODERATE INFECTION CAUSED BY SUSCEPTIBLE STRAINS
- U-925 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA
- U-926 MGT SPECIFIC BACTERIAL INFECTIONS. TREATMENT PTS W/ COMMUNITY ACQUIRED PNEUMONIA OR BACTERIAL SINUSITIS DUE TO CONFIRMED, OR SUSPECTED B-LACTAMASE PRODUCING PATHOGENS & S. PNEUMONIAE WITH REDUCED SUSCEPTIBILITY TO PENICILLIN (MIC=2MC/ML)
- U-927 METHOD FOR INCREASING TEAR PRODUCTION
- U-928 TREATMENT OF BACTERIAL INFECTIOUS DISEASE
- U-929 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER TREATABLE WITH AN SSRI
- U-930 TREATMENT OF IDIOPATHIC THROMBOCYTOPENIC PURPURA (ITP)
- U-931 RELIEF OF MODERATE TO SEVERE ACUTE PAIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-932 PYLERA CAPSULES, IN COMBINATION WITH OMEPRAZOLE ARE INDICATED FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI
- U-933 FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI. THE ERADICATION OF HELICOBACTER PYLORI HAS BEEN SHOWN TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- U-934 IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELL TO THE PERIPHERAL BLOOD FOR COLLECTION AND SUBSEQUENT AUTOLOGOUS TRANSPLANTATION WITH NON-HODGKINS LYMPHOMA AND MULTIPLE MYELOMA
- U-935 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER
- U-936 USE IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELLS TO PERIPHERAL BLOOD FOR COLLECTION & SUBSEQUENT AUTOLOGOUS TRANSPLANTATION IN PATIENTS WITH NON-HODGKIN'S LYMPHOMA & MULTIPLE MYELOMA
- U-937 TREATMENT OF PROSTATE CANCER
- U-938 TREATMENT OF HAIR LOSS AND HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS
- U-939 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING AND STIMULATING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS
- U-940 METHOD TO TREAT AIDS-RELATED KAPOSI'S SARCOMA
- U-941 METHOD TO TREAT OVARIAN CANCER
- U-942 METHOD TO TREAT MULTIPLE MYELOMA
- U-943 GNRH ANTAGONIST INDICATED FOR TREATMENT OF PATIENTS WITH ADVANCED PROSTATE CANCER
- U-944 TREATMENT OF PATIENTS WITH B-CELL CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-945 SEDATIVE-HYPNOTIC AGENT INDICATED FOR MONITORED ANESTHESIA CARE (MAC) SEDATION
- U-946 TREATMENT OF BREAST CANCER
- U-947 WHEN PATIENTS ARE UNABLE TO TAKE THE ORAL FORMULATIONS, PREVACID IV, FOR INJECTION IS INDICATED AS AN ALTERNATIVE FOR THE SHORT-TERM TREATMENT (UP TO 7 DAYS) OF ALL GRADES OF EROSIIVE ESOPHAGITIS
- U-948 TREATMENT OF DIABETES MELLITUS
- U-949 HEALING OF ALL GRADES OF EROSIIVE ESOPHAGITIS (EE) FOR UP TO 8 WEEKS
- U-950 MAINTAIN HEALING OF EROSIIVE ESOPHAGITIS (EE) FOR UP TO 6 MONTHS
- U-951 TREATMENT OF HEARTBURN ASSOCIATED WITH NON-EROSIVE GASTROESOPHAGEAL REFLUX DISEASE (GERD) FOR 4 WEEKS
- U-952 USE AS AN ANALGESIC
- U-953 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION
- U-954 CHRONIC MANAGEMENT OF HYPERURICEMIA IN PATIENTS WITH GOUT. NOT RECOMMENDED FOR THE TREATMENT OF ASYMPTOMATIC HYPERURICEMIA
- U-955 PROPHYLACTIC TREATMENT OF MIGRAINE
- U-956 TREATMENT OF PATIENTS WITH H. PYLORI INFECTION AND DUODENAL ULCER DISEASE
- U-957 A METHOD OF TREATING CANCER IN A PATIENT COMPRISING ADMINISTERING IXABEPILONE OR PHARMACEUTICAL COMPOSITIONS COMPRISING IXABEPILONE
- U-958 METHOD OF TREATING PATIENT COMPRISING MIXING FIRST AND SECOND VIALS OF PRODUCT COMPRISING LYOPHILIZED IXABEPILONE TO PROVIDE AN EPOTHILONE ANALOG SOLUTION, DILUTING SOLUTION WITH A SUITABLE DILUENT TO PREPARE INTRAVENOUS FORMULATION FOR PT
- U-959 METHOD OF TREATING CANCER, IV ADMIN, LYOPHILIZED IXABEPILONE DILUTED, EVERY WEEK OR 3 WEEKS; LYOPHILIZED IXABEPILONE WITH SOLVENT(DEHYDRATED ETHANOL) DILUTED TO CONCENTRATION OF 0.1MG/ML TO 0.9MG/ML
- U-960 METHOD OF TREATING CANCER IN A PATIENT COMPRISING INTRAVENOUSLY ADMINISTERING TO THE PATIENT IXABEPILONE DILUTED IN A PARENTERAL DILUENT
- U-961 METHOD OF TREATING BREAST CANCER BY ADMINISTERING IXABEPILONE; A METHOD OF TREATING A CANCER RESPONSIBLE TO MICROTUBULE STABILIZATION BY ADMINISTERING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

IXABEPILONE

- U-962 SYMBYAX IS INDICATED FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-963 PROZAC AND OLANZAPINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-964 ZYPREXA ZYDIS AND FLUOXETINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-965 USE OF IXABEPILONE IN COMBINATION WITH CAPECITABINE IN TREATMENT OF METASTASIS BREAST CANCER
- U-966 TREATMENT OF ASTHMA (MAINTENANCE AND PROPHYLACTIC THERAPY)
- U-967 A METHOD OF REVERSING SOFT-TISSUE ANESTHESIA I.E. ANESTHESIA OF THE LIP AND TONGUE, AND THE ASSOCIATED FUNCTIONAL DEFICITS RESULTING FROM AN INTRAORAL SUBMUCOSAL INJECTION OF A LOCAL ANESTHETIC
- U-968 A METHOD FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-969 TREATMENT OF MIGRAINE
- U-970 TOPICAL TREATMENT OF LICE INFESTATIONS
- U-971 INDICATED FOR THE ACUTE TREATMENT OF ADULTS WITH SCHIZOPHRENIA
- U-972 MONOTHERAPY OR AS ADJUNCTIVE THERAPY TO LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER
- U-973 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH PIOGLITAZONE AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON PIOGLITAZONE OR METFORMIN ALONE
- U-974 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN
- U-975 TREATMENT OF PULMONARY HYPERTENSION
- U-976 IMPROVEMENT OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES
- U-977 TREATMENT OF ACUTE, UNCOMPLICATED MALARIA INFECTION DUE TO PLASMODIUM FALCIPARUM IN PATIENTS OF 5KG BODYWEIGHT AND ABOVE
- U-978 METHOD OF TREATING HYPONATREMIA
- U-979 RELIEF OF MUSCLE SPASM
- U-980 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN
- U-981 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI-INFLAMMATORY, ANALGESIC, AND ANTIPYRETIC ACTIVITY
- U-982 A METHOD OF TREATING OSTEOPOROSIS
- U-983 METHOD OF TREATING OSTEOPOROSIS IN A POST-MENOPAUSAL WOMAN AT RISK FOR FRACTURE
- U-984 METHOD FOR THE TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND AT RISK FOR BONE FRACTURE
- U-985 TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)
- U-986 TREATMENT OF PATIENTS INFECTED WITH PEDICULUS HUMANUS CAPITIS (HEAD LICE AND THEIR OVA) OF THE SCALP HAIR
- U-987 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH CHRONIC KIDNEY DISEASE ON DIALYSIS
- U-988 TREATMENT OF RHINITIS COMPRISING THE NASAL APPLICATION OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT 7541350
- U-989 FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA
- U-990 TREATMENT OF PROTOZOAL INFECTION
- U-991 TREATMENT OR PROPHYLAXIS OF THROMBOSIS OR EMBOLISMS
- U-992 REDUCTION OF THE RISK OF CARDIOVASCULAR HOSPITALIZATION
- U-993 METHOD OF TREATING INFERTILITY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-994 METHOD OF TREATMENT OF OSTEOPOROSIS WHEREIN THE OSTEOPOROSIS IS STEROID-INDUCED

U-995 METHOD FOR TREATING TYPE II DIABETES BY ADMINISTERING SAXAGLIPTIN

U-996 AN ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL (TC), LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES, AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIX DYSLIPIDEMIA

U-997 TREATMENT OF MAJOR DEPRESSIVE DISORDER BY DOSING AT INTERVALS OF 24 HOURS

U-998 ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL, LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA

U-999 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS

U-1000 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH HYPERLIPIDEMIAS

U-1001 METHOD FOR DELIVERING DRUG TO LUNG OF MAMMAL, COMPRISING ADMINISTERING DRUG PRODUCT BY INHALATION. TREATING A MAMMAL HAVING A CONDITION CAPABLE OF TREATMENT BY INHALATION, COMPRISING ADMINISTERING TO THE LUNG THE DRUG PRODUCT BY INHALATION

U-1002 METHOD OF TREATING INFLAMMATORY CONDITIONS

U-1003 A METHOD OF MYOCARDIAL PERFUSION IMAGING AND INCREASING CORONARY BLOOD FLOW

U-1004 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA

U-1005 METHOD OF TREATING A STAPHYLOCOCCAL INFECTION

U-1006 NEW COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND ADOLESCENTS (12 YEARS OF AGE AND OLDER)

U-1007 METHOD OF TREATING GOUT FLARES

U-1008 APPLICATION OF ANTISEPTIC WITH MOISTURIZERS FOR SURGICAL AND HEALTHCARE PERSONNEL SKIN DISINFECTION

U-1009 METHOD FOR ADMINISTRATION OF TESTOSTERONE

U-1010 TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA DUE TO TETRA HYDROBIOPTERIN RESPONSIVE PHENYLKETONURIA. KUVAN SHOULD BE TAKEN ORALLY WITH FOOD TO INCREASE ABSORPTION

U-1011 USE OF GRANISETRON TRANSDERMAL SYSTEM TO TREAT/PREVENT CHEMOTHERAPY INDUCED NAUSEA AND VOMITING

U-1012 METHOD FOR TREATING INSOMNIA WHILE REDUCING THE RISK OF AN ADVERSE DRUG INTERACTION

U-1013 METHOD OF USING RIBAVIRIN IN COMBINATION WITH PEGYLATED INTERFERON ALPHA-2B TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-1014 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTERFERON ALPHA-2B (PEGYLATED AND NONPEGYLATED) TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-1015 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA

U-1016 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-EXPERIENCED ADULT PATIENTS, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO AN NNRTI AND OTHER ANTIRETROVIRAL AGENTS

U-1017 A METHOD OF TREATING NASAL AND NON-NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

U-1018 TREATMENT OF PULMONARY HYPERTENSION BY INHALATION

U-1019 TREATMENT OF PULMONARY HYPERTENSION

U-1020 METHOD OF USING COLCHICINE FOR THE PROPHYLAXIS OF GOUT FLARES

U-1021 SHORT-TERM TREATMENT (4-8 WEEKS) OF ACTIVE BENIGN GASTRIC ULCER

U-1022 FOR THE PREPARATION OF SKIN PRIOR TO SURGERY; HELPS REDUCE BACTERIA THAT CAN POTENTIALLY CAUSE SKIN INFECTION

U-1023 TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE

U-1024 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP

U-1025 TREATING FREQUENT HEARTBURN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1026 A METHOD OF TREATING HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS.

U-1027 REDUCTION OF ELEVATED PLASMA STEROL AND/OR STANOL LEVELS IN A MAMMAL

U-1028 A METHOD OF DISTRIBUTING SODIUM OXYBATE UNDER CONTROL OF A CENTRAL PHARMACY

U-1029 METHOD FOR TREATING ACUTE ELEVATIONS OF BLOOD PRESSURE IN HUMAN SUBJECT IN NEED THEREOF

U-1030 IMPROVEMENT OF WALKING IN PATIENTS WITH MULTIPLE SCLEROSIS (MS)

U-1031 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA

U-1032 USE OF ROSUVASTATIN CALCIUM FOR THE PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE IN INDIVIDUALS WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH INCREASED RISK FACTORS

U-1033 TOPICAL TREATMENT OF ACNE VULGARIS

U-1034 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS

U-1035 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1036 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH INSULIN

U-1037 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH A PPAR-GAMMA AGONIST

U-1038 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN AND A PPAR-GAMMA AGONIST

U-1039 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN

U-1040 INHIBITION OF THROMBIN IN A PATIENT

U-1041 TREATMENT OF DISORDERS RESPONSIVE TO GROWTH HORMONE

U-1042 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART

U-1043 MANAGEMENT OF MODERATE TO SEVERE PAIN

U-1044 TOPICAL TREATMENT OF SCALP PSORIASIS

U-1045 MAINTENANCE TREATMENT IN PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHO HAVE NOT PROGRESSED ON 1ST-LINE TREATMENT WITH PLATINUM-BASED CHEMOTHERAPY

U-1046 MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES PLATINUM-BASED CHEMOTHERAPY

U-1047 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA (SBCC)

U-1048 WORKS THROUGH THE INDUCTION OF INTERFERON AND OTHER CYTOKINES

U-1049 PROPHYLAXIS OF ORGAN REJECTION IN ADULT PATIENTS AT LOW-MODERATE IMMUNOLOGIC RISK RECEIVING A RENAL TRANSPLANT

U-1050 USE OF METAXALONE FOR TREATMENT OF MUSCULOSKELETAL CONDITIONS

U-1051 TREATMENT OF OROPHARYNGEAL CANDIDIASIS

U-1052 RELIEF OF SIGNS AND SYMPTOMS OF ARTHRITIS AND RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER

U-1053 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER

U-1054 ONYCHOMYCOSIS OF THE TOENAIL CAUSED BY TRICOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES, ONCE DAILY USE FOR 12 CONSECUTIVE WEEKS

U-1055 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH A THIAZOLIDINEDIONE (TZD) AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A TZD OR METFORMIN ALONE

U-1056 TREATMENT OF PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE

U-1057 TREATMENT OF INFLAMMATION AND PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE

U-1058 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

U-1059 ADJUNCTIVE THERAPY TO DIET TO PATIENTS WITH HYPERTRIGLYCERIDEMIA

U-1060 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH ELEVATED CHOLESTEROL AND/OR LIPID LEVELS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1061 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH MIXED DYSLIPIDEMIA

U-1062 ADMINISTRATION OF APPROVED PRODUCT FOR TREATMENT OF ALZHEIMER'S DISEASE

U-1063 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA

U-1064 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA

U-1065 METHOD OF TREATING ANDROGEN RESPONSIVE OR MEDICATED CONDITION IN A MAMMAL BY ADMINISTERING A SAFE & EFFECTIVE AMOUNT OF DUTASTERIDE OR A PHARMACEUTICALLY ACCEPTABLE SOLVATE THEREOF.. CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY

U-1066 METHOD OF TREATING AN ANDROGEN RESPONSE OR MEDIATED DISEASE IN A MAMMAL BY ADMINISTERING AN EFFECTIVE ANDROGEN RESPONSIVE OR MEDICATED DISEASE AMOUNT OF DUTASTERIDE..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERPLASIA

U-1067 TREATMENT OF CANCER

U-1068 TREATMENT OF ASTHMA

U-1069 A METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING AN EXCLUSIVE COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION

U-1070 A METHOD TO CONTROL ABUSE OF A SENSITIVE DRUG BY CONTROLLING WITH A COMPUTER PROCESSOR THE DISTRIBUTION OF THE SENSITIVE DRUG VIA AN EXCLUSIVITY CENTRAL PHARMACY THAT MAINTAINS A CENTRAL DATABASE

U-1071 METHOD OF TREATING BLADDER DYSFUNCTION WITH ONCE A DAY TROSPIUM SALT FORMULATION

U-1072 THE MANAGEMENT OF MODERATE TO SEVERE CHRONIC PAIN IN PATIENTS REQUIRING A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC FOR AN EXTENDED PERIOD OF TIME

U-1073 USE FOR THE TREATMENT OF ASTHMA AND COPD

U-1074 USE OF EXENATIDE MAY RESULT IN REDUCTION IN BODY WEIGHT

U-1075 USE FOR THE TREATMENT OF ASTHMA

U-1076 REDUCE CHRONIC SEVERE DROOLING (I.E., SIALORRHEA) IN PATIENTS WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING

U-1077 PRETREATMENT OF PATIENTS WITH VITAMIN B12 AND FOLIC ACID PRIOR TO PEMETREXED DISODIUM ADMINISTRATION

U-1078 TREATMENT OF ACNE

U-1079 REVLIMID (LENALIDOMIDE) IS INDICATED FOR THE TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)

U-1080 METHOD TO TREAT PULMONARY HYPERTENSION BY ADMINISTERING AMBRISENTAN TO A PATIENT

U-1081 LUMIGAN IS A PROSTAGLANDIN ANALOG INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1082 USE OF A COMBINATION OF TOBRAMYCIN AND DEXAMETHASONE TO TREAT OCULAR INFLAMMATION WHERE AN INFECTION OR RISK OF INFECTION EXISTS

U-1083 ACUTE TREATMENT OF MIGRAINE ATTACKS, WITH OR WITHOUT AURA, AND THE TREATMENT OF CLUSTER HEADACHE EPISODES

U-1084 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER

U-1085 METHOD FOR TREATING IRRITABLE BOWEL SYNDROME AND METHOD FOR TREATING ABDOMINAL DISCOMFORT ASSOCIATED WITH IRRITABLE BOWEL SYNDROME

U-1086 TREATMENT OF AUTOIMMUNE DISEASE

U-1087 DETECTION OF NON-MUSCLE INVASIVE PAPILLARY CANCER OF THE BLADDER BY PHOTODYNAMIC CYSTOSCOPY

U-1088 RELIEF OF MUSCLE SPASM

U-1089 INHIBITION OF THROMBIN

U-1090 LO LOESTRIN FE IS INDICATED FOR THE PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION

U-1091 ASSESSMENT OF BRONCHIAL HYPERRESPONSIVENESS IN PATIENTS 6 YEARS OF AGE OR OLDER WHO DO NOT HAVE CLINICALLY APPARENT ASTHMA

U-1092 TREATMENT OF BREAST CANCER

U-1093 TREATMENT OF PSEUDOBLBAR AFFECT

U-1094 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1095 METHOD OF TREATING OCULAR INFLAMMATION

U-1096 TREATMENT OF PATIENTS WITH METASTATIC BREAST CANCER

U-1097 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH SAXAGLIPTIN AND METFORMIN IS APPROPRIATE

U-1098 METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF TREATING HYPERCALCEMIA

U-1099 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA

U-1100 REDUCTION OF EXCESS ABDOMINAL FAT IN HIV-INFECTED PATIENTS WITH LIPODYSTROPHY

U-1101 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY

U-1102 METHOD OF TREATING CATAPLEXY IN PATIENTS WITH NARCOLEPSY

U-1103 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE

U-1104 USE OF TRAMADOL FOR THE MANAGEMENT OF MODERATE TO MODERATELY SEVERE CHRONIC PAIN

U-1105 TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS FOUR (4) YEARS OF AGE AND OLDER

U-1106 TREATING HYPERTRIGLYCERIDEMIAS WITH REDUCTION OF FOOD EFFECT

U-1107 TREATING HYPERCHOLESTEROLEMIAS WITH REDUCTION OF FOOD EFFECT

U-1108 TREATING TYPE 2 DIABETES MELLITUS WITH EXENATIDE BY STIMULATING INSULIN RELEASE

U-1109 TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL) IN CONNECTION WITH A SPECIAL PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)

U-1110 METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING A COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION

U-1111 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1112 METHOD OF MR IMAGING OF A MAMMAL

U-1113 TREATMENT AND PROPHYLAXIS OF INFLUENZA

U-1114 TREATMENT WITH GABAPENTIN, INCLUDING TREATMENT OF NEUROPATHIC PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH POSTHERPETIC NEURALGIA

U-1115 TREATMENT TO REDUCE THE RISK OF COPD EXACERBATIONS IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS

U-1116 METHOD OF ADMINISTERING COLCHICINE TO FAMILIAL MEDITERRANEAN FEVER PATIENTS

U-1117 TREATMENT OF BREAST CANCER

U-1118 USE FOR THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA

U-1119 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING

U-1120 TO REDUCE GASTROINTESTINAL SIDE EFFECTS ADMINISTER WITH A MEAL; AS STARTING DOSE ADMINISTER ONCE DAILY WITH EVENING MEAL

U-1121 METHOD OF TREATING TRAVELERS' DIARRHEA

U-1122 TREATMENT OF SECONDARILY INFECTED TRAUMATIC SKIN LESIONS DUE TO S. AUREUS AND S. PYOGENES

U-1123 TREATMENT OF ALCOHOL DEPENDENCE

U-1124 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE, FOLLOWING OPIOID DETOXIFICATION

U-1125 METHOD FOR THE DETECTION OF NEUROENDOCRINE TUMORS

U-1126 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE RECEIVED PRIOR CHEMOTHERAPY CONTAINING DOCETAXEL

U-1127 TREATMENT OF PATENT DUCTUS ARTERIOSUS

U-1128 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION IN COMBINATION WITH PEGINTERFERON ALFA AND RIBAVIRIN IN ADULT PATIENTS (>=18 YEARS OF AGE) WITH COMPENSATED LIVER DISEASE

U-1129 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT

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PATENT USE

- NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1130 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR A NIGHT WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1131 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1132 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1133 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1134 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1135 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A) AND INCREASE OF HDL-C
- U-1136 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1137 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1138 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1139 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1140 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1141 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1142 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1143 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR A T NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1144 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1145 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1146 REDUCTION IN TG WITH REDUCED FLUSHING BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1147 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1148 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1149 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1150 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, TG, LP(A), AND INCREASE OF HDL-C
- U-1151 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, LP(A), AND INCREASE OF HDL-C
- U-1152 CYANOCOBALAMIN ADMINISTRATION THROUGH NASAL INFUSION
- U-1153 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS, IS INDICATED FOR THE TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTION IN ANTIRETROVIRAL TREATMENT-NAIVE ADULT PATIENTS, AS SET FORTH IN THE LABELING, INCLUDING I&U SECTION
- U-1154 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMORS, RENAL CELL CARCINOMA AND ADVANCED PANCREATIC NEUROENDOCRINE TUMORS, WITH SUNITINIB

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1155	USE OF THALIDOMIDE IN TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL)
U-1156	TO REDUCE BLOOD PHENYLALANINE (PHE) LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA (HPA)
U-1157	RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES IN ADULTS AND CHILDREN 2 YEARS OF AGE AND OLDER AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
U-1158	RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
U-1159	RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES, SWELLING OF THE NASAL PASSAGES AND SINUS CONGESTION AND PRESSURE IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
U-1160	RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER AND 12 YEARS OF AGE AND OLDER
U-1161	FOR THE TREATMENT AND PROPHYLAXIS OF GOUT FLARES & THE TREATMENT OF FAMILIAL MEDITERRANEAN FEVER
U-1162	TREATMENT OF SEBORRHEIC DERMATITIS OF THE SCALP
U-1163	METHOD OF TREATING THROMBOSIS
U-1164	METHOD OF TREATING AN ARGATROBAN TREATABLE CONDITION
U-1165	USE FOR THE TREATMENT OF MULTIPLE MYELOMA
U-1166	A METHOD FOR TREATMENT OF GOUT FLARES DURING PROPHYLAXIS
U-1167	PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT)
U-1168	THE LONG TERM, ONCE-DAILY MAINTENANCE BRONCHODILATOR TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
U-1169	MANAGEMENT OF BREAKTHROUGH PAIN IN CANCER PATIENTS 18 YEARS OF AGE AND OLDER WHO ARE RECEIVING AND TOLERANT TO OPIOID THERAPY FOR THEIR UNDERLYING PERSISTENT CANCER PAIN
U-1170	TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
U-1171	REDUCTION OF THE RATE OF THROMBOTIC EVENTS IN PATIENTS WITH ACUTE CORONARY SYNDROME
U-1172	TO REDUCE ELEVATED TOTAL-C, APO B, AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE IN COMBINATION WITH A STATIN
U-1173	TO REDUCE ELEVATED TOTAL-C, LDL-C, APO B AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE ALONE OR IN COMBINATION WITH A STATIN OR WITH FENOFIBRATE
U-1174	ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION, 0.9% SODIUM CHLORIDE INJECTION, OR FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO ADMINISTRATION
U-1175	REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
U-1176	TREATMENT OR PREVENTION OF STROKE
U-1177	REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
U-1178	RELIEF OF MODERATE TO SEVERE CHRONIC PAIN
U-1179	TREATMENT OF A CANCER MEDIATED BY AN ANAPLASTIC LYMPHOMA KINASE (ALK)
U-1180	TREATMENT OF THE FOLLOWING INFECTIONS: COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS
U-1181	A METHOD OF TREATING OR PREVENTING OCULAR PAIN IN A PATIENT
U-1182	TREATMENT OF CYCLIC HEAVY MENSTRUAL BLEEDING
U-1183	A METHOD FOR ADMINISTERING FOLLICLE STIMULATING HORMONE (FSH) FOR OVARIAN FOLLICLE OR TESTICULAR STIMULATION IN THE HUMAN
U-1184	TREATMENT OF ERECTILE DYSFUNCTION AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1185 TREATMENT OF OPIOID-INDUCED CONSTIPATION

U-1186 ADMINISTRATION OF AN INHALABLE POWDER COMPRISING TIOTROPIUM VIA DEVICE

U-1187 TREATMENT OF PATHOLOGICAL STATE BY ANTAGONIZING BRADYKININ RECEPTOR INCLUDING TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA (HAE)

U-1188 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE

U-1189 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH METFORMIN

U-1190 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH INSULIN

U-1191 METHOD OF TX TYPE 2 DM IN PTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBO WITH AN AGENT ACTING ON AN ATP-DEPENDENT CHANNEL IN BETA CELLS SUCH AS A SULFONYLUREA(INCL GLIPIZIDE, GLIMEPIRIDE & GLYBURIDE)

U-1192 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A SULFONYLUREA (SUCH AS GLIPIZIDE, GLIMEPIRIDE AND GLYBURIDE)

U-1193 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A PPAR-GAMMA AGONIST (SUCH AS PIOGLITAZONE AND ROSIGLITAZONE)

U-1194 METHOD FOR TREATING INSOMNIA

U-1195 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5, WHICH MAY RESULT IN RENAL OSTEODYSTROPHY, WHILE AVOIDING HYPERPHOSPHATEMIA

U-1196 RELIEF OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS AND OSTEOARTHRITIS AND TO DECREASE RISK OF DEVELOPING UPPER GASTROINTESTINAL ULCERS IN PATIENTS WHO ARE TAKING IBUPROFEN FOR THOSE INDICATIONS

U-1197 METHOD OF TREATMENT OF CHILDREN WITH CENTRAL PRECOCIOUS PUBERTY

U-1198 RECTIV IS A NITRATE VASODILATOR INDICATED FOR THE TREATMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH CHRONIC ANAL FISSURE

U-1199 TREATMENT AND PREVENTION OF POSTMENOPAUSAL OR GLUCOCORTICOID-INDUCED OSTEOPOROSIS AND TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

U-1200 REDUCING THE RISK OF STROKE AND SYSTEMIC EMBOLISM

U-1201 FOR THE TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS

U-1202 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME

U-1203 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT

U-1204 TREATMENT OF UVEITIS

U-1205 TREATMENT OF MACULAR EDEMA

U-1206 DELIVERING AN OCULAR IMPLANT AS DESCRIBED IN THE DOSAGE AND ADMINISTRATION SECTION OF THE APPROVED LABELING OF OZURDEX

U-1207 INFANT USE AGED 1 MONTH TO LESS THAN ONE YEAR, GERD AND EROSIIVE ESOPHAGITIS

U-1208 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS

U-1209 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER

U-1210 USE OF REVLIMID (LENALIDOMIDE) WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO REVLIMID (LENALIDOMIDE)

U-1211 USE OF REVLIMID (LENALIDOMIDE) TO INHIBIT THE SECRETION OF PRO-INFLAMMATORY CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA

U-1212 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA AND TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)

U-1213 TOPICAL TREATMENT OF SEBORRHEIC DERMATITIS IN IMMUNOCOMPETENT PATIENTS 12 YEARS OF AGE AND OLDER

U-1214 METHOD FOR RELIEVING CONSTIPATION IN A HUMAN PATIENT THAT COMPRISES

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MCG+/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT

U-1215 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)

U-1216 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA

U-1217 METHOD OF INCREASING HAIR GROWTH

U-1218 METHOD OF STIMULATING HAIR GROWTH

U-1219 METHOD OF INCREASING THE NUMBER OF HAIRS

U-1220 TREATMENT OF RENAL CELL CARCINOMA

U-1221 TO STIMULATE THE IMMUNE SYSTEM TO INDUCE T CELL PROLIFERATION

U-1222 TO INHIBIT THE PROLIFERATIVE ACTIVITY OF NEOPLASTIC CELLS

U-1223 METHOD FOR TREATING TYPE 2 DIABETES USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENATIDE

U-1224 REDUCTIONS IN BODY WEIGHT ARE OBSERVED WITH EXENATIDE

U-1225 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING PARTIAL LARGE OR SMALL BOWEL RESECTION SURGERY WITH PRIMARY ANASTOMOSIS

U-1226 A METHOD OF PROVIDING A PREDETERMINED CONCENTRATION OF NITRIC OXIDE TO A PATIENT

U-1227 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE

U-1228 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE ALONE OR IN COMBINATION WITH INSULIN

U-1229 TREATMENT OF MILDLY TO MODERATELY ACTIVE ULCERATIVE COLITIS IN MALE PATIENTS

U-1230 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT

U-1231 TREATMENT OF MODERATE-TO-SEVERE PRIMARY RESTLESS LEG SYNDROME IN ADULTS

U-1232 USE AS ANTICOAGULANT IN PTS W/ UNSTABLE ANGINA UNDERGOING PTCA; W/ PROVISIONAL USE OF GLYCOPROTEIN IIB/IIIA INHIBITOR, AS ANTICOAGULANT IN PTS UNDERGOING PCI AND FOR PTS W/, OR AT RISK OF, HIT/HITTS UNDERGOING PCI.INTENDED FOR USE W/ASPIRIN

U-1233 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION, ADMINISTERED WITH FOOD

U-1234 FOR REDUCING TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES, AND TREATING HYPERTRIGLYCERIDEMIA

U-1235 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION

U-1236 USE OF THALOMID (THALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA

U-1237 COMBO W/ OTHER ANTIRETROVIRALS FOR TX OF HIV-1 IN ANTIRETROVIRAL TX-EXPERIENCED PT 6 YEARS UP, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITOR AND OTHER ANTIRETROVIRALS

U-1238 TREATMENT OF ANEMIA DUE TO CHRONIC KIDNEY DISEASE

U-1239 MAGNETIC RESONANCE IMAGING OF THE LIVER

U-1240 TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION

U-1241 MANAGEMENT OF MODERATE TO SEVERE PAIN BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED

U-1242 PREVENTION OF RESPIRATORY DISTRESS (RDS) IN PREMATURE INFANTS

U-1243 WITH DRY HANDS, GENTLY REMOVE THE SUPRENZA (PHENTERMINE HYDROCHLORIDE ODT) TABLET FROM THE BOTTLE. IMMEDIATELY PLACE THE SUPRENZA TABLET ON TOP OF THE TONGUE WHERE IT WILL DISSOLVE, THEN SWALLOW WITH OR WITHOUT WATER

U-1244 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH SULFONYLUREA

U-1245 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH PIOGLITAZONE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1246 SINGLE DOSE ADMINISTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA

U-1247 MANAGEMENT OF POSTHERPETIC NEURALGIA (PHN) IN ADULTS

U-1248 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL MEDICATION ON THE SAME KNEE

U-1249 TREATMENT OF MALE PATIENT HAVING A DISEASE OR CONDITION RESPONSIVE TO A TERATOGENIC DRUG

U-1250 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY OR SPINAL CORD INJURY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA

U-1251 A METHOD OF CONTROLLING POSTOPERATIVE OCULAR PAIN AND BURNING/STINGING IN A PATIENT

U-1252 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE

U-1253 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY

U-1254 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY CONTROLLING WEIGHT GAIN

U-1255 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY

U-1256 TREATMENT OF SEBORRHEIC DERMATITIS

U-1257 TREATMENT OF OPHTHALMIC DISORDERS

U-1258 VISUALIZATION DURING VITRECTOMY PROCEDURES

U-1259 PROPHYLAXIS OF HIV-1 INFECTION

U-1260 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY

U-1261 REDUCTION OF THE RISK OF HOSPITALIZATION FOR ATRIAL FIBRILLATION

U-1262 USE OF QSYMIA (PHENTERMINE AND TOPIRAMATE) FOR WEIGHT MANAGEMENT, INCLUDING, BUT NOT LIMITED TO EFFECTING WEIGHT LOSS, TREATING OBESITY, AND/OR TREATING OVERWEIGHT

U-1263 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) OR CHRONIC BRONCHITIS

U-1264 TREATMENT OF A RESPIRATORY DISEASE

U-1265 PATENTED METHOD OF USING REPAGLINIDE IN COMBINATION WITH METFORMIN AS INDICATED FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1266 METHOD OF TREATING MIDDLE-OF-THE-NIGHT INSOMNIA

U-1267 TREATMENT OF RHEUMATOID ARTHRITIS BY DELAYED RELEASE FORMULATION OF 1MG OR 2MG OF PREDNISONE

U-1268 TREATMENT OF PULMONARY, GASTROINTESTINAL AND/OR RHEUMATOLOGICAL DISEASES OR CONDITIONS BY USE OF DELAYED RELEASE FORMULATIONS OF 1MG OR 2MG PREDNISONE

U-1269 TREATMENT OF RHEUMATOLOGIC, ALLERGIC, PULMONARY, GASTROINTESTINAL, DERMATOLOGIC DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 5MG PREDNISONE TABLET

U-1270 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH INSULIN (WITH OR WITHOUT METFORMIN AND/OR PIOGLITAZONE)

U-1271 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE (PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES

U-1272 TREATMENT OF SIGNS AND SYMPTOMS OF PARKINSON'S DISEASE BY APPLICATION OF CLAIMED TRANSDERMAL SYSTEM

U-1273 TREATMENT OF RESTLESS LEGS SYNDROME BY APPLICATION OF CLAIMED TRANSDERMAL DELIVERY SYSTEM

U-1274 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY DUE TO CYSTIC FIBROSIS OR OTHER CONDITIONS

U-1275 TREATMENT OF CHRONIC HEPATITIS B IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER

U-1276 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1277 METHOD OF INCREASING EYELASH GROWTH INCLUDING LENGTH, THICKNESS, DARKNESS AND/OR NUMBER OF EYELASHES BY ADMINISTERING BIMATOPROST TO AN EYELID MARGIN

U-1278 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULTS

U-1279 TREATMENT OF HIV INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE

U-1280 USE OF A CALCIPOTRIENE CONTAINING FOAM FOR THE TREATMENT OF PSORIASIS

U-1281 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE PREVIOUSLY RECEIVED DOCETAXEL

U-1282 PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING

U-1283 A METHOD OF TREATING CHRONIC MYELOGENOUS LEUKEMIA

U-1284 A METHOD OF TREATING A NEOPLASM

U-1285 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS

U-1286 A METHOD OF REDUCING THE RISK OF PULMONARY EDEMA IN PATIENTS IN NEED OF TREATMENT WITH INHALED NITRIC OXIDE

U-1287 METHOD OF REDUCING TG LEVELS IN PATIENT SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA

U-1288 TREATMENT OF ERECTILE DYSFUNCTION BY ADMINISTERING A FILM-COATED TABLET

U-1289 MANAGEMENT OF MODERATE TO SEVERE ACUTE PAIN

U-1290 TREATMENT OF LUNG CANCER

U-1291 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL) IN PATIENTS WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE (15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION

U-1292 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 1, 2, OR 5 MG PREDNISONE TABLET

U-1293 A METHOD OF LOWERING INTRAOCULAR PRESSURE IN A PATIENT WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1294 METHOD OF TREATING GLAUCOMA IN A PATIENT

U-1295 A METHOD OF TREATING A PATIENT WITH GLAUCOMA OR OCULAR HYPERTENSION

U-1296 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION

U-1297 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION BY INHIBITING ENDOTHELIN RECEPTORS

U-1298 ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES

U-1299 TREATMENT OF PATIENTS WITH LEUKEMIA INCLUDING CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)

U-1300 TREATMENT OF PATIENTS WITH TYROSINE KINASE INHIBITOR (TKI) RESISTANT OR INTOLERANT CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)

U-1301 TREATMENT OF DEEP VEIN THROMBOSIS (DVT)

U-1302 TREATMENT OF PULMONARY EMBOLISM (PE)

U-1303 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM

U-1304 USE OF ONCE-A-DAY AMOXICILLIN PRODUCT TO TREAT TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES

U-1305 TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS, AND TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER, CO-ADMINISTERED WITH RITONAVIR (PREZISTA/RITONAVIR) AND WITH OTHER ANTIRETROVIRAL AGENTS

U-1306 TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY

U-1307 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE ADULT PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY

U-1308 MULTIPLE MYELOMA

U-1309 BONE METASTASES

U-1310 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1311 METHOD OF TREATING CYSTIC FIBROSIS

U-1312 USE FOR THE TREATMENT OF HYPERGLYCEMIA

U-1313 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1314 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER

U-1315 THE LONG TERM TREATMENT OF PROPHYLACTIC MANAGEMENT OF OCULAR HYPERTENSION AND GLAUCOMA

U-1316 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES

U-1317 TREATMENT OF HYPERCHOLESTEROLEMIA, HYPERLIPIDEMIA AND HYPERLIPOPROTEINEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

U-1318 TREATMENT OF HYPERCHOLESTEROLEMIA BY DECREASING THE AMOUNT OR ACTIVITY OF MICROSOMAL TRIGLYCERIDE TRANSFER PROTEIN IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

U-1319 SYMPTOMATIC RELIEF OF NON-INFECTIOUS DIARRHEA

U-1320 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME WHO ARE DEPENDENT ON PARENTERAL SUPPORT

U-1321 TREATMENT OF PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS

U-1322 METHOD OF REDUCING OCULAR HYPERTENSION

U-1323 REDUCING THE RISK OF STROKE

U-1324 MANAGEMENT OF CYSTIC FIBROSIS PATIENTS

U-1325 INDUCTION OF REMISSION IN PATIENTS WITH ACTIVE, MILD TO MODERATE ULCERATIVE COLITIS

U-1326 METHOD OF INDUCING CONTRACEPTION IN A FEMALE OF REPRODUCTIVE AGE WHO HAS NOT YET REACHED PREMENOPAUSE

U-1327 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF, USING A FLOWABLE HYDROGEL FORMULATION

U-1328 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF

U-1329 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER

U-1330 METHODS OF TREATING LIPID METABOLISM AND GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1331 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1332 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1333 METHODS OF LOWERING ELEVATED POST PRANDIAL BLOOD GLUCOSE LEVELS COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR

U-1334 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-1335 METHODS OF MODIFYING GLUCOSE METABOLISM AND TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR AND ONE OR MORE OTHER THERAPEUTIC AGENTS SUCH AS METFORMIN

U-1336 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR AND METFORMIN

U-1337 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING ALOGLIPTIN

U-1338 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING A COMPOUND SUCH AS ALOGLIPTIN

U-1339 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

U-1340 METHODS OF TREATING LIPID METABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN

U-1341 METHODS OF TREATING GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN

U-1342 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN

U-1343 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN

U-1344 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN PREPARATION

U-1345 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH A DOSAGE UNIT COMPRISING 24MICROG+/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT

U-1346 USE OF FEBUXOSTAT FOR THE MANAGEMENT OF HYPERURICEMIA IN PATIENTS SUFFERING FROM GOUT AND, WHEN USED WITH THEOPHYLLINE WITHOUT THE NEED FOR DOSE ADJUSTMENT OF THEOPHYLLINE

U-1347 TREATMENT OF A SKIN DISORDER

U-1348 TREATMENT OF OSTEOARTHRITIS

U-1349 TREATMENT OF JUVENILE RHEUMATOID ARTHRITIS

U-1350 TREATMENT OF ANKYLOSING SPONDYLITIS

U-1351 TREATMENT OF ACUTE PAIN

U-1352 TREATMENT OF PRIMARY DYSMENORRHEA

U-1353 ADJUNCTIVE THERAPY TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LOW DENSITY LIPOPROTEIN-CHOLESTEROL, APOLIPOPROTEIN B, TOTAL CHOLESTEROL, AND NON-HIGH DENSITY LIPOPROTEIN CHOLESTEROL IN PTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

U-1354 INHIBITION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN HYPERSTIMULATION WITH FSH

U-1355 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHOD FOR TREATING A RESPIRATORY DISEASE IN A CHILD

U-1356 TREATMENT OF NASAL SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER. TREATMENT OF NASAL SYMPTOMS ASSOCIATED W PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER

U-1357 TREATMENT OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHODS FOR TREATING A RESPIRATORY DISEASE IN A CHILD

U-1358 TREATMENT OF BACTERIAL INFECTIONS IN THE NASAL PASSAGE OF ADULT PATIENTS AND HEALTH CARE WORKERS WITH METHICILLIN RESISTANT S. AUREUS

U-1359 USE OF POMALIDOMIDE TO INHIBIT THE SECRETION OF PRO-INFLAMMATION CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA

U-1360 USE OF POMALIDOMIDE FOR THE TREATMENT OF MULTIPLE MYELOMA

U-1361 USE OF POMALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO POMALIDOMIDE

U-1362 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED-RELEASE 1,2, OR 5MG PREDNISONE TABLET

U-1363 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING/STINGING FOLLOWING CORNEAL SURGERY

U-1364 MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)

U-1365 PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT

U-1366 TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY TO

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

ANOVULATORY INFERTILE WOMEN

- U-1367 METHOD OF ADMINISTERING FSH FOR THE TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY IN ANOVULATORY INFERTILE WOMEN
- U-1368 TREATMENT OF SOLID EXCRETORY SYSTEM TUMORS; ADVANCED RENAL CELL CARCINOMA (RCC), AFTER FAILURE OF TREATMENT WITH SUNITINIB OR SORAFENIB
- U-1369 TREATMENT OF VAGINAL SYMPTOMS OF UROGENITAL ATROPHY BY ORALLY ADMINISTERING OSPEMIFENE WITH FOOD TO ENHANCE BIOAVAILABILITY OF OSPEMIFENE
- U-1370 TREATMENT OF DYSpareunia ASSOCIATED WITH MENOPAUSE
- U-1371 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH ELEVATED INTRAOCULAR PRESSURE OR GLAUCOMA
- U-1372 ADMINISTRATION WITHOUT FOOD FOR TREATMENT OF HIV-1 INFECTION
- U-1373 METHOD OF TREATING ACETAMINOPHEN OVERDOSE WITH ACETYLCYSTEINE SOLUTIONS
- U-1374 TREATMENT OF PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML)
- U-1375 ADASUVE IS A TYPICAL ANTIPSYCHOTIC INDICATED FOR THE ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I DISORDER IN ADULTS
- U-1376 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS
- U-1377 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA
- U-1378 TREATMENT OF A NITROGEN METABOLISM DISORDER
- U-1379 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS
- U-1380 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS WHEREIN THE PATIENT HAS CARDIOVASCULAR DISEASE
- U-1381 USE OF PRASUGREL AND ASPIRIN IN PATIENTS REQUIRING THE REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS
- U-1382 TREATMENT OF NAUSEA AND VOMITING OF PREGNANCY IN WOMEN WHO DO NOT RESPOND TO CONSERVATIVE MANAGEMENT
- U-1383 DOSAGE ADJUSTMENT OF A NITROGEN SCAVENGING DRUG IN THE TREATMENT OF A UREA CYCLE DISORDER
- U-1384 METHOD OF TREATING MULTIPLE SCLEROSIS
- U-1385 METHOD OF TREATING AN AUTOIMMUNE DISEASE SELECTED FROM AUTOIMMUNE POLYARTHRITIS AND MULTIPLE SCLEROSIS BUT NOT TREATING PSORIATIC ARTHRITIS
- U-1386 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF A PERSON IN NEED THEREOF
- U-1387 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEVERE HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1388 TREATMENT OF PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEVERE HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1389 ELLA IS A PROGESTERONE AGONIST/ANTAGONIST EMERGENCY CONTRACEPTION INDICATED FOR THE PREVENTION OF PREGNANCY FOLLOWING UNPROTECTED INTERCOURSE OR A KNOWN OR SUSPECTED CONTRACEPTIVE FAILURE. ELLA CAN BE TAKEN WITH OR WITHOUT FOOD
- U-1390 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF AN ADULT MALE SUBJECT IN NEED THEREOF
- U-1391 METHOD FOR TREATING OPIOID-INDUCED CONSTIPATION
- U-1392 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH OPIOID-INDUCED CONSTIPATION
- U-1393 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION
- U-1394 METHOD FOR RELIEVING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION THAT COMPRISES ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MICROG+/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1395 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- CONSTIPATION WITH A DOSAGE UNIT COMPRISING 24MICROG +/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1396 TREATMENT OF ADVANCED HORMONE RECEPTOR POSITIVE, HER2-NEGATIVE BREAST CANCER IN COMBINATION WITH EXEMESTANE AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
- U-1397 USE AS AN ANTISEPTIC FOR THE PREPARATION OF A PATIENT'S SKIN PRIOR TO SURGERY
- U-1398 METHOD OF TREATING CHRONIC HEPATITIS C
- U-1399 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS BY ADMINISTERING A TOTAL DAILY DOSE IN TWO DIVIDED DOSES
- U-1400 FOR THE TREATMENT OF PRIMARY HYPERLIPIDEMIA, MIXED HYPERLIPIDEMIA OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1401 INDICATED FOR LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PTS WITH A HISTORY OF EXACERBATIONS
- U-1402 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR INDOLENT B-CELL NON-HODGKIN LYMPHOMA (NHL)
- U-1403 FIRST-LINE TREATMENT OF METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) WITH EGFR EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- U-1404 METHOD FOR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION BY OPENING CIC CHANNELS
- U-1405 THERAPEUTIC TREATMENT OF BONE METASTASES
- U-1406 TREATMENT OF MELANOMA
- U-1407 TREATMENT OF NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH + CML)
- U-1408 TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE OR OLDER
- U-1409 TREATMENT OF HIV-1 BY ONCE DAILY ADMINISTRATION
- U-1410 TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES
- U-1411 THIS DRUG IS ADMINISTERED BY SUBLINGUAL ROUTE TO HUMANS FOR MAINTENANCE TREATMENT OF OPIOID DEPENDENCE
- U-1412 TREATMENT OF ATOPIC DERMATITIS
- U-1413 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO INFUSION
- U-1414 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MANTLE CELL LYMPHOMA (MCL)
- U-1415 TREATING A PATIENT HAVING A CONDITION SUSCEPTIBLE TO TREATMENT WITH METHYLPHENIDATE, SUCH AS ADHD, BY ADMINISTERING THE FORMULATION RECITED IN CLAIMS 1 OR 2
- U-1416 USE OF FENOFIBRATE FOR REDUCING ELEVATED TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES
- U-1417 USE FOR TREATMENT OF HELICOBACTER INFECTIONS
- U-1418 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAFV600E MUTATION AS DETECTED BY AN FDA APPROVED TEST
- U-1419 TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE LIFE THREATENING
- U-1420 METHOD OF ONCE A DAY ADMINISTRATION
- U-1421 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE
- U-1422 METHOD OF TREATING PATIENTS NEEDING AN IRON SUPPLEMENT
- U-1423 AMYVID IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET) IMAGING OF THE BRAIN TO ESTIMATE BETA-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT
- U-1424 LONG-TERM, ONCE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS
- U-1425 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1426 USE FOR TREATMENT OF DIAPER DERMATITIS COMPLICATED BY CANDIDIASIS

U-1427 ALKYLATING DRUG INDICATED FOR THE TOPICAL TREATMENT OF STAGE IA AND IB MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN DIRECTED THERAPY

U-1428 TOPICAL TREATMENT OF FACIAL ERYTHEMA OF ROSACEA

U-1429 TREATMENT OF PATIENTS WITH BREAST CANCER WHOSE TUMORS OVEREXPRESS THE HER2 RECEPTOR

U-1430 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL AND PERENNIAL ALLERGIC RHINITIS

U-1431 METHOD OF TREATING HYPERGLYCEMIA TO IMPROVE GLYCEMIC CONTROL IN A PATIENT BY ORAL ADMIN OF ONCE A DAY OSMOTIC DOSAGE FORM OF GLIPIZIDE WITH POLYETHYLENE OXIDE, HYDROXYPROPYLMETHYLCELLULOSE, CELLULOSE ACETATE, AND SODIUM CHLORIDE

U-1432 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX

U-1433 IMPROVEMENTS OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS

U-1434 TREATMENT OF PANCREATIC CANCER

U-1435 COMBINATION USE OF TOPICAL DICLOFENAC ON THE KNEE AND ADMINISTRATION OF AN ORAL NSAID.

U-1436 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL AGENT SELECTED FROM SUNSCREEN AND INSECT REPELLANT

U-1437 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE DILUENT FOR FLOLAN OR STERILE DILUENT FOR EPOPROSTENOL SODIUM PRIOR TO ADMINISTRATION

U-1438 ZINGO INTRADERMAL INJECTION SYSTEM IS A DRUG DELIVERY SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION

U-1439 METHOD OF TREATING AN AFFECTIVE DISORDER SUCH AS DEPRESSION

U-1440 USE OF INGENOL MEBUTATE TO TREAT ACTINIC KERATOSIS

U-1441 A METHOD OF TREATING OR REDUCING OCULAR PAIN AND BURNING/STINGING

U-1442 SUBCUTANEOUS INJECTION OF METHOTREXATE

U-1443 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS

U-1444 A DOSING REGIMEN OF AGGRASTAT (TIROFIBAN HYDROCHLORIDE) (25MCG/KG FOLLOWED BY 0.15MCG/KG/MIN INFUSION) TO REDUCE THE RATE OF THROMBOTIC CORONARY EVENTS ASSOCIATED WITH ACUTE CORONARY SYNDROME (ACS) IN PATIENTS WITH NON-ST ELEVATION ACS

U-1445 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 1% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION

U-1446 METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING ADMINISTERING MACITENTAN IN COMBINATION WITH A COMPOUND HAVING PHOSPHODIESTERASE-5 INHIBITORY PROPERTIES

U-1447 TREATING PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIA

U-1448 TREATING SEVERE HYPERTRIGLYCERIDEMIA

U-1449 METHOD OF ALLEVIATING A SKIN CONDITION

U-1450 TREATMENT OF ALLERGIC RHINITIS SYMPTOMS

U-1451 APPROVED INDICATIONS: APTIOM (ESLICARBAZEPINE ACETATE) IS INDICATED AS ADJUNCTIVE TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH EPILEPSY. PATENT CLAIMS: IN A METHOD OF TREATING A SUBJECT AFFLICTED WITH EPILEPSY

U-1452 METHOD FOR CHRONIC WEIGHT MANAGEMENT

U-1453 A METHOD OF TREATING HYPOXIC RESPIRATORY FAILURE BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT

U-1454 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1455 TREATMENT OF PERIANAL WARTS

U-1456 TREATMENT OF MANTLE CELL LYMPHOMA

U-1457 A METHOD OF PURGING A NITRIC OXIDE DELIVERY SYSTEM

U-1458 A METHOD OF REDUCING INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1459 TREATMENT OF CARCINOMA OF THE THYROID

U-1460 TREATMENT OF HERPES LABIALIS

U-1461 A METHOD OF GENERATING AN INJECTABLE FOAM OF CONTROLLED DENSITY AND BUBBLE SIZE

U-1462 A METHOD OF USING A SCLEROSING AGENT FOR THE TREATMENT OF INCOMPETENT GREAT SAPHENOUS VEINS, ACCESSORY SAPHENOUS VEINS AND VISIBLE VARICOSITIES OF THE GREAT SAPHENOUS (GSV) SYSTEM ABOVE AND BELOW THE KNEE

U-1463 A METHOD OF INTRAVENOUS INJECTION USING ULTRASOUND GUIDANCE, ADMINISTERED VIA A SINGLE CANNULA INTO THE LUMEN OF THE TARGET INCOMPETENT TRUNK VEINS OR BY DIRECT INJECTION INTO VARICOSITIES

U-1464 TREATMENT OF OPIOID DEPENDENCE/SUBLINGUAL OR BUCCAL APPLICATION

U-1465 USE OF THALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO THALIDOMIDE

U-1466 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER

U-1467 METHOD OF TREATING HEPATITIS C

U-1468 CONTROL OF PHOSPHOROUS LEVELS IN PATIENTS

U-1469 USE OF PHOSLYRA FOR REDUCTION OF SERUM PHOSPHOROUS IN PATIENTS

U-1470 FOR THE TREATMENT OF HEPATITIS C

U-1471 A METHOD FOR TREATING CARDIOVASCULAR DISEASE COMPRISING ADMINISTERING A RECONSTITUTED LYOPHILIZED PHARMACEUTICAL COMPOSITION COMPRISING EPOPROSTENOL, ARGININE AND SODIUM HYDROXIDE.

U-1472 INTENSIVE CARE UNIT SEDATION, INCLUDING SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES

U-1473 MANAGEMENT OF RISK OF DRONEDARONE/BETA-BLOCKER INTERACTION IN PATIENTS IN SINUS RHYTHM WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF

U-1474 A METHOD FOR THE TREATMENT OF A PATIENT SUFFERING FROM A DISEASE TREATABLE WITH ROTIGOTINE, COMPRISING APPLYING THE CLAIMED TRANSDERMAL DELIVERY SYSTEM (TDS) TO THE SKIN OF THE PATIENT

U-1475 USE OF ORENITRAM FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1).

U-1476 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA.

U-1477 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL PRESCRIPTION MEDICATION ON THE SAME KNEE

U-1478 METHOD OF REDUCING TG LEVELS IN PATIENT ON STATIN THERAPY SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA

U-1479 INCREASE TEAR PRODUCTION TO TREAT PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).

U-1480 TREATMENT OF ADVANCED RENAL CELL CARCINOMA

U-1481 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE

U-1482 DICLOFENAC POTASSIUM FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1483 INCREASE TEAR PRODUCTION IN PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).

U-1484 COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND CHILDREN (6 YEARS OF AGE AND OLDER)

U-1485 TREATING A SUBJECT UNDERGOING ABDOMINAL SURGERY BY ADMINISTERING ALVIMOPAN TO ACCELERATE THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1486 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER

U-1487 METHOD OF INCREASING EYELASH GROWTH

U-1488 USE OF TOPICAL DICLOFENAC FOR TREATING PAIN

U-1489 USE OF TOPICAL DICLOFENAC ON A JOINT FOR TREATING OSTEOARTHRITIS

U-1490 FOR USE IN PATIENTS HAVING SYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER, WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE

U-1491 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA

U-1492 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER

U-1493 METHOD FOR PREVENTING ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS

U-1494 SUBLINGUAL OR BUCCAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE

U-1495 RISK REDUCTION OF REBLEEDING IN PTS FOLLOWING THERAPEUTIC ENDOSCOPY FOR ACUTE BLEEDING GASTRIC OR DUODENAL ULCERS IN ADULTS.

U-1496 METHOD TO TREAT HEMANGIOMA.

U-1497 NEURACEQ IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET) IMAGING OF THE BRAIN TO ESTIMATE P-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT

U-1498 METHOD OF TREATING PATIENTS WITH GASTRIC RETENTIVE DOSAGE FORM

U-1499 MANAGEMENT OF ACUTE PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA

U-1500 TESTOSTERONE REPLACEMENT THERAPY IN ADULT MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE; PRIMARY HYPOGONADISM (CONGENITAL OR ACQUIRED); HYPOGONADOTROPIC HYPOGONADISM (CONGENITAL OR ACQUIRED).

U-1501 PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM

U-1502 PROPHYLAXIS OF PULMONARY EMBOLISM

U-1503 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN

U-1504 USE OF OTEZLA (APREMILAST) FOR INHIBITING PDE4

U-1505 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS

U-1506 TREATMENT OF PATIENTS WITH GASTROINTESTINAL STROMAL TUMOR (GIST), INCLUDING BUT NOT LIMITED TO PATIENTS PREVIOUSLY TREATED WITH IMATINIB AND PATIENTS WITH GIST HAVING RESISTANCE TO A KIT TYROSINE KINASE INHIBITOR

U-1507 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN

U-1508 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING A PLURALITY OF COMPOSITE SUBUNITS AS CLAIMED

U-1509 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING A GASTRIC ACID REDUCER

U-1510 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED.

U-1511 TREATMENT OF HYPERTRIGLYCERIDEMIA

U-1512 REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS

U-1513 TREATMENT OF HIV-1 INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS

U-1514 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER BY BUCCAL OR SUBLINGUAL ADMINISTRATION OF FENTANYL

U-1515 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULT PATIENTS.

U-1516 METHOD OF TREATING CHRONIC IDIOPATHIC CONSTIPATION IN ADULT PATIENTS.

U-1517 TREATMENT OF BACTERIAL INFECTIONS USING A TWO-DOSE REGIMEN OF DALBAVANCIN.

U-1518 MAINTAINING PUPIL SIZE BY PREVENTING INTRAOPERATIVE MIOSIS AND REDUCING POSTOPERATIVE OCULAR PAIN

U-1519 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT WITH IRRITABLE BOWEL SYNDROME

U-1520 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1521 MAINTENANCE TREATMENT OF OPIOID DEPENDENCE

U-1522 TREATMENT OF TYPE 2 DIABETES MELLITUS IN A PATIENT, WHEREIN GLYCEMIC CONTROL (HBA1C < 7.0%) IS NOT ACHIEVABLE USING ONE OR MORE OF INSULIN, METFORMIN, PIOGLITAZONE, OR ROSIGLITAZONE

U-1523 METHOD OF INDUCING TOPICAL ANESTHESIA IN THE EYE

U-1524 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE

U-1525 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY

U-1526 THE TREATMENT OF PATIENTS WITH TRAVELERS' DIARRHEA (TD) OR THE REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE

U-1527 FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY

U-1528 A METHOD OF LOWERING INTRAOCULAR PRESSURE

U-1529 ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)

U-1530 USE OF ARIPIRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION

U-1531 METHOD FOR TRANSDERMAL DELIVERY OF TESTOSTERONE

U-1532 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY PATIENTS WITH SODIUM OXYBATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED.

U-1533 PULMONARY ADMINISTRATION OF PARTICLES COMPRISING A DIKETOPIPERAZINE AND INSULIN.

U-1534 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH A DIKETOPIPERAZINE.

U-1535 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH MICROPARTICLES OF A DIKETOPIPERAZINE.

U-1536 ADMINISTRATION OF A COMPOSITION COMPRISING A DIKETOPIPERAZINE AND INSULIN.

U-1537 TREATMENT OF A PATIENT HAVING DIABETES MELLITUS WITH A PRANDIAL RAPID ACTING INSULIN.

U-1538 ADMINISTRATION OF FDKP MICROPARTICLES COMPRISING INSULIN.

U-1539 PULMONARY ADMINISTRATION OF AN INSULIN COMPOSITION COMPRISING FDKP AT THE BEGINNING OF A MEAL TO A PATIENT ALSO BEING TREATED WITH A LONG-ACTING INSULIN.

U-1540 BUTRANS IS A PARTIAL OPIOID AGONIST PRODUCT INDICATED FOR THE MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.

U-1541 TREATMENT OF PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX (TSC) WHO HAVE SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) THAT REQUIRES THERAPEUTIC INTERVENTION BUT CANNOT BE CURATIVELY RESECTED.

U-1542 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA AND/OR NON-HODGKINS LYMPHOMA

U-1543 TREATMENT OF A PATIENT BY ADMINISTERING THE FORMULATION RECITED IN CLAIM 1 OR CLAIM 23

U-1544 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA (PTCL).

U-1545 A METHOD OF TRANSDERMALLY DELIVERING TESTOSTERONE

U-1546 FOR USE IN THE TREATMENT OF MALIGNANT HYPERTHERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTHERMIA IN PATIENTS AT HIGH RISK.

U-1547 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), CHRONIC BRONCHITIS OR EMPHYSEMA

U-1548 FOR THE LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS

U-1549 FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA

U-1550 METHOD OF TREATING METASTATIC PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS.

U-1551 METHOD OF TREATING PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS, IN THE ABSENCE OF INTERFERON ALPHA.

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1552 FOR HEALING OF ALL GRADES OF EROSION ESOPHAGITIS (EE)

U-1553 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN

U-1554 FOR THE TREATMENT OF HEARTBURN ASSOCIATED WITH SYMPTOMATIC NON-EROSIVE GASTROESOPHAGEAL DISEASE (GERD)

U-1555 MANAGEMENT OF MODERATE TO SEVERE PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.

U-1556 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE

U-1557 A METHOD OF TESTOSTERONE REPLACEMENT THERAPY COMPRISING THE STEP OF NASALLY ADMINISTERING TO A PATIENT IN NEED OF SUCH TREATMENT AN EFFECTIVE AMOUNT OF TESTOSTERONE GEL FORMULATION.

U-1558 FOR THE TREATMENT OF PATIENTS WITH RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA OR [RELAPSED] SMALL LYMPHOCYTIC LYMPHOMA

U-1559 INDICATED FOR THE ONCE-DAILY MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN PATIENTS AGED 12 YEARS OF AGE AND OLDER

U-1560 A METHOD OF DISRUPTING LEUKOCYTE FUNCTION, INCLUDING AS AN INHIBITOR OF PI3KDELTA KINASE

U-1561 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS

U-1562 TREATMENT OF PATIENTS WITH HEPATIC ENCEPHALOPATHY (HE)

U-1563 A METHOD OF TRANSDERMAL ADMINISTRATION OF A PHYSIOLOGICALLY ACTIVE AGENT TO A SUBJECT.

U-1564 A METHOD OF TREATING GAUCHER'S DISEASE

U-1565 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE

U-1566 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER

U-1567 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER

U-1568 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE

U-1569 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS

U-1570 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS USING A SINGLE DOSE

U-1571 TREATMENT OF GAUCHER DISEASE TYPE 1

U-1572 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION.

U-1573 USE OF RUXOLITINIB (JAKAFI) FOR INHIBITING JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2.

U-1574 A METHOD OF CATALYZING THE HYDROLYSIS OF GLUCOCEREBROSIDE TO GLUCOSE AND CERAMIDE.

U-1575 PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY

U-1576 TREATMENT OF LEUKEMIA

U-1577 CONTROL OF SERUM PHOSPHOROUS LEVELS

U-1578 TREATMENT OF ACUTE OTITIS MEDIA

U-1579 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER

U-1580 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAD RECEIVED PRIOR DOCETAXEL CHEMOTHERAPY

U-1581 IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA.

U-1582 TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1583 FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY

U-1584 USE OF NALTREXONE AND BUPROPION IN A LAYERED FORMULATION FOR CHRONIC WEIGHT MANAGEMENT FOR AFFECTING WEIGHT LOSS

U-1585 USE OF NALTREXONE AND BUPROPION BASED ON AN ESCALATING DOSE SCHEDULE

U-1586 FOR EFFECT ON BLOOD GLUCOSE PARAMETERS IN PATIENTS WITH INSULIN RESISTANCE

U-1587 SINGLE-DOSE INFILTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA.

U-1588 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC).

U-1589 METHOD OF USE FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA

U-1590 KUVAN IS INDICATED TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA

U-1591 TREATMENT OF ASTHMA IN PATIENTS AGED 12 YEARS AND OLDER

U-1592 TO REDUCE SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE

U-1593 MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA, AND REDUCTION OF EXACERBATIONS IN COPD PATIENTS.

U-1594 DILATION OF THE PUPIL

U-1595 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIASIS

U-1596 LAMICTAL IS AN ANTIEPILEPTIC DRUG (AED) INDICATED FOR: EPILEPSY-ADJUNCTIVE THERAPY IN PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE: (1.1) PARTIAL SEIZURES PRIMARY GENERALIZED TONIC-CLONIC SEIZURES

U-1597 TREATMENT OF DIABETIC MACULAR EDEMA

U-1598 METHOD OF ADMINISTRATION OF CONTROLLED RELEASE OXYMORPHONE

U-1599 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI-INFLAMMATORY, ANALGESIC, AND ANTIPYRETIC ACTIVITY

U-1600 DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS

U-1601 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS

U-1602 METHOD OF ADMINISTERING PIRFENIDONE CAPSULES TO TREAT A FIBROTIC CONDITION

U-1603 METHOD FOR ADMINISTERING PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE

U-1604 METHOD FOR ADMINISTERING PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH A STRONG INHIBITOR OF CYP1A2

U-1605 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING OR AVOIDING ANOTHER STRONG CYP1A2 INDUCER

U-1606 METHOD FOR ADMINISTERING PIRFENIDONE WHILE AVOIDING OR DISCONTINUING CONCOMITANT USE OF A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME INVOLVED IN PIRFENIDONE METABOLISM

U-1607 METHOD OF ADMINISTERING A DOSAGE FORM THAT INCLUDES A GRANULATE FORMULATION OF PIRFENIDONE TO TREAT A FIBROTIC CONDITION

U-1608 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF A FIBROSIS CONDITION

U-1609 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS

U-1610 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN USE OF PIRFENIDONE

U-1611 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING A STRONG CYP1A2 INDUCER

U-1612 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY AVOIDING SMOKING OR BY AVOIDING ANOTHER STRONG CYP1A2 INDUCER

U-1613 DOSAGE MODIFICATION IN TREATMENT WITH PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH CIPROFLOXACIN

U-1614 USE OF TOPICAL DICLOFENAC SODIUM FOR TREATING PAIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1615 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL

U-1616 NASAL ADMINISTRATION OF A TESTOSTERONE GEL TO A PATIENT TO TREAT THE PATIENT FOR A CONDITION ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE

U-1617 METHOD OF TREATING MEDULLARY THYROID CANCER

U-1618 A METHOD OF TREATING A PATIENT SUFFERING FROM A PAIN ASSOCIATED SLEEP DISTURBANCE COMPRISING ADMINISTERING A LIQUID COMPOSITION FORMULATED INSIDE A SOFT GEL CAPSULE, AS CLAIMED, TO THE PATIENT

U-1619 TREATMENT OF IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)

U-1620 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX, WITH A SUBSTANTIALLY NON-IMMUNOGENIC CARBOHYDRATE COMPONENT, IN ABOUT 15 MINUTES OR LESS.

U-1621 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A COMPLEXING AGENT.

U-1622 FOR THE TREATMENT OF POLYCYTHEMIA VERA

U-1623 USE OF EXENATIDE MAY RESULT IN REDUCTION IN APPETITE.

U-1624 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA, ADVANCED RENAL CELL CARCINOMA, OR DIFFERENTIATED THYROID CARCINOMA.

U-1625 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE POOR METABOLIZERS OF CYP2D6

U-1626 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING

U-1627 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN ADULTS

U-1628 METHOD OF TREATING DISORDERS WITH AN ETIOLOGY COMPRISING OR ASSOCIATED WITH EXCESS GH-SECRETION

U-1629 METHOD OF TREATING ACROMEGALY

U-1630 TREATMENT IN COMBINATION WITH A CORTICOID SUCH AS PREDNISONE OF PROSTATE CANCER PREVIOUSLY TREATED WITH DOCETAXEL

U-1631 TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA.

U-1632 TREATMENT OF SCHIZOPHRENIA, WITH EFFICACY IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA

U-1633 USE OF ARIPIPRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA

U-1634 TREATMENT OF BRCA MUTATED OVARIAN CANCER USING PARP INHIBITOR

U-1635 USE OF RITONAVIR AS A POTENT CYP3A INHIBITOR TO INCREASE PLASMA DRUG CONCENTRATION OF PARITAPREXIR AND OVERALL DRUG EXPOSURE FOR TREATMENT OF HCV INFECTION

U-1636 USE OF DASABUVIR TO INHIBIT VIRAL REPLICATION FOR THE TREATMENT OF HCV INFECTION.

U-1637 TREATMENT OF HCV INFECTION USING PARITAPREXIR, OMBITASVIR, RITONAVIR, AND DASABUVIR WITH RIBAVIRIN.

U-1638 TREATMENT OF HCV INFECTION USING PARITAPREXIR

U-1639 USE OF NALTREXONE AND BUPROPION IN EXTENDED-RELEASE FORM FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY

U-1640 TREATMENT OF MODERATE TO SEVERE CHRONIC PAIN BY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED

U-1641 MEMANTINE HCL/DONEPEZIL HCL COMBINATION FOR THE TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE

U-1642 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND FOR WHOM METFORMIN THERAPY IS INAPPROPRIATE BY ADMINISTERING LINAGLIPTIN

U-1643 TREATING CUSHING'S SYNDROME

U-1644 TREATMENT OF OVERACTIVE BLADDER BY APPLICATION OF OXYBUTYRIN CHLORIDE GEL TO SKIN

U-1645 TREATMENT OF PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1646 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

U-1647 TREATMENT OF PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

U-1648 TREATMENT OF PATIENTS WITH PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION

U-1649 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM

U-1650 TREATMENT OF WALDENSTROM'S MACROGLOBULINEMIA

U-1651 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN

U-1652 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN AND METFORMIN

U-1653 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT METFORMIN)

U-1654 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT INSULIN OR A SULFONYLUREA)

U-1655 A METHOD TO ACCELERATE THE TIME TO GASTROINTESTINAL RECOVERY BY ADMINISTERING ABOUT 12 MG OF ALVIMOPAN TO THE PATIENT FROM ABOUT 30 TO 60 MINUTES PRIOR TO SURGERY

U-1656 METHOD OF IRON ADMINISTRATION TO TREAT PATIENTS IN NEED OF IRON REPLACEMENT

U-1657 METHOD FOR PROVIDING POST COITAL CONTRACEPTION TO A WOMAN BY ADMINISTERING ABOUT 30 MG OF ULIPRISTAL ACETATE WITHIN ABOUT 120 HOURS AFTER INTERCOURSE, WHEREIN THE WOMAN IS OVERWEIGHT HAVING A BMI OF 25 TO 29.99

U-1658 TREATMENT OF ER-POSITIVE, HER2-NEGATIVE ADVANCED BREAST CANCER IN COMBINATION WITH LETROZOLE AS INITIAL ENDOCRINE-BASED THERAPY FOR METASTATIC DISEASE IN POSTMENOPAUSAL WOMEN

U-1659 MANAGEMENT OF PAIN

U-1660 TREATMENT OF HIV-1 INFECTION IN ADULTS WITH NO DARUNAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS

U-1661 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS ALSO TAKING LOW DOSE ASPIRIN

U-1662 A METHOD OF TREATING OCULAR PAIN

U-1663 TREATMENT OF HIV-1 INFECTION

U-1664 TREATMENT OF BACTERIAL VAGINOSIS WITH METRONIDAZOLE GEL

U-1665 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER BY ADMINISTERING THE COMPOSITION OF CLAIM 1

U-1666 PALLIATIVE TREATMENT OF PROSTATE CANCER

U-1667 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL ALLERGIC RHINITIS

U-1668 METHOD OF TREATING DEPRESSION OR MAJOR DEPRESSIVE DISORDER

U-1669 TREATMENT OF MULTIPLE MYELOMA, IN COMBINATION WITH BORTEZOMIB AND DEXAMETHASONE

U-1670 NATROBA TOPICAL SUSPENSION IS A PEDICULICIDE INDICATED FOR THE TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS SIX (6) MONTHS OF AGE AND OLDER.

U-1671 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH CONJUNCTIVITIS

U-1672 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTION

U-1673 TREATMENT OF COMPLICATED URINARY TRACT INFECTION, INCLUDING PYELONEPHRITIS

U-1674 DOSAGE MODIFICATION TO REDUCE RISKS ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE

U-1675 USE OF TROKENDI XR FOR THE TREATMENT OF EPILEPSY

U-1676 METHODS FOR TREATING BACTERIAL INFECTIONS

U-1677 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS (IPF)

U-1678 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL

U-1679 TREATMENT OF ACUTE OTITIS EXTERNA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1680 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS

U-1681 TREATMENT OF PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC

U-1682 TREATMENT OF BACTERIAL VAGINOSIS

U-1683 TREATMENT FOR CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION

U-1684 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA

U-1685 DOSAGE MODIFICATION TO REDUCE THE RISK ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE

U-1686 A METHOD TO REDUCE WITHDRAWAL SYMPTOMS, INCLUDING NICOTINE CRAVING, ASSOCIATED WITH SMOKING CESSATION

U-1687 TREATMENT OF HCV INFECTION USING OMBITASVIR

U-1688 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1689 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1690 METHOD FOR REDUCTION OF SUBMENTAL FAT

U-1691 INDICATED FOR THE ONCE-DAILY INHALED TREATMENT FOR ASTHMA IN ADULTS AGED 18 YEARS AND OLDER

U-1692 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1693 METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS

U-1694 A METHOD FOR TREATING HEART FAILURE IN A HUMAN USING A CRYSTALLINE FORM OF IVABRADINE HYDROCHLORIDE

U-1695 METHOD FOR TREATING THYROID CARCINOMA INCLUDING DIFFERENTIATED THYROID CANCER

U-1696 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA

U-1697 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A DIKETOPIPERAZINE.

U-1698 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS

U-1699 A METHOD FOR TREATING ACUTE LYMPHOBLASTIC LEUKEMIA

U-1700 A METHOD FOR TREATING PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA

U-1701 A METHOD FOR TREATING LEUKEMIA RESULTING FROM A MUTATION IN THE BCR-ABL KINASE DOMAIN

U-1702 TREATMENT OF COPD

U-1703 TREATMENT OF RESPIRATORY COMPLAINTS

U-1704 USE FOR TREATMENT IN PATIENTS WITH DIABETES

U-1705 USE FOR TREATMENT IN PATIENTS WITH HYPERGLYCEMIA

U-1706 TREATMENT OF TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE WHEREIN THE COMBINED THERAPEUTIC EFFECT IS GREATER THAN THE ADDITIVE EFFECT OF ADMINISTERING EACH AGENT ALONE

U-1707 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS AND SYMPTOMS THEREOF.

U-1708 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS.

U-1709 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE).

U-1710 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH FLUVOXAMINE

U-1711 FOR THE TREATMENT OF PATIENTS WITH CLL, FL OR SLL

U-1712 MEKINIST IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA

U-1713 TAFINLAR IN COMBINATION WITH TRAMETINIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA

U-1714 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 6 YEARS AND OLDER

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
- U-1715 P2Y12 PLATELET INHIBITOR FOR USE AS ADJUNCT TO PERCUTANEOUS CORONARY INTERVENTION TO REDUCE RISK OF VARIOUS DISEASES/CONDITIONS IN PATIENTS NOT TREATED WITH A P2Y12 PLATELET INHIBITOR AND NOT GIVEN A GLYCOPROTEIN IIB/IIIA INHIBITOR
- U-1716 TREATMENT OF COUGH AND SYMPTOMS ASSOCIATED WITH UPPER RESPIRATORY ALLERGIES OR A COMMON COLD WITH CODEINE PHOSPHATE AND CHLORPHENIRAMINE MALEATE ORALLY ADMINISTERED EXTENDED RELEASE TABLETS
- U-1717 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE
- U-1718 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO HAVE THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE.
- U-1719 ACUTE TREATMENT OF MIGRAINE
- U-1720 METHOD OF PROVIDING A THERAPEUTICALLY EFFECTIVE AND STABLE MEDIAN BLOOD PLASMA LEVEL OF LEVODOPA
- U-1721 USE OF RUXOLITINIB (JAKAFI) FOR BLOCKING SIGNAL TRANSDUCTION OF JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2
- U-1722 TREATMENT OF BASAL CELL CARCINOMA
- U-1723 TREATMENT OF HEART FAILURE
- U-1724 METHOD OF INHIBITING HEPATITIS C VIRUS
- U-1725 METHOD OF INHIBITING HEPATITIS C VIRUS WITH DAKLINZA AND AT LEAST ONE ADDITIONAL COMPOUND HAVING ANTI-HCV ACTIVITY
- U-1726 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH CORONARY HEART DISEASE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1727 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA
- U-1728 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH STABLE NYHA CLASS III HEART FAILURE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1729 REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT)
- U-1730 REDUCE THE RISK OF RECURRENT PULMONARY EMBOLISM
- U-1731 TEMPORARY RELIEF OF MINOR ACHES AND PAINS
- U-1732 TEMPORARY REDUCTION OF FEVER
- U-1733 TREATMENT/PREVENTION OF CARDIOVASCULAR DISEASE
- U-1734 USE OF FLIBANSERIN OR A PHARMACEUTICALLY ACCEPTABLE ACID ADDITION SALT THEREOF TO TREAT HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD)
- U-1735 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER WITH INTRAVENOUS IBUPROFEN SUCH THAT MEAN ARTERIAL BLOOD PRESSURE DOES NOT INCREASE THE DOSAGE INTERVAL
- U-1736 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
- U-1737 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE BEING TREATED WITH FLUOXETINE
- U-1738 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)
- U-1739 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND -THE-CLOCK, LONG-TERM OPIOID TREATMENT, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-1740 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY
- U-1741 PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY
- U-1742 ROLAPITANT IS APPROVED FOR THE PREVENTION OF DELAYED NAUSEA AND VOMITING (I.E., EMESIS) ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1743 FOR THE PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY

U-1744 PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING

U-1745 FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA

U-1746 MONOTHERAPY OR ADJUNCTIVE THERAPY FOR TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH EPILEPSY

U-1747 FOR CLAIMS 1-3,6-13,16-24 AND 26-32: METHOD OF TREATING ADHD

U-1748 FOR CLAIMS 1-4,6-14,16-24 AND 26-32: METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS

U-1749 ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

U-1750 TREATMENT OF SCHIZOPHRENIA AND/OR ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH CARIPRAZINE

U-1751 TREATMENT OF PATIENTS WITH METASTATIC COLORECTAL CANCER WHO HAVE BEEN PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY

U-1752 PROPHYLAXIS OF ORGAN REJECTION

U-1753 TREATMENT OF HCV INFECTION USING DASABUVIR

U-1754 FOR THE TREATMENT OF PULMONARY HYPERTENSION (PAH) IN COMBINATION WITH TADALAFIL

U-1755 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS

U-1756 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF

U-1757 INHIBITION ON PI3K KINASE

U-1758 METHOD OF TREATING ALLERGIC REACTION VIA INJECTION

U-1759 METHOD OF REVERSING THE ANTICOAGULANT EFFECT OF DABIGATRAN USING IDARUCIZUMAB

U-1760 RISK-REDUCTION OF NSAID GASTRIC ULCER IN PATIENTS REQUIRING CHRONIC NSAID TREATMENT

U-1761 PLAQUE PSORIASIS

U-1762 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1763 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1764 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1765 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1766 TREATMENT OF HYPERKALEMIA

U-1767 USE OF CALCIPOTRIENE FOAM FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS AGED 18 YEARS AND OLDER

U-1768 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN

U-1769 TREATMENT OF PAIN BY TRANSMUCOSAL DELIVERY OF BUPRENORPHINE

U-1770 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN NEGATIVE SYMPTOMS AND/OR COGNITIVE DYSFUNCTION OF SCHIZOPHRENIA

U-1771 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION OR 0.9% SODIUM CHLORIDE INJECTION PRIOR TO ADMINISTRATION

U-1772 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN

U-1773 LONG-TERM MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-1774 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1775 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES INCLUDING PSORIASIS

U-1776 METHOD OF USING COBIMETINIB FOR THE TREATMENT OF MELANOMA

U-1777 TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY

U-1778 METHOD FOR TREATING MULTIPLE MYELOMA

U-1779 METHOD FOR TREATING MULTIPLE MYELOMA WITH ONE OR MORE OTHER THERAPEUTIC AGENTS

U-1780 METHOD FOR TREATING CANCER, INCLUDING MULTIPLE MYELOMA

U-1781 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER IN PATIENTS REQUIRING NSAID TREATMENT

U-1782 FOR HEAD LICE INFESTATIONS

U-1783 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM AS CLAIMED

U-1784 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM TRIHYDRATE AS CLAIMED

U-1785 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM FORMULATION AS CLAIMED

U-1786 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS WHILE MANAGING THE RISK OF TERIFLUNOMIDE AND ROSUVASTATIN INTERACTION BY LIMITING THE ROSUVASTATIN DOSE TO NO MORE THAN 10MG AND/OR ADMINISTERING ABOUT HALF THE NORMAL DOSE

U-1787 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY

U-1788 TREATMENT OF PATIENT HAVING DIABETES MELLITUS VIA ORAL INHALATION OF FDKP MICROPARTICLES COMPRISING INSULIN

U-1789 METHOD OF ADMINISTERING AN ETHANOL-FREE TAXANE LIQUID NANODISPERSION FORMULATION TO A SUBJECT COMBINING THE FORMULATION WITH AN AQUEOUS MEDIUM TO PROVIDE AN ETHANOL-FREE TAXANE DILUTED SOLUTION

U-1790 FOR USE IN TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR NON-HODGKIN'S LYMPHOMA

U-1791 EMERGENCY TREATMENT OF ADULT & PEDIATRIC PATIENTS FOLLOWING FLUOROURACIL OR CAPECITABINE OVERDOSE, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING CARDIAC OR CNS TOXICITY OR UNUSUALLY SEVERE ADVERSE REACTIONS WITHIN 96 HOURS

U-1792 TREATMENT OF OTIC INFECTION OR INFLAMMATION

U-1793 TREATMENT OF PEDIATRIC PATIENTS WITH OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT

U-1794 REVERSAL OF DRUG-INDUCED NEUROMUSCULAR BLOCK

U-1795 REVERSAL OF NEUROMUSCULAR BLOCKAGE INDUCED BY ROCURONIUM BROMIDE OR VECURONIUM BROMIDE

U-1796 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA

U-1797 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING SELEXIPAG

U-1798 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING SELEXIPAG IN COMBINATION WITH THE ENDOTHELIN RECEPTOR ANTAGONIST MACITENTAN

U-1799 METHOD OF INCREASING GROWTH OF HAIR INCLUDING EYELASHES

U-1800 A METHOD OF TREATING OCULAR PAIN AND/OR ENHANCING OCULAR COMFORT

U-1801 REDUCTION OF SERUM URIC ACID LEVELS

U-1802 TREATMENT OF GOUT

U-1803 TREATMENT OF HYPERURICEMIA

U-1804 ACHIEVING A THERAPEUTIC BENEFIT IN A SUBJECT WITH GOUT

U-1805 USE OF DEXLANSOPRAZOLE IN PATIENTS TAKING CLOPIDOGREL WITHOUT MEANINGFUL CYP2C19 INTERACTIONS

U-1806 COADMINISTERING WITH ALLOPURINOL TO REDUCE SERUM URIC ACID (SUA) BELOW 4 MG/DL; BELOW 6MG/DL IN PATIENTS HAVING URIC ACID DEPOSITS; AND/OR BELOW 6MG/DL WITH SUA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

	INTRADAY CHANGE MORE THAN 50% AND/OR ADVERSE EVENT RATE LESS THAN 15%
U-1807	TREATMENT OF PEDIATRIC PATIENTS 8 TO 17 YEARS OF AGE WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
U-1808	USE OF NALTREXONE AND BUPROPION FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER
U-1809	METHOD OF DRUG DELIVERY VIA THE NASAL CAVITY
U-1810	TREATMENT OF PAIN IN PATIENTS WITH HEPATIC IMPAIRMENT
U-1811	TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATIONS AFTER CONFIRMING THE PRESENCE OF BRAF V600E MUTATION
U-1812	TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA
U-1813	TREATMENT OF PATIENTS INFECTED WITH HEPATITIS C VIRUS
U-1814	METHOD OF TREATING GLAUCOMA OR ELEVATED INTRAOCULAR PRESSURE
U-1815	TREATMENT OF PARTIAL-ONSET SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY
U-1816	TREATMENT OF A UREA CYCLE DISORDER
U-1817	PEDIATRIC USE AGES 1 MONTH TO 2 YEARS, GERD AND EROSIIVE ESOPHAGITIS
U-1818	TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH LETROZOLE AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN, OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
U-1819	MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE
U-1820	METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 3% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION
U-1821	METHOD FOR CONTRACEPTION TO A WOMAN COMPRISING ADMINISTERING TO THE WOMAN 30MG OF ULIPRISTAL ACETATE MORE THAN 72 HOURS AND UP TO 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
U-1822	TREATMENT OF SCHIZOPHRENIA OR BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA AND/OR BIPOLAR DISORDER
U-1823	A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY COMPENSATING LONG-TERM SENSITIVITY DRIFT OF ELECTROCHEMICAL GAS SENSORS USED IN SYSTEMS FOR DELIVERING THERAPEUTIC NITRIC OXIDE TO A PATIENT
U-1824	A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT
U-1825	METHOD OF USING VISMODEGIB TO TREAT CANCER IN A MAMMAL
U-1826	TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCICLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
U-1827	A METHOD OF PROVIDING A SUBJECT WITH THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET ACCORDING TO CLAIM 1
U-1828	INCREASING MEAN ARTERIAL BLOOD PRESSURE IN ADULT PATIENTS WITH HYPOTENSION ASSOCIATED WITH SEPTIC SHOCK
U-1829	EMERGENCY TREATMENT OF ALLERGIC REACTIONS (TYPE I), INCLUDING ANAPHYLAXIS
U-1830	INDUCTION AND MAINTENANCE OF MYDRIASIS DURING INTRAOCULAR SURGERY
U-1831	METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A CRYSTALLINE FORM OF SELEXIPAG
U-1832	IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE
U-1833	REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
U-1834	TREATMENT OF POSTOPERATIVE INFLAMMATION AND PREVENTION OF OCULAR PAIN IN PATIENTS UNDERGOING CATARACT SURGERY
U-1835	TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION,

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

U-1836 TREATMENT OF HEREDITARY TYROSINEMIA TYPE 1 (HT-1) IN COMBINATION WITH DIETARY RESTRICTION OF TYROSINE AND PHENYLALANINE

U-1837 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN ALONE OR IN COMBINATION WITH INSULIN, METFORMIN, A THIAZOLIDINEDIONE, GLYBURIDE OR METFORMIN PLUS A SULFONYLUREA

U-1838 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN IN COMBINATION WITH METFORMIN

U-1839 COMPOSITION AND METHOD FOR PROVIDING A REDUCTION IN SIDE EFFECTS FOR HUMAN PATIENTS IN NEED OF ACETYLCYSTEINE THERAPY

U-1840 TREATMENT OF HCV INFECTION USING PARITAPREVRIR, OMBITASVIR, RITONAVIR, AND DASABUVIR, WITHOUT RIBAVIRIN

U-1841 USE IN THE LONG-TERM, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-1842 METHOD OF TREATING EPILEPSY

U-1843 TREATMENT OF PSYCHOSIS

U-1844 TREATMENT OF PARKINSON'S DISEASE PSYCHOSIS

U-1845 TREATMENT OF PSYCHOSIS OR A SYMPTOM THEREOF

U-1846 TREATMENT OF A NEURODEGENERATIVE DISEASE OR A SYMPTOM THEREOF

U-1847 METHOD OF TREATING A BACTERIAL INFECTION

U-1848 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN

U-1849 METHOD OF TREATING PULMONARY HYPERTENSION BY ADMINISTERING TREPROSTINIL OR A SALT THEREOF BY INHALATION USING A DEVICE

U-1850 METHOD OF ADMINISTERING LEVETIRACETAM

U-1851 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES

U-1852 METHOD OF TREATING TYPE 2 DIABETES

U-1853 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND, OPTIONALLY, A SULFONYLUREA

U-1854 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC)

U-1855 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS

U-1856 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT HOMOZYGOUS FOR THE UGT1A1*28 ALLELE

U-1857 TO INCREASE BLOOD PRESSURE IN ADULTS WITH VASODILATORY SHOCK (E.G., POST-CARDIOTOMY OR SEPSIS) WHO REMAIN HYPOTENSIVE DESPITE FLUIDS AND CATECHOLAMINES

U-1858 TREATMENT OF PLAQUE PSORIASIS

U-1859 TREATMENT OF SCHIZOPHRENIA, ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER, ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER, AND TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER

U-1860 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION

U-1861 USE OF AN INHALER TO ADMINISTER DRY POWDER MEDICAMENT

U-1862 TREATMENT OF POST-MYOCARDIAL INFARCTION

U-1863 TREATMENT OF STROKE

U-1864 TREATMENT OF MYOCARDIAL INFARCTION

U-1865 TREATMENT OF THROMBOTIC STROKE

U-1866 TREATMENT OF STABLE AND UNSTABLE ANGINA

U-1867 METHOD OF INHIBITING PLATELET AGGREGATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1868 TREATMENT OF ARTERIAL THROMBOTIC COMPLICATIONS SELECTED FROM THE GROUP CONSISTING OF UNSTABLE ANGINA, THROMBOTIC OR EMBOLIC STROKE, TRANSIENT ISCHAEMIC ATTACKS, PERIPHERAL VASCULAR DISEASE AND MYOCARDIAL INFARCTION

U-1869 TREATMENT OF AN ARTERIAL THROMBOTIC COMPLICATION IN A PATIENT WITH CORONARY ARTERY, CEREBROVASCULAR OR PERIPHERAL VASCULAR DISEASE

U-1870 ZINGO IS A POWDER INTRADERMAL SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION

U-1871 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH STAGE 3 OR 4 CHRONIC KIDNEY DISEASE USING CONTROLLED RELEASE, ORAL 25-HYDROXYVITAMIN D

U-1872 USE OF SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN TREATING PATIENTS HAVING 25-HYDROXYVITAMIN D INSUFFICIENCY OR DEFICIENCY

U-1873 ADMINISTRATION OF 25-HYDROXYVITAMIN D3 BY CONTROLLED RELEASE

U-1874 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING OMEPRAZOLE ACCORDING TO CLAIMS 1-8

U-1875 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING S-OMEPRAZOLE TRIHYDRATE ACCORDING TO CLAIMS 1-3

U-1876 METHOD OF ANESTHETIZING AT LEAST A PORTION OF THE MAXILLARY DENTAL ARCH

U-1877 METHOD OF TREATING PULMONARY HYPERTENSION BY ORALLY ADMINISTERING A FORMULATION OF A PHARMACEUTICALLY ACCEPTABLE SALT OF TREPROSTINIL

U-1878 FOR OPIOID DEPENDENCE

U-1879 METHOD OF DIAGNOSING TUMORS USING POSITRON EMISSION TOMOGRAPHY

U-1880 TREATMENT OF SIGNS AND SYMPTOMS OF DRY EYE DISEASE (DED)

U-1881 IMPROVEMENT IN GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR

U-1882 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANALGESIC AND ANTIPYRETIC ACTIVITY

U-1883 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS (GIST)

U-1884 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1885 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1886 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1887 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

U-1888 USE OF CONTROLLED RELEASE 25-HYDROXYVITAMIN D IN TREATING SECONDARY HYPERPARATHYROIDISM IN PATIENTS HAVING CHRONIC KIDNEY DISEASE

U-1889 TREATMENT OF HCV INFECTION USING DASABUVIR/OMBITASVIR/PARITAPREVIR/RITONAVIR FIXED DOSE COMBINATION

U-1890 OTC USE: ALLERGY SYMPTOM RELIEVER; TEMPORARILY RELIEVES THESE SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES; NASAL CONGESTION, RUNNY NOSE, SNEEZING, ITCHY NOSE, AND (ITCHY WATER EYES (AGES 12 AND UP))

U-1891 TREATMENT OR PREVENTION OF NAUSEA AND VOMITING

U-1892 METHOD OF TREATING LEFT VENTRICULAR DYSFUNCTION

U-1893 METHOD OF TREATING MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER IN PEDIATRIC PATIENTS

U-1894 COMBINATION TREATMENT WITH A GLITAZONE FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS

U-1895 METHOD OF TREATING PROSTATE CANCER

U-1896 SUPPLEMENT FOR VITAMIN B12 DEFICIENCIES

U-1897 METHOD OF TREATING ACS USING ANGIOPLASTY WITH AGGRASTAT (TIROFIBAN)

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

HYDROCHLORIDE)

U-1898 METHOD OF INHIBITING PLATELET AGGREGATION WITH AGGRASTAT (TIROFIBAN HYDROCHLORIDE)

U-1899 TREATMENT OF PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN

U-1900 TREATMENT OF THE SIGNS AND SYMPTOMS OF DRY EYE DISEASE (DED)

U-1901 TREATMENT OF SCHIZOAFFECTIVE DISORDER AS A MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS

U-1902 TREATMENT OR SECONDARY PREVENTION OF CARDIOVASCULAR DISEASE, CARDIOVASCULAR EVENTS, OR CEREBROVASCULAR EVENTS AND RISK-REDUCTION OF ASPIRIN-ASSOCIATED GASTRIC ULCERS

U-1903 USE OF NALOXONE HYDROCHLORIDE FOR EMERGENCY TREATMENT OF KNOWN OR SUSPECTED OPIOID OVERDOSE, AS MANIFESTED BY RESPIRATORY AND/OR CENTRAL NERVOUS SYSTEM DEPRESSION.

U-1904 (I) TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY; (II) RESTORING/INCREASING FUNCTIONAL DYSTROPHIN PROTEIN; OR (III) INDUCING SKIPPING; EACH OF (I)-(III) IN PATIENTS HAVING A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING

U-1905 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, THE PATIENT HAVING A R117H MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE

U-1906 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, SUCH AS A PATIENT HAVING A G551D MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE

U-1907 USE OF A DELIVERY DEVICE TO ADMINISTER A DOSE OF NALOXONE

U-1908 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND FORM I LUMACAFTOR

U-1909 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND LUMACAFTOR

U-1910 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING THE DOSAGE UNIT OF CLAIM 1 OF U.S. PATENT NO. 8,716,338

U-1911 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS USING IVACAFTOR AND LUMACAFTOR

U-1912 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING A DOSAGE UNIT AS DEFINED IN CLAIM 1 OF U.S. PATENT NO. 9,192,606

U-1913 TREATMENT OF PEDIATRIC PATIENTS WITH BILATERAL OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT

U-1914 IN COMBINATION WITH RITUXIMAB, FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)

U-1915 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND WHO ARE INELIGIBLE FOR METFORMIN THERAPY BY ADMINISTERING LINAGLIPTIN

U-1916 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY (CINV)

U-1917 TREATMENT OF EXOCRINE PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN

U-1918 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING

U-1919 RESTORING AN MRNA READING FRAME TO INDUCE DYSTROPHIN PROTEIN PRODUCTION IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING

U-1920 USE OF EXTENDED RELEASE ORAL 25-HYDROXYVITAMIN D3 IN TREATING SECONDARY HYPERPARATHYROIDISM IN ADULT PATIENTS HAVING CHRONIC KIDNEY DISEASE STAGE 3 OR STAGE 4

U-1921 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE BY PROVIDING AN ABUSE-DETERRENT ORAL CONTROLLED RELEASE COMBINATION DRUG PRODUCT

U-1922 INTRAVAGINAL PRASTERONE (DEHYDROEPIANDROSTERONE) AT A DAILY DOSE OF 6.5MG FOR THE TREATMENT OF DYSpareunia, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1923 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY BASAL INSULIN OR LIXISENATIDE BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE

U-1924 KYPROLIS IS INDICATED IN COMBINATION WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY

U-1925 USE OF AN AUTO INJECTOR TO ADMINISTER NALOXONE HCL

U-1926 METHOD OF TREATING, REDUCING THE INCIDENCE OF, OR PREVENTING AN ISCHEMIC EVENT IN A PATIENT UNDERGOING PCI BY ADMINISTERING INTRAVENOUSLY 30 UG/KG BOLUS BEFORE PCI AND CONTINUOUS INFUSION OF 4 UG/KG/MIN FOR AT LEAST 2 HOURS OR THE DURATION OF THE PCI

U-1927 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB

U-1928 RUBRACA IS INDICATED AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES.

U-1929 TREATMENT OF DIABETES MELLITUS WITH AN INHALED INSULIN TO IMPROVE GLYCEMIC CONTROL USING A DRY POWDER INHALATION SYSTEM COMPRISING AN INHALER, A CARTRIDGE AND A DRY POWDER MEDICAMENT COMPRISING INSULIN IN A SINGLE INHALATION

U-1930 METHOD OF AEROSOLIZING/DEAGGLOMERATING AN INSULIN DRY POWDER FOR USE IN TREATING DIABETES MELLITUS VIA ORAL INHALATION USING AN INHALER WITH A CARTRIDGE CONTAINING THE INSULIN DRY POWDER.

U-1931 PROPHYLAXIS OR TREATMENT OF VENOUS AND ARTERIAL THROMBOTIC DISEASE

U-1932 METHOD OF TREATING MILD TO MODERATE ATOPIC DERMATITIS.

U-1933 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT SURGERY

U-1934 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH A STRONG CYP1A2 INHIBITOR

U-1935 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH A HISTORY OF MYOCARDIAL INFARCTION

U-1936 TREATMENT OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION

U-1937 TREATMENT OF MYOCARDIAL INFARCTION IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION

U-1938 TREATMENT OF STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION

U-1939 ADMINISTRATION ONCE DAILY WITHIN TWO HOURS AFTER WAKING IN THE MORNING FOR IMPROVEMENT OF GLYCEMIC CONTROL IN A TYPE 2 DIABETES PATIENT

U-1940 IMPROVEMENT IN THE APPEARANCE OF MODERATE TO SEVERE CONVEXITY OR FULLNESS ASSOCIATED WITH SUBMENTAL FAT IN ADULTS BY MEANS OF REDUCING SUBMENTAL FAT VOLUME AS DESCRIBED IN THE APPROVED LABELING

U-1941 TREATMENT OF INFANTILE-ONSET SPINAL MUSCULAR ATROPHY

U-1942 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INCREASING EXON-7 INCLUSION IN SMN2 MRNA

U-1943 TREATMENT OF SPINAL MUSCULAR ATROPHY

U-1944 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INHIBITING AN SMN2 PRE-MRNA INTRONIC SPLICING SILENCER SITE

U-1945 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN.

U-1946 TREATMENT OF SMALL LYMPHOCYTIC LYMPHOMA

U-1947 TREATMENT OF MARGINAL ZONE LYMPHOMA

U-1948 A METHOD FOR TREATING CHRONIC MYELOID LEUKEMIA

U-1949 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)

U-1950 TREATMENT OF PATIENTS WITH ADVANCED (METASTATIC) NON-SMALL CELL LUNG CANCER WHOSE DISEASE PROGRESSED DURING OR AFTER PLATINUM-BASED CHEMOTHERAPY

U-1951 TREATMENT OF ONYCHOMYCOSIS OF A TOENAIL

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1952 FOR USE IN THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA

U-1953 REDUCE THE RISK OF STROKE IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS

U-1954 TREATMENT OF DEEP VEIN THROMBOSIS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS

U-1955 TREATMENT OF PULMONARY EMBOLISM WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS

U-1956 FOLLOWING INITIAL 6 MONTHS TREATMENT FOR DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE), REDUCTION IN THE RISK OF RECURRENCE OF DVT AND OF PE WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS

U-1957 PROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM IN PATIENTS UNDERGOING KNEE OR HIP REPLACEMENT SURGERY, WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS

U-1958 FOR THE TREATMENT OF GENOTYPE 1, 2, 3 OR 4 CHRONIC HEPATITIS C VIRUS (HCV) INFECTION AS A COMPONENT OF A COMBINATION ANTIVIRAL TREATMENT REGIMEN WITH RIBAVIRIN

U-1959 TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM

U-1960 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF SCHIZOPHRENIA IN ADULTS

U-1961 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES (AGES 10 TO ADULT)

U-1962 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: MAINTENANCE MONOTHERAPY IN ADULTS

U-1963 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: AS ADJUNCTIVE TREATMENT TO LITHIUM OR VALPROATE IN ADULTS

U-1964 ELEVATION OF INTRACELLULAR CGMP RESULTING IN INCREASED INTESTINAL FLUID AND ACCELERATED TRANSIT

U-1965 FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) IN COMBINATION WITH TADALAFIL, WHEREIN THE WEIGHT RATIO OF AMBRISENTAN TO TADALAFIL IS ABOUT 1:2 TO ABOUT 1:3

U-1966 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES IN PEDIATRIC PATIENTS AGE 10-17

U-1967 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE THERAPY WITH ONE OR MORE CONVENTIONAL ANTIHYPERGLYCEMIC AGENTS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN

U-1968 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WHO HAVE NOT BEEN PREVIOUSLY TREATED WITH AN ANTIHYPERGLYCEMIC AGENT BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN

U-1969 TOPICAL TREATMENT OF ONYCHOMYCOSIS OF THE TOENAIL(S) DUE TO TRICHOPHYTON RUBRUM AND TRICHOPHYTON MENTAGROPHYTES

U-1970 TREATMENT OF ONYCHOMYCOSIS OF A TOENAIL CAUSED BY TRICHOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES

U-1971 FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA

U-1972 FOR THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA

U-1973 METHOD OF TREATING CYSTIC FIBROSIS USING N-(5-HYDROXY-2,4-DITERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE AND 3-(6-(1-2,2-DIFLUOROBENZO[D][1,3]DIOXOL-5-YL) CYCLOPROPANECARBOXAMIDO)-3-METHYLPYRIDIN-2-YL)BENZOIC ACID

U-1974 TREATMENT OF HALLUCINATIONS AND DELUSIONS ASSOCIATED WITH PARKINSON'S DISEASE PSYCHOSIS

U-1975 METHOD OF INCREASING EYELASH GROWTH WITH BIMATOPROST

U-1976 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO HAVE INADEQUATE CONTROL WITH DAPAGLIFLOZIN

U-1977 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO ARE ALREADY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

TREATED WITH DAPAGLIFLOZIN AND SAXAGLIPTIN

U-1978 TREATMENT OF ADVANCED PROSTATE CANCER WITH A REDUCED LIKELIHOOD OF CAUSING A GONADOTROPHIN RELEASING HORMONE AGONIST SIDE-EFFECT

U-1979 THE TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY

U-1980 A METHOD OF TREATING NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS

U-1981 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER

U-1982 USE OF REVLIMID (LENALIDOMIDE) FOR TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA DUE TO LOW-OR INTERMEDIATE-1-RISK MYELODYSPLASTIC SYNDROMES ASSOCIATED WITH A DELETION 5Q ABNORMALITY WITH OR WITHOUT ADDITIONAL CYTOGENETIC ABNORMALITIES

U-1983 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB

U-1984 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE

U-1985 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)

U-1986 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE, WHEREIN THOSE PATIENTS HAVE NOT RECEIVED PREVIOUS TREATMENT FOR MULTIPLE MYELOMA

U-1987 METHOD OF CONTROLLING GLYCEMIA IN DIABETICS BY ADMINISTERING AN INITIAL DOSE OF INSULIN-FDKP WITH A MEAL; DETERMINING BLOOD GLUCOSE LEVEL 1-2 HRS AFTER AND ADMINISTERING A SUPPLEMENTAL DOSE OF INSULIN-FDKP IF POSTPRANDIAL GLUCOSE LEVEL IS >140 MG/DL

U-1988 METHOD TO TREAT INFANTILE HEMANGIOMA

U-1989 INTRAVITREAL TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)

U-1990 INTRAVITREAL TREATMENT OF DIABETIC MACULAR EDEMA

U-1991 REDUCTION OF MORTALITY IN ACUTE MYOCARDIAL INFARCTION

U-1992 USE OF TROKENDI XR FOR PROPHYLACTIC TREATMENT OF MIGRAINE

U-1993 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING "OFF" EPISODES

U-1994 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) IN ADULTS

U-1995 TREATMENT OF TARDIVE DYSKINESIA

U-1996 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1997 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN AND/OR A PPAR-GAMMA AGONIST AND/OR SULFONYLUREA AND/OR INSULIN

U-1998 TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

U-1999 CHRONIC IDIOPATHIC CONSTIPATION

U-2000 MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS

U-2001 USE FOR THE TREATMENT OF ASTHMA IN PATIENTS 6 YEARS OF AGE AND OLDER

U-2002 USE FOR MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

U-2003 A METHOD OF POSITIONING AN INTRAUTERINE SYSTEM BY HOLDING AN INSERTER HANDLE WITH ONE HAND, ADVANCING THE INSERTER THROUGH THE CERVIX AND INTO THE UTERUS, AND RETRACTING A SLIDER ON THE HANDLE TO RELEASE THE INTRAUTERINE SYSTEM

U-2004 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2005	REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH GENERALIZED TONIC-CLONIC SEIZURES
U-2006	REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH MIXED SEIZURE PATTERNS THAT INCLUDE PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY, GENERALIZED TONIC-CLONIC SEIZURES, OR OTHER PARTIAL OR GENERALIZED SEIZURES
U-2007	TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WHO ARE FLT3 MUTATION-POSITIVE, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION CHEMOTHERAPY
U-2008	TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)
U-2009	METHOD OF TREATING POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE.
U-2010	ACUTE TREATMENT OF MIGRAINE BY DELIVERING A POWDERED SUBSTANCE COMPRISING SUMATRIPTAN VIA A BREATH-POWERED DELIVERY DEVICE
U-2011	TREATMENT OF MIGRAINE VIA DELIVERY OF SUMATRIPTAN VIA THE NASAL CAVITY
U-2012	A METHOD FOR TREATING OVARIAN CANCER BY ADMINISTERING RUCAPARIB, WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA MUTATION
U-2013	TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)
U-2014	A METHOD OF TREATING SECONDARY HYPERPARATHYROIDISM (SHPT)
U-2015	SODIUM THIOSULFATE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM NITRITE FOR THE TREATMENT OF ACUTE CYANIDE POISONING
U-2016	TREATMENT FOR ONYCHOMYCOSIS THAT IS TINEA UNGUIUM
U-2017	TREATMENT OF OPIOID DEPENDENCE
U-2018	MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF
U-2019	METHOD OF DELIVERING TO A PATIENT WITH DIABETES MELLITUS IN A SINGLE INHALATION, GREATER THAN 75% OF A DRY POWDER DOSE COMPRISING INSULIN AND FUMARYL DIKETOPIPERAZINE USING A HIGH RESISTANCE TO FLOW DRY POWDER INHALER.
U-2020	MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
U-2021	METHOD OF ADMINISTERING LEVETIRACETAM UNDER FASTED CONDITIONS
U-2022	METHOD OF ADMINISTERING LEVETIRACETAM UNDER FED CONDITIONS
U-2023	A METHOD OF INCREASING THE BIOAVAILABILITY OF GUAIFENESIN IN A SOLUTION CONTAINING 54% TO 66% BY WEIGHT OF PROPYLENE GLYCOL AND GLYCEROL, WHEREIN THE METHOD INCREASES THE CMAX BY AT LEAST 1.5 AND/OR INCREASES THE AUC (0-INF) BY AT LEAST 1.4
U-2024	METHOD FOR TRANSDERMALLY DELIVERING A DRUG TO A USER IN NEED THEREOF
U-2025	TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER
U-2026	TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.
U-2027	TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST.
U-2028	TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA IN ADULTS
U-2029	PREVENTING CONDITION CHARACTERIZED BY UNDESIRED THROMBOSIS
U-2030	PROPHYLAXIS OF VENOUS THROMBOSIS
U-2031	TAFINLAR IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
U-2032	TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2033 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS
- U-2034 INHIBITING COAGULATION
- U-2035 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM
- U-2036 A METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING PARENTERALLY ADMINISTERING A FORMULATION COMPRISING A) 0.1 TO 5% W/V OF TREPROSTINIL OR A PHARMACEUTICALLY ACCEPTABLE SALT THEREOF AND B) A CITRATE BUFFER
- U-2037 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- U-2038 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- U-2039 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING AN NS5A INHIBITOR
- U-2040 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1A OR 3 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING SOFOSBUVIR WITHOUT AN NS5A INHIBITOR
- U-2041 TREATMENT OF PARTIAL-ONSET SEIZURES
- U-2042 DISCONTINUING ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE
- U-2043 EXTENDED ADJUVANT TREATMENT OF ADULT PATIENTS WITH EARLY STAGE HER2-OVEREXPRESSED/AMPLIFIED BREAST CANCER, TO FOLLOW ADJUVANT TRASTUZUMAB BASE THERAPY
- U-2044 DOSE REDUCTION OF PIRFENIDONE BY ABOUT ONE HALF DURING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TWICE DAILY (1500 MG/DAY) TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER
- U-2045 ADMINISTRATION OF PIRFENIDONE AND AVOIDING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER
- U-2046 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6
- U-2047 ADMINISTERING PIRFENIDONE CONCURRENTLY WITH FLUVOXAMINE, THE PIRFENIDONE AT A DOSE OF ABOUT 801 MG/DAY TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2048 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2049 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE
- U-2050 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2051 DISCONTINUING SMOKING TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2052 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INDUCER TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2053 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2, INCLUDING CIGARETTE SMOKE, TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2054 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2 TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2055 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 LIVER ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2056 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY, FOLLOWING BY ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2057 DOSING 2403 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2058 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400MG/DAY DOSE, FOLLOWED BY ADMINISTERING 2403MG/DAY IN TREATMENT OF IPF
- U-2059 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE IN TREATMENT OF IPF
- U-2060 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN AT LEAST 1600MG/DAY IN TREATMENT OF IPF
- U-2061 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2062 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY DOSE IN TREATMENT OF IPF
- U-2063 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2064 DOSING AT LEAST 1602 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2065 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2066 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE, FOLLOWED BY FULL DAILY DOSE
- U-2067 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2068 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE
- U-2069 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING A SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2070 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN SUB-1600 MG/DAY, THEN AT LEAST 1602 MG/DAY
- U-2071 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2072 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2073 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING FULL DAILY DOSE IN TREATMENT OF IPF
- U-2074 DOSING 1602 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2075 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF
- U-2076 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING 801 MG/DAY FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF
- U-2077 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE THEN FULL DAY DAILY DOSE IN TREATMENT OF IPF

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PATENT USE

- U-2078 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN LIMITS, THEN SUB-2400MG/DAY DOSE, THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2079 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF FIBROSIS AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2080 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF IPF AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2081 DISCONTINUING USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6 AND THEN ADMINISTERING PIRFENIDONE
- U-2082 MODIFYING PIRFENIDONE ADMINISTRATION FROM A DOSE OF ABOUT 2400 MG/DAY DOWNWARD BY ABOUT 1600 MG/DAY WHILE CO-ADMINISTERING FLUVOXAMINE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2083 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY 801 MG/DAY, DOSE, THEN 1602 MG/DAY IN TREATMENT OF IPF
- U-2084 TREATMENT OF SEVERE CHRONIC PAIN VIA INTRATHECAL INFUSION OF ZICONOTIDE IN PATIENTS ALSO RECEIVING MORPHINE
- U-2085 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH RIFAMPIN
- U-2086 A METHOD FOR ADMINISTERING ESTRADIOL COMPRISING A MONOLITHIC TRANSDERMAL DRUG DELIVERY SYSTEM CONSISTING OF (I) A BACKING LAYER AND (II) A SINGLE ADHESIVE POLYMER MATRIX LAYER AS CLAIMED IN US PATENT NO. 9730900
- U-2087 TREATMENT OF RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION
- U-2088 TREATMENT OF PARTIAL-ONSET SEIZURES WITH OR WITHOUT SECONDARILY GENERALIZED SEIZURES IN PATIENTS WITH EPILEPSY
- U-2089 TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY
- U-2090 FOR THE TREATMENT OF ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- U-2091 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT NOT HOMOZYGOUS FOR THE UGT1A1*28 ALLELE
- U-2092 METHOD FOR CONFIRMING DOSE DELIVERY
- U-2093 TREATMENT OF TYPE II SPINAL MUSCULAR ATROPHY
- U-2094 TREATMENT OF TYPE III SPINAL MUSCULAR ATROPHY
- U-2095 MITOSOL IS AN ANTIMETABOLITE INDICATED AS AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY. IT IS INTENDED FOR TOPICAL APPLICATION TO THE SITE OF GLAUCOMA FILTRATION SURGERY
- U-2096 SOTYLIZE IS INDICATED FOR THE MAINTENANCE OF NORMAL SINUS RHYTHM [DELAY IN TIME TO RECURRENCE OF ATRIAL FIBRILLATION/ATRIAL FLUTTER (AFIB/AFL)] IN PATIENTS WITH SYMPTOMATIC AFIB/AFL WHO ARE CURRENTLY IN SINUS RHYTHM
- U-2097 TREATMENT OF DMD IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2098 INCREASING PRODUCTION OF FUNCTIONAL DYSTROPHIN PROTEIN IN DMD PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2099 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING BRONCHITIS AND/OR EMPHYSEMA
- U-2100 INDICATED FOR THE ONCE-DAILY TREATMENT OF ASTHMA IN PATIENTS 18 YEARS AND OLDER
- U-2101 MAINTENANCE TREATMENT OF RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY

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PATENT USE

- U-2102 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY BASED ON AN FDA-APPROVED COMPANION DIAGNOSTIC FOR LYNPARZA
- U-2103 MAINTENANCE TREATMENT OF BRCA-MUTATED RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-2104 TREATMENT OF HYPERURICEMIA ASSOCIATED WITH GOUT IN PATIENTS WHO HAVE NOT ACHIEVED TARGET SERUM URIC ACID LEVELS WITH A MEDICALLY APPROPRIATE DAILY DOSE OF ALLOPURINOL ALONE
- U-2105 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING IMMEDIATE RELEASE LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2106 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2107 TREATMENT OF LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA REFRACTORY TO RADIOACTIVE IODINE TREATMENT
- U-2108 TREATMENT OF HORMONE RECEPTOR POSITIVE HER2-NEGATIVE ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY
- U-2109 CAROSPIR IS INDICATED FOR TREATMENT OF NYHA CLASS III-IV HEART FAILURE AND REDUCED EJECTION FRACTION TO INCREASE SURVIVAL, MANAGE EDEMA, AND TO REDUCE THE NEED FOR HOSPITALIZATION FOR HEART FAILURE
- U-2110 METHOD FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS WITH MODERATE RENAL IMPAIRMENT WHO ARE OBESE, OR OVERWEIGHT AND HAVE AT LEAST ONE WEIGHT RELATED COMORBID CONDITION
- U-2111 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1-5 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1-5
- U-2112 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 6 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 6
- U-2113 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 7
- U-2114 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 9 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 9
- U-2115 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 10 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 10
- U-2116 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 12 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 12
- U-2117 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 14-15 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 14-15
- U-2118 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 16-18 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 16-18
- U-2119 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 19
- U-2120 TREATMENT OF PATIENTS 18 YEARS OF AGE AND OLDER WITH COMPLICATED URINARY TRACT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

INFECTIONS CAUSED BY SUSCEPTIBLE MICROORGANISMS

- U-2121 TREATMENT OF PARTIAL-ONSET SEIZURES IN A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ABSENCE SEIZURES
- U-2122 USE FOR REDUCING EXACERBATIONS OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2123 TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY WHO HAVE BEEN PREVIOUSLY TREATED WITH OXCARBAZEPINE
- U-2124 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- U-2125 THE TREATMENT OF AN INFLAMMATORY DISORDER OF THE RESPIRATORY TRACT BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE AND A LONG-ACTING BETA2 ADRENORECEPTOR AGONIST
- U-2126 USE OF FLUTICASONE FUROATE FOR THE TREATMENT OF AN INFLAMMATORY OR ALLERGIC CONDITIONS, INCLUDING CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2127 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2128 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHOLINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLIDINIUM, VIA INHALATION
- U-2129 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHOLINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLIDINIUM, VIA TOPICAL APPLICATION
- U-2130 TREATMENT OF PARTIAL ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY
- U-2131 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH COMPARABLE EFFICACY, AND A REDUCTION IN SPECIFIED ADVERSE EVENTS, COMPARED TO BRIMONIDINE 0.2% TID
- U-2132 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY
- U-2133 METHOD OF DELIVERING FLUTICASONE PROPIONATE TO A NASAL AIRWAY
- U-2134 THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE AND A LONG-ACTING BETA2 ADRENORECEPTOR
- U-2135 AS MONOTHERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY AND PRIOR CHEMOTHERAPY IN THE METASTATIC SETTING
- U-2136 TREATMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-2137 TREATMENT OF POSTHERPETIC NEURALGIA
- U-2138 TOPICAL TREATMENT OF ACTINIC KERATOSIS OF THE FACE OR SCALP USING MORE THAN ONE TREATMENT COURSE OF INGENOL MEBUTATE
- U-2139 TREATMENT OF TYPE 2 DIABETES MELLITUS IN COMBINATION WITH EXENATIDE
- U-2140 METHOD OF TREATING PARTIAL ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2141 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6
- U-2142 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR RECURRENT DVT AND/OR AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
- U-2143 AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS, TO REDUCE THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS AND/OR PULMONARY EMBOLISM IN CERTAIN PATIENTS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS
- U-2144 REDUCTION OF INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-2145 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-2146 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A ROTATING DRIVE SLEEVE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2147 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ORALLY ADMINISTERING 20MG OF TASIMELTEON ONCE DAILY BEFORE BEDTIME

U-2148 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY MEASURING AND DISPLAYING AN INDICATION OF THE CALCULATED DELIVERY CONCENTRATION OF NITRIC OXIDE AS COMPARED TO THE DESIRED DELIVERY CONCENTRATION OF NITRIC OXIDE

U-2149 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON

U-2150 TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE

U-2151 METHOD OF TREATING PAIN OR INFLAMMATION WITH AN INJECTABLE CONTROLLED OR SUSTAINED RELEASE FORMULATION OF TRIAMCINOLONE ACETONIDE

U-2152 TREATMENT OF PAIN ASSOCIATED WITH IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)

U-2153 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2154 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2155 REDUCING BODY WEIGHT IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2156 REDUCING HBA1C IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4

U-2157 TREATING TYPE 2 DIABETES MELLITUS BY STIMULATING INSULIN RELEASE

U-2158 DECREASING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY USING A SUSTAINED-RELEASE COMPOSITION

U-2159 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA

U-2160 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 5 MG OF MELOXICAM

U-2161 TREATMENT OF NAUSEA AND VOMITING, INCLUDING THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY OR MODERATELY EMETOGENIC CANCER CHEMOTHERAPY

U-2162 FOR CLEANSING THE LARGE INTESTINE AS A PREPARATION FOR COLONOSCOPY

U-2163 TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCICLIB OR ABEMACICLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

U-2164 ZELBORAF IS INDICATED FOR THE TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION

U-2165 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 10 MG OF MELOXICAM

U-2166 TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

U-2167 METHOD OF USING A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER

U-2168 METHOD OF USING A LOGIC CIRCUIT TO STABILIZE BATTERY VOLTAGE SUPPLIED TO A SENSOR EMBEDDED WITH A TABLET AND THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER

U-2169 METHOD OF USING A RECEIVER TO IDENTIFY A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT

U-2170 METHOD OF USING A RECEIVER TO RECEIVE A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT

U-2171 ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RCC FOLLOWING NEPHRECTOMY

U-2172 METHOD TO TREAT SEVERE ALLERGIC EMERGENCIES IN PATIENTS WEIGHING 7.5 TO 15 KG (16.5 TO 33 LBS)

U-2173 TREATING OPIOID DEPENDENCE BY ADMINISTERING BUPRENORPHINE

U-2174 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH

U-2175 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE MONTHLY

U-2176 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE

U-2177 TREATING OPIOID ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE

U-2178 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE COMPOSITION WITH 28 DAY DOSE DURATION

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U-2179 IN SITU FORMATION OF SOLID BUPRENORPHINE COMPOSITION

U-2180 TREATING ADDICTION WITH 100 MG OR 300 MG DOSE OF BUPRENORPHINE

U-2181 TREATING OPIOID DEPENDENCY BY SUBCUTANEOUSLY ADMINISTERING BUPRENORPHINE

U-2182 IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS

U-2183 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 13

U-2184 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 13, AND 14

U-2185 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15 AND 27

U-2186 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15, 27, AND 28

U-2187 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29 AND 39

U-2188 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29, 39, AND 40

U-2189 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41 AND 52

U-2190 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41, 52, AND 53

U-2191 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54 AND 64

U-2192 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54, 64, AND 65

U-2193 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66 AND 75

U-2194 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66, 75, AND 76

U-2195 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77 AND 87

U-2196 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77, 87, AND 88

U-2197 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89 AND 99

U-2198 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89, 99, AND 100

U-2199 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA

U-2200 COMBINATION TREATMENT WITH INSULIN GLARGINE WITH OR WITHOUT METFORMIN FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS

U-2201 TREATMENT OF BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN BIPOLAR DISORDER

U-2202 OZEMPIC IS INDICATED AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC

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CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

- U-2203 A METHOD OF PROVIDING A SUBJECT WITH A THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET AS CLAIMED
- U-2204 TREATING PATIENTS WITH ACUTE PROMYELOCYTIC LEUKEMIA (APL) WHO ARE REFRACTORY TO, OR HAVE RELAPSED FROM, RETINOID AND ANTHRACYCLINE CHEMOTHERAPY, AND WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- U-2205 TREATMENT OF SEBORRHEIC KERATOSES THAT ARE RAISED
- U-2206 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE
- U-2207 TREATING ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE
- U-2208 TREATING ADDICTION BY ONCE PER MONTH ADMINISTRATION OF BUPRENORPHINE
- U-2209 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH
- U-2210 TREATING OPIOID ADDICTION BY 100 MG OR 300 MG DOSE BUPRENORPHINE
- U-2211 TREATING OPIOID ADDICTION BY ADMINISTRATION OF BUPRENORPHINE
- U-2212 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2213 REDUCING HBA1C IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2214 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES
- U-2215 ERTUGLIFLOZIN IN COMBINATION WITH SITAGLIPTIN AND IN FURTHER COMBINATION WITH METFORMIN AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2216 ERTUGLIFLOZIN AND SITAGLIPTIN IN COMBINATION AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2217 TREATING HIGH OUTPUT SHOCK WITH ANGIOTENSIN II BY INCREASING MEAN ARTERIAL PRESSURE IN PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE
- U-2218 MAINTAINING MEAN ARTERIAL PRESSURE OF ABOUT 65 MMHG OR HIGHER WITH ANGIOTENSIN II IN SHOCK PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE
- U-2219 TREATMENT OF CHRONIC SMALL LYMPHOCYTIC LEUKEMIA
- U-2220 A METHOD FOR THE DIAGNOSIS OF ADULT GROWTH HORMONE DEFICIENCY BY MEASURING THE LEVEL OF GROWTH HORMONE AFTER ORAL ADMINISTRATION OF MACIMORELIN
- U-2221 TREATING REFRACTORY HYPOTENSION WITH ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A PATIENT RECEIVING VASOPRESSOR
- U-2222 RELIEVES REDNESS OF THE EYE DUE TO MINOR EYE IRRITATIONS
- U-2223 METHOD OF TREATING ANGINA PECTORIS
- U-2224 TREATMENT OF DYSKINESIA AND INCREASING ON TIME WITHOUT TROUBLESOME DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2225 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN PATIENTS WITH HEPATIC IMPAIRMENT
- U-2226 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN PATIENTS WITH RENAL IMPAIRMENT
- U-2227 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN GERIATRIC PATIENTS
- U-2228 TREATMENT OF SMALL LYMPHOCYTIC LEUKEMIA
- U-2229 IN COMBINATION WITH TRETINOIN, TREATING ADULTS AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH NEWLY-DIAGNOSED LOW-RISK ACUTE PROMYELOCYTIC LEUKEMIA (APL) CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-A GENE EXPRESSION
- U-2230 IRRITABLE BOWEL SYNDROME WITH CONSTIPATION
- U-2231 TREATING REFRACTORY HYPOTENSION WITH ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A PATIENT RECEIVING VASOPRESSOR
- U-2232 TREATMENT OF PSORIATIC ARTHRITIS USING A DOSAGE TITRATION SCHEDULE

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U-2233	TREATMENT OF PSORIATIC ARTHRITIS WITH APREMILAST USING A DOSAGE TITRATION SCHEDULE AND A SECOND ACTIVE AGENT
U-2234	USE OF IVACAFTOR FOR TREATING CYSTIC FIBROSIS IN A PATIENT WITH A MILD TO MODERATE CF PHENOTYPE WITH AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
U-2235	USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC HIGH-RISK CASTRATION-SENSITIVE PROSTATE CANCER
U-2236	REDUCING THE RISK OF PRETERM BIRTH IN WOMEN WITH A SINGLETON PREGNANCY WHO HAVE A HISTORY OF SINGLETON SPONTANEOUS PRETERM BIRTH
U-2237	TREATMENT OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)
U-2238	METHOD OF IMPROVING GLYCEMIC CONTROL IN PATIENTS WITH DIABETES MELLITUS BY ADMINISTERING A MIXTURE OF INSULIN DEGLUDEC AND INSULIN ASPART DURING OR AROUND THE TIME OF THE LARGEST MEAL OF THE DAY
U-2239	REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH A REDUCTION IN SPECIFIED ADVERSE EVENTS, COMPARED TO BRIMONIDINE 0.2% TID
U-2240	REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH COMPARABLE EFFICACY TO BRIMONIDINE 0.2% TID
U-2241	TREATMENT OF SMALL LYMPHOCYTIC LYMPHOMA WITH 17P DELETION
U-2242	TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION
U-2243	TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA WITH 17P DELETION
U-2244	A METHOD OF TREATING BACTERIAL INFECTIONS IN HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) PATIENTS COMPRISING ADMINISTERING A BACTERICIDALLY EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
U-2245	A METHOD OF TREATING A BACTERIAL INFECTION IN HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) PATIENTS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
U-2246	TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVING AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
U-2247	TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF PATIENTS WITH A MILD TO MODERATE CLINICAL PHENOTYPE OF CYSTIC FIBROSIS HAVING AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
U-2248	TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR THE F508DEL MUTATION AND A SECOND MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR
U-2249	MANAGEMENT OF ACUTE PAIN SEVERE ENOUGH TO REQUIRE AN OPIOID ANALGESIC AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
U-2250	DETECTION OF CARCINOMA IN THE BLADDER BY PHOTODYNAMIC CYSTOSCOPY
U-2251	IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
U-2252	THE TREATMENT OF ACUTE OTITIS EXTERNA IN PATIENTS 6 MONTHS OF AGE AND OLDER DUE TO PSEUDOMONAS AERUGINOSA AND STAPHYLOCOCCUS AUREUS
U-2253	PROPHYLACTIC TREATMENT OF NAUSEA AND VOMITING, INCLUDING PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED CHEMOTHERAPY
U-2254	USE OF POMALIDOMIDE WITH DEXAMETHASONE FOR PATIENTS WITH MULTIPLE MYELOMA AFTER AT LEAST TWO PRIOR THERAPIES INCLUDING LENALIDOMIDE AND A PROTEASOME INHIBITOR AND DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETING THE LAST THERAPY
U-2255	TREATING SECONDARY HYPERPARATHYROIDISM IN CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND THE SUSTAINED RELEASE IS OVER AT LEAST 10 HOURS
U-2256	TREATING SECONDARY HYPERPARATHYROIDISM IN CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE

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PATENT USE

- LEVEL AND CMAX IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2257 TREATING SHPT IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE SERUM PARATHYROID HORMONE LEVEL AND CHANGE IN SERUM CONCENTRATION OF CALCIFEDIOL IN DOSE INTERVAL IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2258 TREATING SECONDARY HYPERPARATHYROIDISM IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND CMAX24HR/C24HR IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2259 TREATING SECONDARY HYPERPARATHYROIDISM IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND TMAX IS INCREASED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2260 METHOD OF REDUCING THE RISK OF PERIPROCEDURAL MYOCARDIAL INFARCTION, AND STENT THROMBOSIS IN A PATIENT UNDERGOING PCI BY ADMINISTERING INTRAVENOUSLY 30 UG/KG BOLUS BEFORE PCI AND THEN A CONTINUOUS INFUSION
- U-2261 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MILD TO MODERATE PAIN OR MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS
- U-2262 MODIFIED DOSING REGIMEN FOR THE REDUCTION OF FEVER
- U-2263 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MODERATE TO SEVERE PAIN WITH ADJUNCTIVE OPIOID ANALGESICS
- U-2264 METHODS OF TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ARTERIOSIS WITH AQUEOUS COMPOSITION
- U-2265 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HEC AND MEC IN ADULT AND PEDIATRIC PATIENTS
- U-2266 METHODS OF MAKING AQUEOUS COMPOSITION AND TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ARTERIOSIS WITH AQUEOUS COMPOSITION
- U-2267 METHOD FOR RELIEVING THE PAIN ASSOCIATED WITH POST-HERPETIC NEURALGIA
- U-2268 DISCONTINUING A STRONG CYP1A2 INDUCER TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2269 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400MG/DAY DOSE THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2270 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, THEN SUB-2400MG/DAY DOSE, THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2271 THERAPEUTIC TREATMENT OF PATIENTS WITH CASTRATION-RESISTANT PROSTATE CANCER, SYMPTOMATIC BONE METASTASES AND NO KNOWN VISCERAL METASTATIC DISEASE
- U-2272 TREATMENT OF NASAL POLYPS IN PATIENTS ≥ 18 YEARS OF AGE WHO HAVE HAD ETHMOID SINUS SURGERY USING A CORTICOSTEROID-ELUTING (MOMETASONE FUROATE) IMPLANT
- U-2273 A METHOD FOR TREATING EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER, WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA MUTATION
- U-2274 MAINTAINING SERUM 25-HYDROXYVITAMIN D AT A LEVEL OF AT LEAST 30 NG/ML WITH ORAL, SUSTAINED RELEASE 25-HYDROXYVITAMIN D
- U-2275 TREATING CYSTIC FIBROSIS PATIENTS AGES 12 AND OLDER, WHO ARE HOMOZYGOUS FOR F508DEL OR HAVE AT LEAST 1 CFTR GENE MUTATION RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS (<30% CRYSTALLINE) IVACAFTOR
- U-2276 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT AGE 6 OR OLDER HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2277 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE
- U-2278 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE IN COMBINATION WITH METFORMIN
- U-2279 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE IN COMBINATION WITH METFORMIN AND A SECOND ORAL ANTIDIABETIC DRUG

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PATENT USE

U-2280 ADJUNCTIVE TREATMENT OF PATIENTS WITH TSC-ASSOCIATED PARTIAL-ONSET SEIZURES

U-2281 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1

U-2282 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 2 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 2

U-2283 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 3-7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 3-7

U-2284 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 8 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 8

U-2285 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 11 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 11

U-2286 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 14 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 14

U-2287 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 16-19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 16-19

U-2288 TREATMENT OF TYPE 2 DIABETES MELLITUS WITH EXENATIDE AS AN ADD-ON TO BASIL INSULIN OR BASAL INSULIN PLUS METFORMIN THERAPY

U-2289 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21L858R MUTATIONS

U-2290 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (45 ML/MIN/1.73 M² ≤ EGFR < 60 ML/MIN/1.73 M²) BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN

U-2291 REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS IN PATIENTS WITH A HISTORY OF MYOCARDIAL INFARCTION (MI) OR WITH PERIPHERAL ARTERIAL DISEASE (PAD)

U-2292 METHOD OF REDUCING THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND CARDIOVASCULAR DISEASE BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN

U-2293 USE IN COMBINATION WITH DEXAMETHASONE IN ADULTS FOR THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF CANCER CHEMOTHERAPY, INCLUDING, BUT NOT LIMITED TO, HIGHLY EMETOGENIC CHEMOTHERAPY

U-2294 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA (ITP) WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT

U-2295 TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER

U-2296 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION

U-2297 IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES PATIENTS BY ADMINISTERING A STARTING DOSE OF 10 MCG FOR 14 DAYS AND INCREASING TO A MAINTENANCE DOSE OF 20 MCG ON DAY 15

U-2298 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS

U-2299 TAFINLAR IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION

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- U-2300 USE IN COMBINATION WITH THE MUSCARINIC ANTAGONIST SOLIFENACIN SUCCINATE FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-2301 USE IN COMBINATION WITH DEXAMETHASONE IN ADULTS FOR THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY EMETOGENIC CANCER CHEMOTHERAPY
- U-2302 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-2303 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-2304 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION
- U-2305 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- U-2306 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM
- U-2307 TREATMENT OF AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE
- U-2308 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- U-2309 USE IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER TO IMPROVE PROCESSING SPEED, AN ASPECT OF COGNITIVE FUNCTION
- U-2310 FOR CLEANSING OF THE COLON IN PREPARATION FOR COLONOSCOPY IN ADULTS
- U-2311 TREATMENT OF HYPERURICEMIA ASSOCIATED WITH GOUT IN PATIENTS WHO HAVE NOT ACHIEVED TARGET SERUM URIC ACID LEVELS WITH A XANTHINE OXIDASE INHIBITOR ALONE
- U-2312 TREATMENT OF HYPERKALEMIA IN ADULTS
- U-2313 METHOD OF REDUCING THE RISK OF CARDIOVASCULAR DEATH, NON-FATAL MYOCARDIAL INFARCTION, AND/OR NON-FATAL STROKE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE BY ADMINISTERING LIRAGLUTIDE
- U-2314 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC LIVER DISEASE WHO IS SCHEDULED TO UNDERGO A PROCEDURE USING DOPTelet
- U-2315 TREATMENT OF MULTIPLE SCLEROSIS IN THE PEDIATRIC PATIENT POPULATION WITH 0.25 MG FINGOLIMOD
- U-2316 TREATMENT OF DYSPAREUNIA
- U-2317 TREATMENT OF A SYMPTOM OF VULVAR AND VAGINAL ATROPHY
- U-2318 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2319 KYPROLIS IS INDICATED IN COMBINATION WITH DEXAMETHASONE OR WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-2320 KYPROLIS IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE OR MORE LINES OF THERAPY
- U-2321 A METHOD OF APPLYING TRYPAN BLUE ONTO AN OUTER SURFACE OF THE ANTERIOR LENS CAPSULE TO FACILITATE REMOVAL OF THE LENS SUBSTANCE
- U-2322 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC)
- U-2323 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-2324 FOR SECONDARY PREVENTION OF CARDIOVASCULAR AND CEREBROVASCULAR EVENTS IN PATIENTS AT RISK OF DEVELOPING ASPIRIN-ASSOCIATED GASTRIC ULCERS
- U-2325 EMERGENCY TREATMENT OF ALLERGIC REACTIONS (TYPE 1), INCLUDING ANAPHYLAXIS; A

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- METHOD OF TREATING ALLERGIC REACTION, ANAPHYLAXIS, ANAPHYLACTIC SHOCK, OR COMBINATION THEREOF BY AN INJECTION OF AT LEAST ONE DOSAGE OF THE INJECTABLE LIQUID PHARMACEUTICAL
- U-2326 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS
- U-2327 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS, COMPRISING MONITORING A PATIENT'S SERUM SODIUM CONCENTRATION
- U-2328 METHOD OF USING PLAZOMICIN TO TREAT BACTERIAL INFECTIONS
- U-2329 METHOD OF ADMINISTERING A LOCAL ANESTHETIC PRIOR TO PERFORMING A DIAGNOSTIC OR SURGICAL PROCEDURE ON A SUBJECT WITH HEPATIC OR RENAL IMPAIRMENT
- U-2330 METHOD OF TREATING MELANOMA
- U-2331 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA
- U-2332 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA MEDIATED BY A B-RAF PROTEIN KINASE
- U-2333 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA WITH A BRAF MUTATION
- U-2334 TREATMENT OF MELANOMA WITH A BRAF MUTATION
- U-2335 TREATMENT OF MELANOMA
- U-2336 TREATMENT OF MELANOMA MEDIATED BY A B-RAF PROTEIN KINASE
- U-2337 INDICATED IN COMBINATION WITH BINIMETINIB FOR THE TREATMENT OF MELANOMA WITH A BRAF MUTATION
- U-2338 MAINTAINING MEAN ARTERIAL PRESSURE OF ABOUT 65 MMHG OR ABOVE WITH ABOUT 1 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN HYPOTENSIVE PATIENTS TREATED WITH VASOPRESSIN OR A VASOPRESSIN ANALOGUE AND REDUCING VASOPRESSIN OR VASOPRESSIN ANALOGUE USE
- U-2339 USE OF A PHARMACEUTICAL COMPOSITION COMPRISING LINAGLIPTIN, METFORMIN AND A BASIC AMINO ACID TO TREAT TYPE 2 DIABETES MELLITUS
- U-2340 TREATMENT OF POSTOPERATIVE INFLAMMATION
- U-2341 METHOD OF RECONSTITUTING A LYOPHILIZED LIPOSOMAL COMPOSITION FOR ADMINISTERING CYTARABINE AND DAUNORUBICIN TO TREAT ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- U-2342 METHOD OF ADMINISTERING A RECONSTITUTED LIPOSOMAL COMPOSITION CONTAINING CYTARABINE AND DAUNORUBICIN TO TREAT ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- U-2343 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR F508DEL AND A SECOND CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2344 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC LIVER DISEASE WHO ARE SCHEDULED TO UNDERGO A PROCEDURE
- U-2345 TREATMENT OF PATIENTS WITH CASTRATION-RESISTANT PROSTATE CANCER (CRPC)
- U-2346 TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 13 KG
- U-2347 TREATMENT OF TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT AND FOR WHOM METFORMIN THERAPY IS INAPPROPRIATE BY ADMINISTERING LINAGLIPTIN WITHOUT DOSE ADJUSTMENT
- U-2348 A METHOD FOR PREVENTION OF PREGNANCY
- U-2349 FOR ONCE-DAILY MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN PATIENTS AGED 5 YEARS AND OLDER
- U-2350 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS ACUTE MYELOGENOUS LEUKEMIA (AML)
- U-2351 TREATMENT OF ACUTE MYELOID LEUKEMIA (AML) WITH AN IDH1 MUTATION
- U-2352 TREATMENT OF HIV-1 INFECTION IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS

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- U-2353 TX OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASES IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TX HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2354 COMBINATION WITH OTHER ANTIRETROVIRALS (ATV) FOR TREATMENT OF HIV-1 IN ATV TREATMENT-EXPERIENCED PATIENTS 2 YEARS AND OLDER WITH EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITOR AND OTHER ATV
- U-2355 IN COMBINATION WITH AN AROMATASE INHIBITOR FOR THE TREATMENT OF PRE/PERIMENOPAUSAL OR POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE-BASED THERAPY
- U-2356 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- U-2357 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- U-2358 TREATMENT OF PATIENTS WITH HORMONE RECEPTOR (HR)-NEGATIVE BREAST CANCER WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA, HER2-NEGATIVE METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING
- U-2359 TREATMENT OF PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE BREAST CANCER WHO SHOULD HAVE BEEN TREATED WITH PRIOR ENDOCRINE THERAPY OR BE CONSIDERED INAPPROPRIATE FOR ENDOCRINE THERAPY
- U-2360 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS
- U-2361 METHOD OF ADMINISTERING A GRANULATE FORMULATION OF 5-METHYL-1-PHENYL-2-(1H)-PYRIDONE AS RECITED IN CLAIM 1, TO TREAT IDIOPATHIC PULMONARY FIBROSIS
- U-2362 TREATMENT OF HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6
- U-2363 ADMINISTRATION OF RISPERIDONE
- U-2364 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY
- U-2365 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE IN ADULTS WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2366 TREATMENT OF LIVER DISEASE THROUGH NUTRITION FOR PATIENTS UNDER THE AGE OF 12
- U-2367 USE FOR PATIENTS WITH PARENTERAL NUTRITION ASSOCIATED CHOLESTASIS OR PARENTERAL NUTRITION ASSOCIATED LIVER DISEASE
- U-2368 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 9 YEARS OF AGE AND OLDER
- U-2369 FOR THE TREATMENT OF GENOTYPE 1, 4, 5 OR 6 CHRONIC HEPATITIS C VIRUS (HCV) INFECTION
- U-2370 FOR TREATMENT-NAIVE GENOTYPE 1 PATIENTS WITH CHRONIC HEPATITIS C VIRUS (HCV) INFECTION FOR A DURATION OF 8-WEEKS
- U-2371 THE TREATMENT OF FABRY PATIENTS
- U-2372 A METHOD OF REDUCING LEFT VENTRICULAR MASS INDEX (LVMI) IN A FABRY PATIENT BY ADMINISTERING MIGALASTAT
- U-2373 A METHOD OF REDUCING PODOCYTE GLOBOTRIAOSYL CERAMIDE (GL-3) IN A FABRY PATIENT BY ADMINISTERING MIGALASTAT
- U-2374 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND IVACAFTOR
- U-2375 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR FORM I AND IVACAFTOR
- U-2376 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-2377 USE OF VITAL DYE FOR FACILITATING SURGICAL PROCEDURES FOR VITREO-RETINAL SURGERY
- U-2378 TREATMENT OF POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS
- U-2379 USE IN IDENTIFICATION OF INTRAOCULAR MEMBRANES TO FACILITATE REMOVAL DURING

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OPHTHALMIC SURGERY

- U-2380 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS IN PATIENTS 18 YEARS OF AGE AND OLDER
- U-2381 TREATMENT IN COMBINATION WITH A GNRH AGONIST OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)
- U-2382 TREATMENT IN COMBINATION WITH A GNRH AGONIST OF HIGH RISK NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)
- U-2383 METHOD OF CONTROLLING GLYCEMIA IN A DIABETIC PATIENT WITH DELAYED OR PROLONGED FOOD ABSORPTION BY ADMINISTERING 50 TO 75% OF A PREDETERMINED DOSE OF INSULIN-FDKP AT MEALTIME, AND ADMINISTERING REMAINDER OF DOSE 30-120 MINUTES AFTER BEGINNING OF MEAL
- U-2384 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 10
- U-2385 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1,10 AND 11
- U-2386 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12 AND 19
- U-2387 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12, 19 AND 20
- U-2388 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 21 AND 28
- U-2389 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 21, 28, AND 29
- U-2390 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 30 AND 41
- U-2391 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 30, 41, AND 42
- U-2392 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 43 AND 50
- U-2393 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 43, 50 AND 51
- U-2394 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY
- U-2395 FOR THE TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY
- U-2396 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 8716338 AND IVACAFTOR
- U-2397 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AND IVACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 9192606
- U-2398 TOPICAL TREATMENT OF PRIMARY AXILLARY HYPERHIDROSIS IN ADULTS AND PEDIATRIC PATIENTS 9 YEARS OF AGE AND OLDER
- U-2399 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 12 YEARS AND OLDER, WITH A F508DEL OR G551D CFTR GENE MUTATION AND A A455E, 2789+5G->A, OR 3849+10KBC->T MUTATION, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF U.S. PATENT 10058546
- U-2400 REDUCING ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION

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- U-2401 A METHOD OF TREATING AMYOTROPHIC LATERAL SCLEROSIS IN A PATIENT IN NEED OF SUCH TREATMENT, SAID METHOD COMPRISING ADMINISTERING TO SAID PATIENT AN EFFECTIVE AMOUNT OF A SUSPENSION ACCORDING TO CLAIM 1
- U-2402 TREATMENT OF SCHIZOPHRENIA BY RAPID AND CONTINUOUS INTRAMUSCULAR INJECTION
- U-2403 TREATMENT OF PSORIASIS USING A DOSAGE TITRATION SCHEDULE
- U-2404 METHOD OF DELIVERING SUMATRIPTAN TO A NASAL CAVITY
- U-2405 A METHOD FOR TREATING A BACTERIAL INFECTION IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE
- U-2406 A METHOD FOR TREATING A PATIENT 9 YEARS OF AGE AND OLDER SUFFERING FROM AN INFLAMMATORY SKIN DISORDER OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE
- U-2407 A METHOD FOR TREATING ACNE IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE CRYSTALLINE SALT
- U-2408 A METHOD FOR TREATING A BACTERIAL INFECTION IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE CRYSTALLINE SALT
- U-2409 A METHOD FOR TREATING ACNE IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING SARECYCLINE HYDROCHLORIDE IN 60 MG, 100 MG OR 150 MG EQUIVALENT DOSES
- U-2410 TREATMENT OF ADULT PATIENTS FOR WHOM TREATMENT WITH BOTH AMLODIPINE FOR HYPERTENSION AND CELECOXIB FOR OSTEOARTHRITIS ARE APPROPRIATE
- U-2411 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 12 YEARS OR OLDER WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE TABLET COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1, 19, OR 21 OF U.S. PATENT NO. 10,076,513 AND IVACAFTOR
- U-2412 FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- U-2413 FOR THE TREATMENT OF PATIENTS WITH FOLLICULAR LYMPHOMA (FL)
- U-2414 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS AS PART OF A COMBINATION DRUG REGIMEN
- U-2415 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS AS PART OF A COMBINATION ANTIBACTERIAL DRUG REGIMEN
- U-2416 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS WITH CYSTIC FIBROSIS AS PART OF A COMBINATION DRUG REGIMEN
- U-2417 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN NON-CYSTIC FIBROSIS ADULTS AS PART OF A COMBINATION ANTIBACTERIAL DRUG REGIMEN
- U-2418 METHOD OF ADMINISTERING TESTOSTERONE ENANTHATE SUBCUTANEOUSLY
- U-2419 METHOD OF OPERATING AN INJECTION DEVICE
- U-2420 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-2421 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2422 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME WHO HAVE BEEN PREVIOUSLY TREATED WITH CLOBAZAM
- U-2423 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME WHO HAVE BEEN PREVIOUSLY TREATED WITH CLOBAZAM
- U-2424 USE IN COMBINATION WITH CLOBAZAM FOR TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME
- U-2425 USE FOR THE TREATMENT OF CONVULSIVE SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2426 USE FOR THE TREATMENT OF CONVULSIVE SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME
- U-2427 USE FOR THE TREATMENT OF DROP SEIZURES IN PATIENTS WITH DRAVET SYNDROME

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U-2428	TREATMENT OF PARTIAL-ONSET SEIZURES WITH OR WITHOUT SECONDARILY GENERALIZED SEIZURES IN PATIENTS WITH EPILEPSY 4 YEARS OF AGE AND OLDER
U-2429	TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
U-2430	TREATMENT OF POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN AMYLOIDOSIS
U-2431	TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS
U-2432	LONG-TERM, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
U-2433	METHOD OF TREATING A BIOLOGICAL RHYTHM DISORDER, SUCH AS INSOMNIA
U-2434	USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
U-2435	REDUCTION OF RISK OF MAJOR CARDIOVASCULAR EVENTS (CV DEATH, MI, AND STROKE) IN CHRONIC CAD OR PAD
U-2436	USE IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER TO IMPROVE TREATMENT EMERGENT SEXUAL DYSFUNCTION (TESD) INDUCED BY PRIOR SEROTONIN REUPTAKE INHIBITOR TREATMENT
U-2437	TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BREAST CANCER SUSCEPTIBILITY GENE (BRCA)-MUTATED (GBRCAM) HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE LOCALLY ADVANCED OR METASTATIC BREAST CANCER
U-2438	CARDIOVASCULAR OUTCOMES TRIAL OF LIRAGLUTIDE 1.8 MG IN PATIENTS WITH TYPE 2 DIABETES AND CARDIOVASCULAR DISEASE
U-2439	TREATMENT OF MENOPAUSE SYMPTOMS, INCLUDING VASOMOTOR SYMPTOMS
U-2440	FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
U-2441	REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS
U-2442	USE FOR THE TREATMENT OF ATONIC SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME
U-2443	USE FOR THE TREATMENT OF ATONIC SEIZURES IN PATIENTS WITH DRAVET SYNDROME
U-2444	TREATMENT OF SUBJECTS HAVING BACTERIAL SKIN OR SKIN STRUCTURE INFECTION
U-2445	TREATMENT IN COMBINATION WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
U-2446	TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
U-2447	TREATMENT OF SEVERE HYPERTRIGLYCERIDEMIA (500 MG/DL) IN ADULT PATIENTS AS AN ADJUNCT TO DIET
U-2448	TREATMENT OF TRAVELERS' DIARRHEA CAUSED BY NON-INVASIVE STRAINS OF ESCHERICHIA COLI IN ADULTS
U-2449	TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTION
U-2450	POSITRON EMISSION TOMOGRAPHY DIAGNOSTIC AGENT IN ADULTS WITH SUSPECTED PROSTATE CANCER RECURRENCE BASED ON ELEVATED BLOOD PROSTATE SPECIFIC ANTIGEN LEVELS FOLLOWING PRIOR TREATMENT
U-2451	TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
U-2452	COMBINATION WITH IMMUNOSUPPRESSIVE THERAPY FOR FIRST-LINE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS AND OLDER WITH SEVERE APLASTIC ANEMIA
U-2453	TREATMENT OF FUNGAL INFECTIONS, INCLUDING BLASTOMYCOSIS, HISTOPLASMOSIS, AND ASPERGILLOSIS
U-2454	USE FOR THE TREATMENT OF DROP SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME
U-2455	USE IN COMBINATION WITH CLOBAZAM FOR TREATMENT OF DROP SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME
U-2456	TREATMENT OF ACUTE MYELOID LEUKEMIA (AML)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2457 REINITIATION OF SCHIZOPHRENIA TREATMENT FOLLOWING A MISSED DOSE MORE THAN 9 MONTHS AGO

U-2458 REINITIATION OF SCHIZOPHRENIA TREATMENT FOLLOWING A MISSED DOSE 4-9 MONTHS AGO

U-2459 TREATMENT OF DYSKINESIA AND DECREASING OFF TIME IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS

U-2460 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF CORONARY ARTERY BYPASS GRAFT IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2461 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF CARDIOVASCULAR BYPASS GRAFT AND VASCULATURE IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2462 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF VESSEL WITH ARTERIOVENOUS MALFORMATION IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2463 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION IN SURGICAL FLAPS IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2464 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF TRANSPLANTED ORGAN OR ATTACHED VESSEL IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2465 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF VESSEL GRAFT IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2466 VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF DONOR ORGAN OR ATTACHED VESSEL IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2467 VISUALIZATION OF EXTRAHEPATIC BILIARY DUCT ATTACHED TO DONOR ORGAN IN PATIENTS 12 YEARS AND OLDER

U-2468 VISUALIZATION OF EXTRAHEPATIC BILIARY DUCT ATTACHED TO TRANSPLANTED ORGAN IN PATIENTS 12 YEARS AND OLDER

U-2469 METHOD OF TREATING CANCEROUS SOLID TUMORS

U-2470 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION

U-2471 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK FUSION GENE IN A PEDIATRIC PATIENT

U-2472 METHOD OF TREATING NEUROBLASTOMA, GLIOMA, THYROID, AND BREAST CANCER SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION

U-2473 METHOD OF TREATING CMN, IFS, HGG, DIPGS, PTC, SOFT TISSUE SARCOMA, AND SPINDLE CELL SARCOMA SOLID TUMORS EXHIBITING AN NTRK GENE FUSION IN A PEDIATRIC PATIENT WITH AN ORAL SOLUTION

U-2474 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION AFTER SURGICAL RESECTION

U-2475 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION IN A PEDIATRIC PATIENT

U-2476 USE OF A DELIVERY DEVICE TO DELIVER A DOSE OF NALOXONE

U-2477 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH CYP1A2 STRONG INHIBITORS

U-2478 METHOD FOR THE INDUCTION OF LOCAL ANESTHESIA PRIOR TO PERFORMING A PROCEDURE ON, THROUGH, OR ADJACENT TO THE MUCOUS MEMBRANES

U-2479 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES

U-2480 MAINTENANCE TREATMENT OF GBRCA- OR SBRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM-BASED CHEMOTHERAPY

U-2481 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY

U-2482 TREATMENT OF HR-NEGATIVE, HER-2 NEGATIVE, GBRCA-MUTATED METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR

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METASTATIC SETTING

- U-2483 TREATMENT OF HR-POSITIVE, HER-2 NEGATIVE, GBRCA-MUTATED METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING, AND WITH ENDOCRINE THERAPY OR ARE INAPPROPRIATE FOR ENDOCRINE THERAPY
- U-2484 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE TREATED WITH CARBIDOPA/LEVODOPA BY INHALATION OF LEVODOPA POWDER PARTICLES
- U-2485 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE TREATED WITH CARBIDOPA/LEVODOPA BY INHALATION OF LEVODOPA POWDER PARTICLES THROUGH A SINGLE BREATH ACTIVATED STEP
- U-2486 INTERMITTENT TREATMENT OF OFF EPISODES IN PATIENTS WITH PARKINSON'S DISEASE WITH A POWDER INHALER
- U-2487 DEXTENZA IS APPROVED FOR THE TREATMENT OF OCULAR PAIN FOLLOWING OPHTHALMIC SURGERY
- U-2488 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB
- U-2489 TREATMENT OF MODERATE TO SEVERE OPIOID USE DISORDER
- U-2490 TREATMENT OF COMPLICATED URINARY TRACT INFECTION (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, AND ENTEROBACTER CLOACAE SPECIES COMPLEX
- U-2491 A METHOD FOR DELIVERING A COMPOSITION TO A MUCUS MEMBRANE
- U-2492 A METHOD FOR DELIVERING A PHARMACEUTICAL AGENT ACROSS A MUCOSAL BARRIER
- U-2493 A METHOD FOR TREATING INFLAMMATION AND/OR OTHER DISORDERS IN AN EYE OF A PATIENT
- U-2494 INDICATED FOR THE TREATMENT OF VENTRICULAR ARRHYTHMIAS, SUCH AS SUSTAINED VENTRICULAR TACHYCARDIA, THAT IN THE JUDGEMENT OF THE PHYSICIAN ARE LIFE-THREATENING
- U-2495 VENTRICULAR FIBRILLATION
- U-2496 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE
- U-2497 TREATMENT OF DRUG-INDUCED EXTRAPYRAMIDAL REACTION IN ADULT PATIENTS
- U-2498 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-2499 METHOD OF REDUCING ADVERSE EFFECTS IN PATIENTS SUFFERING FROM EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY WHO ARE CONCOMITANTLY ADMINISTERED SODIUM OXYBATE AND DIVALPROEX SODIUM
- U-2500 USE OF A DELIVERY DEVICE TO DELIVER A BIOEQUIVALENT DOSE OF A NALOXONE COMPOSITION VIA A NEEDLE
- U-2501 TREATMENT OF PARTIAL-ONSET SEIZURES
- U-2502 TREATMENT OF TREATMENT-RESISTANT DEPRESSION IN ADULT IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-2503 TREATMENT OF ADULTS WITH METASTATIC GASTRIC OR GJA PREVIOUSLY TREATED WITH AT LEAST TWO PRIOR LINES OF CHEMOTHERAPY THAT INCLUDED A FLUOROPYRIMIDINE, A PLATINUM, EITHER A TAXANE OR IRINOTECAN, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
- U-2504 TREATMENT OF HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN IN COMBINATION WITH RIBOCICLIB AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- U-2505 TREATMENT OF PRE/PERIMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-2506 METHOD OF TREATING TESTOSTERONE DEFICIENCY
- U-2507 METHOD OF TREATING ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSSI) CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA
- U-2508 A METHOD OF TREATING BACTERIAL INFECTIONS IN COMPLICATED INTRA-ABDOMINAL INFECTION AND COMPLICATED URINARY TRACT INFECTION, INCLUDING PYELONEPHRITIS,

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PATENT USE

PATIENTS COMPRISING ADMINISTERING A BACTERICIDALLY EFFECTIVE AMOUNT OF AVIBACTAM SODIUM

- U-2509 A METHOD OF TREATING A BACTERIAL INFECTION IN COMPLICATED INTRA-ABDOMINAL INFECTION (CIAI) AND COMPLICATED URINARY TRACT INFECTION (CUTI), INCLUDING PYELONEPHRITIS, PATIENTS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AVIBACTAM SODIUM
- U-2510 A METHOD FOR CONTRACEPTION COMPRISING THE STEP OF ORAL ADMINISTRATION A DOSAGE OF 20 MG TO 30 MG OF ULIPRISTAL ACETATE TO A WOMAN WITHIN 72 HOURS AND UP TO 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-2511 A METHOD OF TREATING MULTIPLE SCLEROSIS BY ADMINISTERING SIPONIMOD USING A TITRATION SCHEME TO REACH A MAINTENANCE DOSE
- U-2512 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH AN EFFECTIVE AMOUNT OF TEZACAFTOR AND IVACAFTOR
- U-2513 MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2514 MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2515 PALBOCICLIB FOR HR-POS. HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER IN COMBO WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY IN POSTMENOPAUSAL WOMEN OR MEN, OR WITH FULVESTRANT IN PTS WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-2516 A METHOD FOR REDUCING SERUM GLUCOSE LEVELS IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2517 A METHOD FOR REDUCING SERUM GLUCOSE LEVELS IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2518 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-2519 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2520 TREATING MS WITH ORAL CLADRIBINE ACC. TO THE STEPS (I) INDUCTION PERIOD WITH ABOUT 1.7 MG/KG-3.5 MG/KG CLADRIBINE; (II) CLADRIBINE-FREE PERIOD OF ABOUT 8-10 MONTHS; (III) MAINTENANCE PERIOD WITH ABOUT 1.7 MG/KG CLADRIBINE; (IV) CLADRIBINE-FREE PERIOD
- U-2521 TREATMENT OF MS WITH A TABLET WITH AN ADMIXTURE OF (A) AN AMORPHOUS INCLUSION COMPLEX OF CLADRIBINE AND HYDROXYPROPYL-B-CYCLODEXTRIN AND (B) AMORPHOUS FREE CLADRIBINE AND CYCLODEXTRIN AS A NON-INCLUSION COMPLEX, CLADRIBINE/CYCLODEXTRIN 1:10-1:16 W/W
- U-2522 TREATING RRMS OR SPMS WITH ORAL CLADRIBINE: (I) 2-4 MONTHS INDUCTION WITH 1.7 MG/KG - 3.5 MG/KG CLADRIBINE; (II) CLADRIBINE-FREE PERIOD OF ABOUT 8-10 MONTHS; (III) 2-4 MONTHS MAINTENANCE WITH ABOUT 1.7 MG/KG CLADRIBINE; (IV) CLADRIBINE-FREE PERIOD
- U-2523 TREATMENT OF MS WITH AN ADMIXTURE OF (A) AN AMORPHOUS INCLUSION COMPLEX OF CLADRIBINE (2CDA) AND CYCLODEXTRIN AND (B) AMORPHOUS FREE 2CDA AND CYCLODEXTRIN AS A NON-INCLUSION COMPLEX, FORMULATED AS A SOLID ORAL FORM, W/O SIGN. AMOUNTS OF CRYST. 2CDA
- U-2524 TREATMENT OF THE CARDIOMYOPATHY OF WILD TYPE OR HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS (ATTR-CM)
- U-2525 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY OR URGE INCONTINENCE
- U-2526 ACUTE TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY (I.E., SEIZURE CLUSTERS, ACUTE REPETITIVE SEIZURES) THAT ARE DISTINCT FROM A PATIENT'S USUAL SEIZURE PATTERN IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
- U-2527 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2528 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS A R117H MUTATION IN THE CFTR GENE

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- U-2529 TREATMENT OF A MODERATE MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2530 TREATMENT OF CF IN A PATIENT AGE 6 MONTHS TO < 6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2531 TREATMENT OF CF IN A PATIENT AGE 6 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-2532 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, OR 6 IN ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG
- U-2533 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML)
- U-2534 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML)
- U-2535 USE IN COMBINATION WITH METHYLPREDNISOLONE FOR THE TREATMENT OF PATIENTS WITH PROSTATE CANCER
- U-2536 FOR TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE
- U-2537 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- U-2538 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LEUKEMIA (SLL) IN COMBINATION WITH A GA101 ANTIBODY SUCH AS OBINUTUZUMAB FOR ONE OR MORE DOSING PERIODS, WHEREIN THE CLL OR SLL IS A CD20-EXPRESSING CANCER
- U-2539 IN COMBINATION WITH FULVESTRANT FOR TREATMENT OF POSTMENOPAUSAL WOMEN, AND MEN, WITH HR-POSITIVE, HER-2-NEGATIVE, PIK3CA-MUTATED, ADVANCED OR METASTATIC BREAST CANCER
- U-2540 TREATMENT OF HORMONE RECEPTOR POSITIVE ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN
- U-2541 REDUCING THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION (MI), AND STROKE IN A PATIENT RECEIVING 75-100 MG ASPIRIN DAILY WITH A HISTORY OF MI BY ADMINISTERING 60 MG TICAGRELOR TWICE DAILY
- U-2542 REDUCING THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN A PATIENT RECEIVING 75-100 MG ASPIRIN DAILY AND HAVING OR WHO HAD ACUTE CORONARY SYNDROME BY ADMINISTERING 60 MG TICAGRELOR TWICE DAILY
- U-2543 TREATMENT OF SCHIZOPHRENIA WITH CARIPRAZINE
- U-2544 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH CARIPRAZINE
- U-2545 TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION) WITH CARIPRAZINE
- U-2546 USE FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2547 METHOD OF PROVIDING CONTRACEPTION IN A WOMAN HAVING A BMI OF 25 KG/M2 OR MORE WITH RESULTANT LIMITED BLEEDING EVENTS PER TREATMENT CYCLE
- U-2548 TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY OR OBSTRUCTIVE SLEEP APNEA (OSA)
- U-2549 CONTROL OF SERUM PHOSPHORUS LEVELS
- U-2550 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PREVIOUSLY TREATED FOLLICULAR LYMPHOMA IN COMBINATION WITH A RITUXIMAB PRODUCT
- U-2551 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PREVIOUSLY TREATED MARGINAL ZONE LYMPHOMA IN COMBINATION WITH A RITUXIMAB PRODUCT
- U-2552 METHOD OF TREATING POSTPARTUM DEPRESSION
- U-2553 PREVENTION OF PREGNANCY IN FEMALES OF REPRODUCTIVE AGE
- U-2554 TREATMENT OF MILD TO MODERATE ACTIVE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON
- U-2555 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY

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- INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-2556 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULTS WHO HAVE INTOLERANCE TO OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON ASSOCIATED WITH HEAVY UTERINE BLEEDING OR A GASTROINTESTINAL DISORDER BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2557 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 GRAMS OF ELEMENTAL IRON
- U-2558 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY
- U-2559 USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- U-2560 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
- U-2561 USE IN COMBINATION WITH CISPLATIN FOR TREATMENT OF UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WITHOUT PRIOR CHEMOTHERAPY TREATMENT
- U-2562 TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER IN COMBINATION WITH PREDNISONE
- U-2563 TREATMENT OF ADVANCED GASTRIC ADENOCARCINOMA IN COMBINATION WITH CISPLATIN AND FLUOROURACIL IN PATIENTS THAT HAVE NOT RECEIVED PRIOR CHEMOTHERAPY
- U-2564 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK IN COMBINATION WITH CISPLATIN AND FLUOROURACIL
- U-2565 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA
- U-2566 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA
- U-2567 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM, WHERE THE PATIENT EXPERIENCES NO REBOUND OR WORSENING OF FACIAL ERYTHEMA POST-TREATMENT
- U-2568 TREATMENT OF HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD)
- U-2569 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH AN EFFECTIVE AMOUNT OF TEZACAFTOR AND IVACAFTOR
- U-2570 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-2571 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-2572 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER, WITH A F508DEL OR G551D CFTR GENE MUTATION AND A A455E, 2789+5G->, OR 3849+10KBC->T MUTATION, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF U.S. PATENT 10058546
- U-2573 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR F508DEL AND A SECOND CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2574 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2575 TREATING CYSTIC FIBROSIS PATIENTS AGES 6 AND OLDER, WHO ARE HOMOZYGOUS FOR F508DEL OR HAVE AT LEAST 1 CFTR GENE MUTATION RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS (<30% CRYSTALLINE) IVACAFTOR
- U-2576 TREATMENT OF COMMUNITY ACQUIRED BACTERIAL PNEUMONIA
- U-2577 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC IMMUNE

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- THROMBOCYTOPENIA WHO HAS HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- U-2578 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC LIVER DISEASE WHO IS SCHEDULED TO UNDERGO A PROCEDURE
- U-2579 REDUCTION IN A SUBJECT'S RISK OF EXPERIENCING A BREAKTHROUGH OVERT HEPATIC ENCEPHALOPATHY (HE) EPISODE
- U-2580 A METHOD OF TREATING TYPE 2 DIABETES COMPRISING ADMINISTERING SEMAGLUTIDE ONCE WEEKLY IN A AMOUNT OF 1.0 MG TO A SUBJECT IN NEED THEREOF
- U-2581 TREATING HYPOTENSION WITH ABOUT 20 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK
- U-2582 FOR THE ORAL PREVENTION/PROPHYLAXIS OF MALARIA IN ADULTS, COMPRISING A THREE-PHASE DOSING REGIMEN CONSISTING OF A LOADING/INITIAL DOSE, A MAINTENANCE/EXPOSURE DOSE, AND A TERMINAL/POST-EXPOSURE DOSE
- U-2583 TREATMENT OF BACTERIAL VAGINOSIS IN ADULT WOMEN
- U-2584 XPROVIO IS INDICATED IN COMBINATION WITH DEXAMETHASONE TO TREAT RELAPSED OR REFRACTORY MULTIPLE MYELOMA (REFRACTORY TO AT LEAST AN ANTI-CD38 MAB, 2 PROTEASOME INHIBITORS AND 2 IMMUNOMODULATORY AGENTS) IN ADULTS WHO RECEIVED AT LEAST 4 PRIOR THERAPIES
- U-2585 TREATMENT OF PARENTERAL NUTRITION-ASSOCIATED CHOLESTASIS IN PATIENTS UNDER THE AGE OF 12
- U-2586 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS, INCLUDING PYELONEPHRITIS (CUTI)
- U-2587 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS (CIAI)
- U-2588 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH DAPAGLIFLOZIN AND METFORMIN
- U-2589 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH BASAL INSULIN OR BASAL INSULIN PLUS METFORMIN
- U-2590 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, A THIAZOLIDINEDIONE, OR COMBINATION OF ANY TWO OF THESE THERAPIES
- U-2591 LOWERING PLASMA GLUCAGON IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING EXENATIDE AS AN ADJUNCT TO DIET AND EXERCISE
- U-2592 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A SUSTAINED-RELEASE EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-2593 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2594 REDUCING FASTING PLASMA GLUCOSE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2595 REDUCING BODY WEIGHT IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2596 REDUCING HBA1C IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION ONCE WEEKLY AS AN ADJUNCT TO DIET AND EXERCISE TO ACHIEVE A MEAN STEADY STATE PLASMA CONCENTRATION OF EXENATIDE AT LEAST 170 PG/ML
- U-2597 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH DAPAGLIFLOZIN AS ADD-ON TO METFORMIN
- U-2598 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN INJECTABLE SUSTAINED RELEASE FORMULATION OF EXENATIDE AS AN ADJUNCT TO DIET AND EXERCISE
- U-2599 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING AN EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE TO PROVIDE A RELEASE PROFILE HAVING A RATIO OF C-MAX TO C-AVG OF ABOUT 3 OR LESS
- U-2600 IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY

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ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE

U-2601 STIMULATING INSULIN RELEASE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE

U-2602 DELAYING GASTRIC EMPTYING IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A PRE-MIXED EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE

U-2603 METHOD OF TREATING IRON DEFICIENCY

U-2604 TREATMENT OF SEVERE HYPOGLYCEMIA IN PATIENTS WITH DIABETES

U-2605 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION RESISTANT PROSTATE CANCER

U-2606 TREATMENT OF ADULT PATIENTS WITH SYMPTOMATIC TENOSYNOVIAL GIANT CELL TUMOR (TGCT) ASSOCIATED WITH SEVERE MORBIDITY OR FUNCTIONAL LIMITATIONS AND NOT AMENABLE TO IMPROVEMENT WITH SURGERY

U-2607 TREATMENT OF ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY MYELOFIBROSIS

U-2608 METHOD OF TREATING SCHIZOPHRENIA

U-2609 A METHOD FOR INDUCING A REGIONAL ANAESTHESIA VIA INTRATHECAL ADMINISTRATION OF A PATENTED PRESERVATIVE FREE SOLUTION FOR INJECTION (WITH A SPECIFIC COMPOSITION, PH, OSMOLALITY AND DENSITY) CONTAINING 9-11 MG/ML CHLOROPROCAINE HCL

U-2610 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTION IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS

U-2611 TREATMENT OF COMPLICATED URINARY TRACT INFECTION IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS

U-2612 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN PATIENTS 10 YEARS OF AGE AND OLDER

U-2613 TREATMENT OF RELAPSING-REMITTING SCLEROSIS (MS)

U-2614 TREATMENT OF MODERATE TO SEVERE DYSPAREUNIA

U-2615 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD

U-2616 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE RHEUMATOID ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO METHOTREXATE

U-2617 TREATMENT OF ROS1-POSITIVE NON-SMALL CELL LUNG CANCER

U-2618 TREATMENT OF SOLID TUMORS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION

U-2619 TREATMENT OF ADULTS WITH COMMUNITY-ACQUIRED BACTERIAL PNEUMONIA CAUSED BY SUSCEPTIBLE MICROORGANISMS

U-2620 USE OF NINTEDANIB FOR SLOWING THE RATE OF DECLINE IN PULMONARY FUNCTION IN PATIENTS WITH SYSTEMIC SCLEROSIS-ASSOCIATED INTERSTITIAL LUNG DISEASE (SSC-ILD)

U-2621 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MILD TO MODERATE PAIN

U-2622 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 2 YEARS AND OLDER

U-2623 A METHOD OF REDUCING OFF TIME FROM L-DOPA THERAPY, COMPRISING ADMINISTERING, TO A HUMAN PATIENT WITH PARKINSON'S DISEASE, AN EFFECTIVE AMOUNT OF ISTRADEFYLLINE, WHEREIN THE PATIENT CURRENTLY RECEIVES SAID L-DOPA THERAPY

U-2624 TREATMENT OF METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC)

U-2625 TOPICAL TREATMENT OF PLAQUE PSORIASIS IN ADULTS

U-2626 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION BY ADMINISTERING TENAPANOR

U-2627 TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 12 YEARS AND OLDER

U-2628 METHOD OF TREATING TYPE 2 DIABETES MELLITUS

U-2629 TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS AS A REPLACEMENT THERAPY IN VIROLOGICALLY SUPPRESSED ADULTS WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO THE INDIVIDUAL COMPONENTS OF DELSTRIGO

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- U-2630 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 AS A REPLACEMENT THERAPY IN VIROLOGICALLY SUPPRESSED ADULTS WITH NO HISTORY OF TREATMENT FAILURE AND NO KNOWN SUBSTITUTIONS ASSOCIATED WITH RESISTANCE TO DORAVIRINE
- U-2631 TREATMENT OF COMPLICATED URINARY TRACT INFECTION
- U-2632 REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS
- U-2633 TREATMENT OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER, PROGRESSED ON: CRIZOTINIB + AT LEAST 1 OTHER ALK INHIBITOR FOR METASTATIC DISEASE; OR ALECTINIB, OR CERITINIB AS FIRST ALK INHIBITOR FOR METASTATIC DISEASE.
- U-2634 METHOD OF TREATMENT IN PATIENTS WITH CONCOMITANT ANGIOEDEMA
- U-2635 TREATMENT OF ACUTE URTICARIA
- U-2636 METHOD OF INCREASING PEAK PLASMA OR ONSET OF PLASMA CONCENTRATION BY INTRAVENOUS INJECTION IN INDIVIDUALS IN NEED OF TREATMENT FOR ACUTE URTICARIA
- U-2637 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE WITH A SINGLE UNIT DOSE OF 10% OXYBUTYNIN CHLORIDE GEL
- U-2638 INCREASE PAIN-FREE LIGHT EXPOSURE IN ADULT PATIENTS WITH A HISTORY OF PHOTOTOXIC REACTIONS FROM ERYTHROPOIETIC PROTOPORPHYRIA (EPP)
- U-2639 METHOD OF ACTIVATING RARGAMMA RECEPTOR
- U-2640 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING
- U-2641 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM IN ACUTELY ILL MEDICAL PATIENTS AT RISK FOR THROMBOEMBOLIC COMPLICATIONS NOT AT HIGH RISK OF BLEEDING WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS
- U-2642 METHOD OF TREATING CANCER BY DETECTING A CREATININE CLEARANCE OF A PATIENT AND ADMINISTERING LONSURF
- U-2643 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS 65 YEARS OF AGE OR OLDER AND SYMPTOMS THEREOF
- U-2644 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS 65 YEARS OF AGE OR OLDER
- U-2645 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2646 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE ONE F508DEL MUTATION AND ONE R117H MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2647 TREATMENT OF NON-NODULAR ACNE VULGARIS
- U-2648 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2649 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR; AND ANOTHER COMPOSITION COMPRISING IVACAFTOR
- U-2650 TREATMENT OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE USING A SOLID COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AMORPHOUS IVACAFTOR, AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-2651 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-2652 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-2653 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF

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ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR

- U-2654 TREATMENT OF REFRACTORY CHRONIC GRAFT-VERSUS-HOST DISEASE
- U-2655 A METHOD OF TREATMENT OF ADVANCED OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER ASSOCIATED WITH HOMOLOGOUS RECOMBINATION DEFICIENCY (HRD) POSITIVE STATUS
- U-2656 TREATMENT OF ADULT PATIENTS WITH ACTIVE PSORIATIC ARTHRITIS
- U-2657 TREATMENT OF PATIENTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY
- U-2658 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCET'S DISEASE
- U-2659 TREATMENT OF ADULT PATIENTS WITH ORAL ULCERS ASSOCIATED WITH BEHCET'S DISEASE USING A DOSAGE TITRATION SCHEDULE
- U-2660 TREATMENT OF H. PYLORI INFECTION IN ADULTS
- U-2661 CHRONIC WEIGHT MANAGEMENT IN ADULT PATIENTS USING AN EXTENDED RELEASE TABLET CONTAINING LORCARSERIN HYDROCHLORIDE HEMIHYDRATE
- U-2662 USE OF CALCIPOTRIENE FOAM FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS AGED 4 YEARS AND OLDER
- U-2663 USE IN SONOHYSTEROSALPINOGRAPHY TO ASSESS FALLOPIAN TUBE PATENCY
- U-2664 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY INDUCING SKIPPING OF EXON 51
- U-2665 TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2666 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA
- U-2667 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA
- U-2668 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2669 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2670 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA
- U-2671 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB
- U-2672 TREATMENT OF ACUTE HEPATIC PORPHYRIA
- U-2673 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY CORRECTING A DEFECTIVE GENE FOR DYSTROPHIN
- U-2674 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING BY RESTORING OR INCREASING FUNCTIONAL DYSTROPHIN PROTEIN PRODUCTION
- U-2675 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS HAVING A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- U-2676 TREATMENT OF SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR, AS RECITED IN CLAIM 1
- U-2677 PROPHYLAXIS OF ORGAN REJECTION IN PATIENTS CONVERTED FROM TACROLIMUS IMMEDIATE-RELEASE FORMULATIONS
- U-2678 PROPHYLAXIS OF ORGAN REJECTION IN DE NOVO TRANSPLANT PATIENT
- U-2679 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II AT AN INITIAL RATE OF ABOUT 20 NG/KG/MIN AND TITRATING DOWN TO ACHIEVE AND/OR MAINTAIN A MAP OF ABOUT 65 MM HG OR ABOVE
- U-2680 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II WITH AN INITIAL RATE OF ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN IN A SUBJECT HAVING REFRACTORY HYPOTENSION OR SEVERE HYPOTENSION
- U-2681 TREATING LOW BLOOD PRESSURE WITH ANGIOTENSIN II WITH AN INITIAL RATE OF ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN IN A SUBJECT HAVING REFRACTORY HYPOTENSION OR SEVERE HYPOTENSION, AND TITRATING THE RATE UP
- U-2682 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100MG OF ACALABRUTINIB TWICE DAILY

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- U-2683 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100MG OF ACALABRUTINIB TWICE DAILY
- U-2684 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH OBINUTUZUMAB
- U-2685 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH OBINUTUZUMAB
- U-2686 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2687 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LEUKEMIA IN COMBINATION WITH OBINUTUZUMAB BY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY
- U-2688 USE OF VASCEPA TO LOWER TRIGLYCERIDES AND LDL-C IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2689 USE OF VASCEPA TO TREAT MIXED DYSLIPIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2690 USE OF VASCEPA TO LOWER TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2691 USE OF VASCEPA TO TREAT HYPERTRIGLYCERIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2692 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2693 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN A MIXED DYSLIPIDEMIA ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2694 USE OF VASCEPA TO LOWER TRIGLYCERIDES IN A MIXED DYSLIPIDEMIA ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON STATIN THERAPY
- U-2695 USE OF VASCEPA TO TREAT MIXED HYPERTRIGLYCERIDEMIA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (\geq 150 MG/DL) AND ON STATIN THERAPY
- U-2696 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH, CORONARY REVASCULARIZATION, AND UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2697 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND/OR UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2698 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH AND/OR CORONARY REVASCULARIZATION IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2699 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT (CORONARY REVASCULARIZATION, UNSTABLE ANGINA, STROKE AND/OR MYOCARDIAL INFARCTION) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS
- U-2700 USE OF VASCEPA TO REDUCE TRIGLYCERIDES IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE (TG) LEVELS (ABOUT 200 MG/DL TO LESS THAN ABOUT 500 MG/DL) AND ON ROSUVASTATIN THERAPY
- U-2701 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CORONARY REVASCULARIZATION AND/OR UNSTABLE ANGINA IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS (TG \geq 150 MG/DL TO ABOUT 500 MG/DL)
- U-2702 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT (CARDIOVASCULAR DEATH, CORONARY REVASCULARIZATION AND/OR UNSTABLE ANGINA) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS
- U-2703 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CV EVENT (CV DEATH, CORONARY REVASCULARIZATION, UNSTABLE ANGINA, STROKE AND/OR MYOCARDIAL INFARCTION) IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND DIABETES MELLITUS
- U-2704 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND

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AT LEAST ONE RISK FACTOR FOR CARDIOVASCULAR DISEASE

- U-2705 METHOD OF USING CAPSAICIN IN COMBINATION WITH A GEL COMPOSITION FOR REMOVAL OF CAPSAICIN FROM A TREATMENT AREA OR UNINTENDED AREA
- U-2706 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF ONSET AND/OR RECURRENCE OF CARDIOVASCULAR EVENTS IN A PATIENT WHO HAS ESCAPED THE UNSTABLE PERIOD AFTER CARDIOVASCULAR ANGIOPLASTY
- U-2707 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE OCCURRENCE OF A CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH HYPERCHOLESTEROLEMIA
- U-2708 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER
- U-2709 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2710 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-2711 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 750 MG OF ELEMENTAL IRON
- U-2712 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 750 MG OF ELEMENTAL IRON
- U-2713 MODULATION OF 5-HYDROXYTRYPTAMINE 2 RECEPTOR ACTIVITY IN SCHIZOPHRENIA
- U-2714 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN RESIDUAL SYMPTOMS OF SCHIZOPHRENIA
- U-2715 TREATMENT OF SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR, AS RECITED IN CLAIM 2
- U-2716 MAINTENANCE TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GBRCA-MUTATED METASTATIC PANCREATIC ADENOCARCINOMA WHOSE DISEASE HAS NOT PROGRESSED ON AT LEAST 16 WEEKS OF A FIRST-LINE PLATINUM-BASED CHEMOTHERAPY REGIMEN
- U-2717 ACUTE TREATMENT OF MIGRAINE WITH HEADACHE, WITH OR WITHOUT AURA IN ADULTS
- U-2718 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA IN ADULTS
- U-2719 TREATMENT OF RELAPSING REMITTING MULTIPLE SCLEROSIS BY DETERMINING VARICELLA ZOSTER VIRUS (VZV) STATUS AND VACCINATING PRIOR TO COMMENCING TREATMENT
- U-2720 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS; TOPICAL TREATMENT OF THE TOENAIL(S) DUE TO TRICHOPHYTON RUBRUM AND TRICHOPHYTON MENTAGROPHYTES
- U-2721 TOPICAL TREATMENT OF TINEA UNGUIUM BY USING AN APPLICATOR FOR APPLYING A SOLUTION FOR TREATING TINEA UNGUIUM TO AN AFFECTED PART OF A PATIENT
- U-2722 METHOD OF INTRAVENOUSLY ADMINISTERING A DILUTED CYSTEINE HYDROCHLORIDE SOLUTION TO A NEONATE IN NEED THEREOF
- U-2723 MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR 1 DISORDER
- U-2724 A METHOD OF ORAL DELIVERY OF TREPROSTINIL COMPRISING ADMINISTERING AN ORAL OSMOTIC PHARMACEUTICAL DOSAGE FORM
- U-2725 A METHOD OF TREATING PULMONARY HYPERTENSION AND PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING AN ORAL OSMOTIC PHARMACEUTICAL DOSAGE FORM
- U-2726 TREATMENT OF UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) HARBORING A PLATELET-DERIVED GROWTH FACTOR RECEPTOR ALPHA (PDGFRA) EXON 18 MUTATION
- U-2727 NASAL ADMINISTRATION OF DIAZEPAM FOR TREATMENT OF INTERMITTENT, STEREOTYPIC EPISODES OF FREQUENT SEIZURE ACTIVITY IN PATIENTS 6 YEARS OF AGE AND OLDER
- U-2728 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION IN PATIENTS WITH NON-SQUAMOUS NON-SMALL CELL LUNG CANCER
- U-2729 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION IN PATIENTS WITH MESOTHELIOMA
- U-2730 METHOD OF TREATING TYPE 2 DIABETES MELLITUS USING A PHARMACEUTICAL COMPOSITION COMPRISING EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN
- U-2731 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT

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- (45 ML/MIN/1.73 M² ≤ EGFR < 60 ML/MIN/1.73 M²) BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN IN COMBINATION WITH LINAGLIPTIN AND METFORMIN
- U-2732 METHOD OF TREATING TYPE 2 DIABETES USING A PHARMACEUTICAL COMPOSITION COMPRISING LINAGLIPTIN, METFORMIN, EMPAGLIFLOZIN AND A BASIC AMINO ACID
- U-2733 METHOD OF TREATING A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE THERAPY WITH METFORMIN USING A PHARMACEUTICAL COMPOSITION COMPRISING EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN
- U-2734 METHOD OF TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON, WHO HAVE NON-HEMODIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE, BY ADMINISTERING FERRIC DERISOMALTOSE
- U-2735 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150MG ELAGOLIX WHILE CO-ADMINISTERING RIFAMPIN
- U-2736 METHOD OF TREATING EPITHELIOID SARCOMA
- U-2737 METHOD OF TREATING EPITHELIOID SARCOMA BY INHIBITING ENHANCER OF ZESTE HOMOLOG 2 (EZH2)
- U-2738 METHOD OF TREATING A LUNG METASTASIS OF EPITHELIOID SARCOMA
- U-2739 INCREASING BLOOD PRESSURE WITH AN INITIAL RATE OF ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK, AND TITRATING THE RATE UP.
- U-2740 INCREASING BLOOD PRESSURE WITH A RATE OF ABOUT 20 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN A HUMAN SUBJECT HAVING SEPTIC SHOCK
- U-2741 TREATMENT OF CLOSTRIDIODES DIFFICILE-ASSOCIATED DIARRHEA (CDAD) IN PATIENTS FROM 6 MONTHS OF AGE AND OLDER
- U-2742 TREATMENT OF SEVERE HYPOGLYCEMIA
- U-2743 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF UNSTABLE ANGINA IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2744 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF STROKE IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2745 TREATMENT OF NEUROBLASTOMAS THAT HAVE A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION
- U-2746 USE OF NEXLIZET AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2747 USE OF NEXLETOL AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2748 USE OF NEXLETOL AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR INHIBITING CHOLESTEROL SYNTHESIS TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2749 USE OF NEXLIZET AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR INHIBITING CHOLESTEROL SYNTHESIS TO LOWER LDL-C IN ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE
- U-2750 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INTRAVENOUS INJECTION
- U-2751 A TRANSDERMAL METHOD OF CONTRACEPTION
- U-2752 METHOD OF USING L-CYSTEINE IN AN ADMIXTURE FOR TREATING PATIENTS NEEDING PARENTERAL NUTRITION
- U-2753 INCREASING SURVIVAL IN METASTATIC CASTRATION-RESISTANT PROSTATE CANCER PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING AS A 3 WEEK CYCLE CABAZITAXEL AFTER 5 MG DEXCHLORPHENIRAMINE, 8 MG DEXAMETHASONE, AND AN H₂-AGONIST
- U-2754 TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING
- U-2755 OCULAR EXAMINATION, INTRAOCULAR PRESSURE MEASUREMENT, OR REMOVAL OF FOREIGN BODIES OR SUTURES, IN ADULT AND PEDIATRIC PATIENTS REQUIRING A DISCLOSING AGENT IN COMBINATION WITH A TOPICAL OPHTHALMIC ANESTHETIC
- U-2756 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CARDIOVASCULAR DEATH IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE

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U-2757 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOPHRENIA IN ADULTS BY ADMINISTERING TWO LOADING DOSES OF PALIPERIDONE PALMITATE FOLLOWED BY MAINTENANCE DOSE(S)

U-2758 DOSING REGIMEN FOR THE TREATMENT OF SCHIZOAFFECTIVE DISORDER IN ADULTS AS A MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS BY ADMINISTERING TWO LOADING DOSES OF PALIPERIDONE PALMITATE FOLLOWED BY MAINTENANCE DOSE(S)

U-2759 REDUCTION OF INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH OPEN ANGLE GLAUCOMA(OAG) OR OCULAR HYPERTENSION (OHT) WITH A BIODEGRADABLE BIMATOPROST IMPLANT

U-2760 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 12 YEARS OF AGE OR OLDER

U-2761 INTRAVENOUS SOTALOL DOSING REGIMEN FOR ACHIEVING STEADY STATE CONCENTRATION (EXPOSURE) FASTER COMPARED TO THE CONVENTIONAL ORAL DOSING IN A FACILITY THAT CAN PROVIDE ELECTROCARDIOGRAPHIC MONITORING

U-2762 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A MAJOR CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CARDIOVASCULAR DISEASE

U-2763 METHOD OF TREATING ADULTS WITH SCHIZOPHRENIA COMPRISING ADMINISTERING ASENAPINE VIA A TRANSDERMAL PATCH

U-2764 TREATMENT OF POST-OPERATIVE INFLAMMATION AND PAIN FOLLOWING OCULAR SURGERY

U-2765 TREATMENT OF HIV-1 INFECTION IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40 KG WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS

U-2766 TX OF HIV1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS & PEDIATRIC PATIENTS AT LEAST 40KG HAVING NO PRIOR ARV TX HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ARV REGIMEN FOR AT LEAST 6 MO

U-2767 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG WHO HAVE NO PRIOR ARV TREATMENT HISTORY

U-2768 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CY P450 MONOOXYGENASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ARV REGIMEN FOR AT LEAST 6 MONTHS

U-2769 DOSING REGIMEN FOR INTRAVENOUS SOTALOL FOR ADMINISTRATION IN A FACILITY THAT CAN PROVIDE CONTINUOUS ELECTROCARDIOGRAPHIC MONITORING AND CARDIAC RESUSCITATION.

U-2770 CUSHING'S DISEASE

U-2771 TREATMENT OF MODERATE TO SEVERE PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE OR OLDER

U-2772 MAINTENANCE TREATMENT OF CHRONIC PULMONARY DISEASE (COPD)

U-2774 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS

U-2775 TREATMENT OF A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN

U-2776 TREATMENT OF A TYPE 2 DIABETES MELLITUS PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN

U-2777 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR

U-2778 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 2 TO 5 YEARS OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR

U-2779 TREATMENT OF SPASTICITY

U-2780 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME

U-2781 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME

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U-2782	USE FOR REDUCING CONVULSIVE SEIZURE FREQUENCY IN PATIENTS WITH LENNOX GASTAUT SYNDROME
U-2783	USE FOR REDUCING CONVULSIVE SEIZURE FREQUENCY IN PATIENTS WITH DRAVET SYNDROME
U-2784	A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) AND WHERE THE MUTANT IDH1 HAS THE ABILITY TO CONVERT ALPHA-KETOGLUTARATE INTO 2-HYDROXYGLUTARATE (2-HG)
U-2785	A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) AND WHERE THE MUTANT IDH1 HAS THE ABILITY TO CONVERT ALPHA-KETOGLUTARATE INTO 2-HYDROXYGLUTARATE (2-HG)
U-2786	METHOD OF PREVENTING PREGNANCY BY INSERTING A VAGINAL SYSTEM CONTAINING 103 MG OF SEGESTERONE ACETATE AND 17.4 MG ETHINYL ESTRADIOL INTO A VAGINA FOR UP TO THIRTEEN 21/7-DAY (IN/OUT) CYCLES
U-2787	METHOD OF CONTRACEPTION BY INSERTING A VAGINAL SYSTEM FOR UP TO 13 21/7-DAY (IN/OUT) CYCLES, WHEREIN EFFICACY REQUIRES THE SYSTEM CANNOT BE OUT OF THE VAGINA FOR MORE THAN 2 CUMULATIVE HOURS IN ANY SUCH CYCLE WITHOUT USING ALTERNATIVE CONTRACEPTION
U-2788	TREATMENT OF BREAST CANCER INCLUDING HER2 (ERBB2)-POSITIVE OR -OVEREXPRESSING BREAST CANCER
U-2789	POTASSIUM PHOSPHATES INJECTION IS INDICATED AS A SOURCE OF PHOSPHORUS IN INTRAVENOUS FLUIDS TO CORRECT HYPOPHOSPHATEMIA IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
U-2790	TREATMENT OF A TREATMENT-NAIVE PATIENT WITH INADEQUATELY CONTROLLED TYPE 2 DIABETES USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN
U-2791	TREATMENT OF ADULT PATIENTS WITH INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE
U-2792	TREATMENT OF A TREATMENT-NAIVE PATIENT WITH INADEQUATELY CONTROLLED TYPE 2 DIABETES USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN
U-2793	A METHOD FOR DELIVERING NITRIC OXIDE TO A PATIENT WITH PULMONARY HYPERTENSION OR HYPOXIA
U-2794	TREATMENT OF TYPE 2 DIABETES MELLITUS WITH 100 MG CANAGLIFLOZIN PER DAY
U-2795	TREATMENT OF TYPE 2 DIABETES MELLITUS WITH 300 MG CANAGLIFLOZIN PER DAY
U-2796	REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 100 MG CANAGLIFLOZIN PER DAY
U-2797	REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 300 MG CANAGLIFLOZIN PER DAY
U-2798	REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 100 MG CANAGLIFLOZIN PER DAY
U-2799	REDUCTION OF RISK OF END STAGE KIDNEY DISEASE, DOUBLING OF SERUM CREATININE, CARDIOVASCULAR DEATH, AND HOSPITALIZATION FOR HEART FAILURE IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS WITH 300 MG CANAGLIFLOZIN PER DAY
U-2800	TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH NEUROFIBROMATOSIS TYPE 1 (NF1) WHO HAVE SYMPTOMATIC, INOPERABLE PLEXIFORM NEUROFIBROMAS (PN)
U-2801	METHOD OF IRON ADMINISTRATION TO TREAT PATIENTS IN NEED OF IRON REPLACEMENT THERAPY
U-2802	BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH CETUXIMAB, FOR THE TREATMENT OF ADULT PATIENTS WITH METASTATIC COLORECTAL CANCER (CRC) WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, AFTER PRIOR THERAPY
U-2803	BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH BINIMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH A BRAF V600E OR V600K MUTATION, AS DETECTED BY AN FDA- APPROVED TEST
U-2804	A METHOD FOR THE IMPROVEMENT OF NEUROLOGICAL OUTCOME BY REDUCING THE INCIDENCE AND SEVERITY OF ISCHEMIC DEFICITS IN ADULT PATIENTS WITH SUBARACHNOID HEMORRHAGE (SAH) FROM RUPTURED INTRACRANIAL BERRY ANEURYSMS
U-2805	TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY

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U-2806 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF FOOD

U-2807 TREATMENT OF MODERATE TO SEVERE MIGRAINE PAIN WITH PAIN FREE AT 2 HOURS POST ADMINISTRATION

U-2808 TREATMENT OF DYSKINESIA, DECREASING OFF TIME, AND INCREASING ON TIME WITHOUT TROUBLESOME DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS

U-2809 FOR THE TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FIBROBLAST GROWTH FACTOR RECEPTOR 2 (FGFR2) FUSION OR OTHER REARRANGEMENT

U-2810 METHOD OF SUPPORTING EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN

U-2811 METHOD OF TREATING PARKINSON'S DISEASE

U-2812 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE

U-2813 USE FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH A MESENCHYMAL-EPITHELIAL TRANSITION (MET) EXON 14 SKIPPING MUTATION

U-2814 A METHOD OF PROPHYLACTIC TREATMENT OF GOUT FLARES IN ADULTS COMPRISES ADMINISTERING TO A PATIENT A LIQUID COLCHICINE ORAL SOLUTION

U-2815 MEASURING TIME-VARYING CHANGE IN BLOOD IN A TISSUE VOLUME USING MODIFIED BEER-LAMBERT LAW IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO-AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY

U-2816 METHOD FOR TREATING INFLUENZA

U-2817 METHOD OF INHIBITING COMT IN THE PERIPHERY

U-2818 METHOD OF REDUCING O-METHYLATION OF L-DOPA

U-2819 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF ADV. EPITHELIAL OVARIAN CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION

U-2820 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF ADV. EPITHELIAL OVARIAN CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH GENOMIC INSTABILITY

U-2821 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF FALLOPIAN TUBE CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION

U-2822 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF FALLOPIAN TUBE CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH GENOMIC INSTABILITY

U-2823 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH A DELETERIOUS OR SUSPECTED DELETERIOUS BRCA MUTATION

U-2824 MAINTENANCE TREATMENT WITH BEVACIZUMAB OF PRIMARY PERITONEAL CANCER IN COMPLETE OR PARTIAL RESPONSE TO FIRST-LINE PLATINUM CHEMOTHERAPY AND HOMOLOGOUS RECOMBINATION DEFICIENCY-POSITIVE WITH GENOMIC INSTABILITY

U-2825 TREATMENT OF "OFF" EPISODES IN PATIENTS WITH PARKINSON'S DISEASE

U-2826 TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER

U-2827 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY

U-2828 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)

U-2829 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS

U-2830 A METHOD FOR TREATING METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC),

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PATENT USE

WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA-MUTATION

- U-2831 TREATMENT OF PARTIAL-ONSET SEIZURES IN A PATIENT WITH REFRACTORY PARTIAL-ONSET SEIZURES
- U-2832 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC HOMOLOGOUS RECOMBINATION REPAIR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER, WHICH HAS PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE
- U-2833 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER, WHICH HAS PROGRESSED FOLLOWING PRIOR TREATMENT WITH ENZALUTAMIDE OR ABIRATERONE
- U-2834 TREATMENT OF ADULTS WITH PERSISTENT/RECURRENT CHRONIC THROMBOEMBOLIC PULMONARY HYPERTENSION (CTEPH), (WHO GROUP 4) AFTER SURGICAL TREATMENT, OR INOPERABLE CTEPH, TO IMPROVE EXERCISE CAPACITY AND WHO FUNCTIONAL CLASS
- U-2835 TREATMENT OF ADULTS WITH PULMONARY HYPERTENSION (PAH), (WHO GROUP 1), TO IMPROVE EXERCISE CAPACITY, WHO FUNCTIONAL CLASS AND TO DELAY CLINICAL WORSENING
- U-2836 TREATMENT OF ADULT PATIENTS WITH SMALL CELL LUNG CANCER (SCLC) WITH DISEASE PROGRESSION ON OR AFTER PLATINUM-BASED CHEMOTHERAPY.
- U-2837 TREATMENT OF ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC)
- U-2838 REDUCTION OF THE RATE OF A FIRST MYOCARDIAL INFARCTION OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE AT HIGH RISK FOR SUCH EVENTS
- U-2839 TREATMENT OF MYOCARDIAL INFARCTION OR STROKE IN PATIENTS WITH CORONARY ARTERY DISEASE AT HIGH RISK FOR SUCH EVENTS
- U-2840 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP)
- U-2841 USE OF VASCEPA WITH HIGH INTENSITY STATIN THERAPY TO REDUCE THE RISK OF A CV EVENT IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND (1) ESTABLISHED CV DISEASE, OR (2) DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-2842 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS)
- U-2843 NASAL ADMINISTRATION OF METOCLOPRAMIDE FOR TREATMENT OF DIABETIC GASTROPARESIS
- U-2844 IN COMBINATION WITH PEMBROLIZUMAB FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- U-2845 A METHOD OF TREATING A HUMAN PATIENT SUFFERING FROM PULMONARY HYPERTENSION
- U-2846 TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE REFRACTORY TO SYSTEMIC THERAPY
- U-2847 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE IN ADULTS
- U-2848 TREATMENT OF TRAVELERS' DIARRHEA (TD) CAUSED BY NONINVASIVE STRAINS OF ESCHERIA COLI IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-2849 METHOD OF TREATING BLEPHAROPTOSIS
- U-2850 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150 MG ELAGOLIX WHILE CO-ADMINISTERING KETOCONAZOLE
- U-2851 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA POSITIVE FOR AN ENHANCER OF ZESTE HOMOLOG 2 (EZH2) MUTATION BY INHIBITING EZH2
- U-2852 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA POSITIVE FOR AN ENHANCER OF ZESTE HOMOLOG 2 (EZH2) MUTATION
- U-2853 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA
- U-2854 METHOD OF TREATING RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA BY INHIBITING EZH2
- U-2855 XPOVIO IS INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY DIFFUSE LARGE B-CELL LYMPHOMA (DLBCL), NOT OTHERWISE SPECIFIED, INCLUDING DLBCL ARISING FROM FOLLICULAR LYMPHOMA, AFTER AT LEAST 2 LINES OF SYSTEMIC THERAPY
- U-2856 INCREASING SURVIVAL IN METASTATIC CASTRATION-RESISTANT PROSTATE CANCER PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING 20 TO 25 MG/M2 CABAZITAXEL AFTER A PREMEDICATION REGIMEN THAT INCLUDES AN H2-ANTAGONIST
- U-2857 USE OF ORAL OCTREOTIDE FOR LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHO HAVE RESPONDED TO AND TOLERATED TREATMENT WITH OCTREOTIDE OR

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PATENT USE

LANREOTIDE

- U-2858 USE IN COMBINATION WITH STIRIPENTOL, VALPROATE, AND CLOBAZAM FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2859 USE OF CARDIAC MONITORING AND RESTRICTED DISTRIBUTION OF FENFLURAMINE TO MITIGATE RISK OF CARDIOVASCULAR TOXICITY IN THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2860 USE IN COMBINATION WITH STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2861 USE IN COMBINATION WITH CANNABIDIOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-2862 USE FOR THE TREATMENT OF FOCAL SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2863 TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS 6 MONTHS OF AGE AND OLDER
- U-2864 METHOD FOR INHIBITING CYTIDINE DEAMINASE BY ADMINISTERING CEDAZURIDINE
- U-2865 TREATMENT OF MYELODYSPLASTIC SYNDROME
- U-2866 TREATMENT OF CHRONIC MYELOMONOCYTIC LEUKEMIA
- U-2867 METHOD FOR INHIBITING DEGRADATION OF A CDA SUBSTRATE BY ADMINISTERING CEDAZURIDINE
- U-2868 TREATMENT OF NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE
- U-2869 IV ADMINISTRATION OF CANGRELOR BEFORE PCI AND CONTINUOUS INFUSION FOR AT LEAST 2 HOURS OR THE DURATION OF THE PCI AND, DURING OR AFTER THE CONTINUOUS INFUSION, ADMINISTRATION OF A LOADING DOSE OF TICAGRELOR OR AN EQUIVALENT THERAPY (PER LABELING)
- U-2870 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 2
- U-2871 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 2 AND 3
- U-2872 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 4 AND 5
- U-2873 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 4, 5, AND 6
- U-2874 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 7 AND 8
- U-2875 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 7, 8, AND 9
- U-2876 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 10 AND 11
- U-2877 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 10, 11, AND 12
- U-2878 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13 AND 14
- U-2879 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13, 14, AND 15
- U-2880 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 16 AND 17
- U-2881 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 16, 17, AND 18

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U-2882	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 19 AND 20
U-2883	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 19, 20, AND 21
U-2884	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 22, 23, AND 24
U-2885	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 25 AND 26
U-2886	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 25, 26, AND 27
U-2887	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 28 AND 29
U-2888	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 28, 29, AND 30
U-2889	USE FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
U-2890	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 2 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2891	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 3 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2892	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 4 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2893	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 5 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2894	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 6 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2895	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2896	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 8 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2897	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1 AND 9
U-2898	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1 AND 10
U-2899	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN

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CLAIMS 1 AND 11

U-2900	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1 AND 13
U-2901	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 15 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2902	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 16 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2903	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 17 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2904	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1, 17 AND 18 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2905	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1 AND 19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
U-2906	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 22 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2907	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 22 AND 23 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2908	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 22 AND 24 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2909	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 22 AND 25 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 22
U-2910	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 26 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2911	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 27 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2912	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 28 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2913	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 29 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2914	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 30 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
U-2915	ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 31 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2916 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 32 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2917 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 33 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS
- U-2918 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 34 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2919 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 36 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2920 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 38 AND WHEREIN THE EFFECTS ARE AS RECITED CLAIM 26
- U-2921 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS
- U-2922 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 39 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2923 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 26 AND 40 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 26
- U-2924 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 42 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 42
- U-2925 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 42 AND 43 WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 42
- U-2926 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 44 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2927 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 45 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2928 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 46 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2929 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 47 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2930 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 48 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2931 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2

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- DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 49 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2932 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 50 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2933 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 51 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2934 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 52 WHEREIN THE EFFECTS ARE AS RECITED IN SAID CLAIMS
- U-2935 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 44 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 44 AND 54
- U-2936 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 56 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2937 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 44 AND 57 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 44
- U-2938 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY WITH A MIXTURE OF SODIUM, POTASSIUM, MAGNESIUM, AND CALCIUM SALTS OF GHB
- U-2939 TREATMENT OF HIV INFECTION IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 40KG USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE
- U-2940 METHOD OF TREATING PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY
- U-2941 A METHOD OF USING AN AEROSOL DELIVERY DEVICE TO AEROSOLIZE GLYCOPYRROLATE FOR THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2942 METHOD OF TREATING ACNE VULGARIS WITH TOPICALLY APPLIED CORTEXOLONE 17A-PROPIONATE
- U-2943 TREATMENT OF RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA
- U-2944 TREATMENT OF RELAPSED OR REFRACTORY SMALL LYMPHOCYTIC LYMPHOMA
- U-2945 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN COMBINATION WITH URSODEOXYCHOLIC ACID (UDCA) IN ADULTS WITH AN INADEQUATE RESPONSE TO UDCA, OR AS MONOTHERAPY IN ADULTS UNABLE TO TOLERATE UDCA
- U-2946 TREATMENT OF COLORECTAL CANCER THAT HAS A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION
- U-2947 KYPROLIS IS INDICATED IN COMBINATION WITH DARATUMUMAB PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-2948 A METHOD OF POSITIONING AN INTRAUTERINE SYSTEM (IUS) BY DETERMINING A DEPTH OF THE UTERUS, HOLDING AN INSERTER HANDLE WITH ONE HAND, INSERTING THE IUS INTO THE UTERUS, AND RETRACTING A SLIDER ON THE HANDLE TO RELEASE THE IUS INTO THE UTERUS
- U-2949 A METHOD FOR INDUCING A POST-SURGICAL ANALGESIA SPARING EFFECT BY IMPLANTING AT THE SURGICAL SITE A COLLAGEN SPONGE CONTAINING BUPIVACAINE HCL WHICH PROVIDES LOCAL ANESTHESIA FOR UP TO 24 HOURS FOLLOWING IMPLANTATION
- U-2950 CONTINUED TREATMENT OF ADULTS WITH ACUTE MYELOID LEUKEMIA WHO ACHIEVED FIRST COMPLETE REMISSION (CR) OR CR WITH INCOMPLETE BLOOD COUNT RECOVERY FOLLOWING INTENSIVE INDUCTION CHEMOTHERAPY AND ARE NOT ABLE TO COMPLETE INTENSIVE CURATIVE THERAPY
- U-2951 USE OF CU-64 DOTATATE WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION

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- OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT PATIENTS
- U-2952 TREATMENT OF ADULT PATIENTS WITH METASTATIC REARRANGED DURING TRANSFECTION (RET) FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST
- U-2953 USE OF FLUTICASONE FUROATE FOR THE TREATMENT OF AN INFLAMMATORY OR ALLERGIC CONDITIONS, INCLUDING CHRONIC OBSTRUCTIVE PULMONARY DISEASE AND ASTHMA
- U-2954 METHOD OF DISPENSING A COMBINATION MEDICAMENT PRODUCT FROM CLAIMED DELIVERY DEVICE, FOR EXAMPLE FOR THE TREATMENT OF ASTHMA OR COPD
- U-2955 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA; AND ASTHMA
- U-2956 METHOD OF TREATING LAMBERT-EATON MYASTHENIC SYNDROME WITH AMIFAMPRIDINE
- U-2957 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 18 YRS AND OLDER, OR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2958 A METHOD FOR CONTRACEPTION, THE METHOD COMPRISING ADMINISTERING A TABLET COMPRISING 20 MG TO 30 MG OF ULIPRISTAL ACETATE TO A WOMAN WITHIN 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-2959 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A THIRD AND FURTHER CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ELEVATED TG LEVELS (≥ 150 MG/DL) AND ESTABLISHED CARDIOVASCULAR DISEASE
- U-2960 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A SECOND OR FURTHER CARDIOVASCULAR (CV) EVENT IN AN ADULT PATIENT WITH ELEVATED TG LEVELS (≥ 150 MG/DL) AND DIABETES MELLITUS AND 2 OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-2961 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF MYOCARDIAL INFARCTION, STROKE, BOTH IN AN ADULT PATIENT WITH TYPE 2 DIABETES MELLITUS
- U-2962 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF CORONARY REVASCULARIZATION IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE
- U-2963 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2964 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS A R117H MUTATION IN THE CFTR GENE
- U-2965 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-2966 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2967 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-2968 USE OF REMIMAZOLAM FOR INDUCTION AND MAINTENANCE OF PROCEDURAL SEDATION IN ADULTS UNDERGOING PROCEDURES LASTING 30 MINUTES OR LESS
- U-2969 TREATMENT OF ADULT PATIENTS WITH CYCLOSPORIN-RESISTANT, STEROID-DEPENDENT/REFRACTORY, OR STEROID RESISTANT CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2970 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-2971 THE TREATMENT OF ADULT PATIENTS WITH METASTATIC RET FUSION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- U-2972 THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET-MUTANT MEDULLARY THYROID CANCER (MTC) WHO REQUIRE SYSTEMIC THERAPY
- U-2973 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS PAPILLARY THYROID CANCER

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- U-2974 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS MEDULLARY THYROID CANCER
- U-2975 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS DIFFERENTIATED THYROID CANCER
- U-2976 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY, WHEREIN THE CANCER IS RECURRENT THYROID CANCER
- U-2977 TREATING ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE REFRACTORY DIFFERENTIATED THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND ARE RADIOACTIVE IODINE-REFRACTORY
- U-2978 TREATMENT OF HIV-1 INFECTION IN ADULT OR PEDIATRIC PATIENTS (≥ 40 KG) WITH < 50 COPIES/ML HIV-1 RNA AFTER ≥ 6 MONTHS ON PRIOR ANTIRETROVIRAL REGIMEN AND NO KNOWN DARUNAVIR OR TENOFOVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
- U-2979 METHOD COMPRISING IV ADMINISTRATION OF CANGRELOR BEFORE PCI THEN CONTINUOUS INFUSION FOR AT LEAST 2 HOURS OR THE DURATION OF PCI AND, DURING OR AFTER CONTINUOUS INFUSION, ADMINISTRATION OF A LOADING DOSE OF TICAGRELOR, OR AN EQUIVALENT METHOD
- U-2980 METHOD OF TREATING AN ALLERGIC REACTION USING AN AUTO-INJECTOR
- U-2981 A METHOD OF TREATING ACUTE MYELOGENOUS LEUKEMIA (AML) IN A SUBJECT BY ADMINISTERING A PHARMACEUTICAL COMPOSITION WHERE THE AML IS CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 AND THE AML IS NEWLY DIAGNOSED
- U-2982 A METHOD OF TREATING ACUTE MYELOGENOUS LEUKEMIA (AML) IN A SUBJECT BY ADMINISTERING A PHARMACEUTICAL COMPOSITION WHERE THE AML IS CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 AND WHERE THE AML IS RELAPSED/REFRACTORY
- U-2983 TREATMENT OF SCHIZOPHRENIA BY RAPID AND CONTINUOUS INJECTION
- U-2984 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (AT LEAST 12 YEARS OF AGE AND 40 KG) REQUIRING HOSPITALIZATION
- U-2985 A METHOD FOR TREATING DRY EYE IN A PATIENT
- U-2986 MANAGEMENT OF ACUTE PAIN BY INTRAVENOUS INJECTION
- U-2987 METHOD OF TREATING LUNG CANCER, UNDIFFERENTIATED SARCOMA, OR COLORECTAL CANCER THAT EXHIBITS AN NTRK GENE FUSION
- U-2988 REDUCTION OF THE RISK OF STROKE IN PATIENTS WITH ACUTE ISCHEMIC STROKE OR HIGH-RISK TRANSIENT ISCHEMIC ATTACK
- U-2989 METHOD OF USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2990 METHOD OF USE FOR TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2991 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A TABLET CONTAINING SELEXIPAG
- U-2992 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A SOLID PREPARATION CONTAINING SELEXIPAG
- U-2993 A METHOD FOR TREATING A SUBJECT HAVING ADHD, SAID METHOD COMPRISING ORALLY ADMINISTERING TO SAID SUBJECT A RACEMIC METHYLPHENIDATE CHEWABLE TABLET AS CLAIMED
- U-2994 REDUCTION OF RISK OF MYOCARDIAL INFARCTION, STROKE OR CARDIOVASCULAR DEATH IN A PATIENT WITH CHRONIC CAD OR PAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-2995 TREATMENT OF PRIMARY HYPEROXALURIA TYPE 1 (PH1)
- U-2996 ADMINISTRATION OF AN EXTENDED RELEASE TABLET FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-2997 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF STROKE IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDES AND ATRIAL FIBRILLATION

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- U-2998 METHOD OF ADMINISTERING DOCETAXEL TO A SUBJECT COMBINING THE DOCETAXEL PRO-EMULSION FORMULATION WITH AN AQUEOUS MEDIUM TO PRODUCE DOCETAXEL EMULSION
- U-2999 METHOD OF USE OF TREATING, AS AN INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY IN PARTIAL ONSET SEIZURE PATIENTS WITH EPILEPSY AGED 17 YEARS OR OLDER
- U-3000 METHOD FOR POST-EXPOSURE PROPHYLAXIS OF INFLUENZA
- U-3001 PROCEDURES IN ADULT AND PEDIATRIC PATIENTS REQUIRING A DISCLOSING AGENT IN COMBINATION WITH A TOPICAL OPHTHALMIC ANESTHETIC.
- U-3002 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC RET FUSION-POSITIVE THYROID CANCER WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY (IF RADIOACTIVE IODINE IS APPROPRIATE)
- U-3003 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON
- U-3004 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH A STRONG CYP1A2 INHIBITOR
- U-3005 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE USE OF TASIMELTEON WITH RIFAMPIN
- U-3006 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH CYP1A2 STRONG INHIBITORS
- U-3007 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD
- U-3008 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY
- U-3009 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY ADMINISTERING TASIMELTEON TO PATIENTS WITH A SMOKING HISTORY
- U-3010 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3011 A METHOD FOR TREATING OCULAR INFLAMMATION
- U-3012 TREATMENT IN COMBINATION WITH ANDROGEN DEPRIVATION THERAPY OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC) THAT IMPROVES METASTASIS FREE SURVIVAL
- U-3013 TREATMENT IN COMBINATION WITH ORCHIECTOMY OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC)
- U-3014 METHOD FOR THE INDUCTION OF LOCAL ANESTHESIA OF THE MUCOUS MEMBRANES
- U-3015 TOPICAL TREATMENT OF ACTINIC KERATOSIS OF THE FACE OR SCALP
- U-3016 ADJUVANT THERAPY AFTER TUMOR RESECTION IN PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS
- U-3017 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY PATIENTS WITH A SALT OF GAMMA-HYDROXYBUTYRATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3018 XPOVIO IS INDICATED IN COMBINATION WITH BORTEZOMIB AND DEXAMETHASONE FOR THE TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3019 TREATMENT OF ADULT PATIENTS WITH ADVANCED PROSTATE CANCER THAT IS SEX-HORMONE-DEPENDENT
- U-3020 TREATMENT OF ADULT PATIENTS WITH ADVANCED PROSTATE CANCER
- U-3021 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE
- U-3022 TREATMENT OF CF IN PATIENTS 12 YEARS AND OLDER WHO HAVE A F508DEL OR G551D CFTR MUTATION AND A 2ND MUTATION SELECTED FROM R117H, A455E, 2789+5G->A, & 3849+10KBC->T, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 10058546
- U-3023 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WHO HAVE A F508DEL OR G551D CFTR MUTATION AND A 2ND MUTATION SELECTED FROM R117H, A455E, 2789+5G->A, AND

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- 3849+10KBC->T, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF US 10058546
- U-3024 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-3025 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-3026 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-3027 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR AT LEAST ONE CFTR MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,206,877
- U-3028 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3029 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3030 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3031 TREATMENT OF CF IN PATIENTS 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3032 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-3033 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3034 TREATMENT OF TRD IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS DURING THE INDUCTION PHASE IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3035 TREATMENT OF DEPRESSIVE SYMPTOMS IN ADULTS WITH MDD WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR BY NASALLY ADMINISTERING 56MG OR 84 MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3036 TREATMENT OF TRD IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE 2X WEEKLY FOR 4 WEEKS DURING THE INDUCTION PHASE FOLLOWED BY A MAINTENANCE PHASE OF 56MG OR 84 MG WEEKLY OR 1X EVERY TWO WEEKS IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3037 A METHOD OF DELIVERING NITRIC OXIDE TO A PATIENT
- U-3038 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INTRAVENOUS INJECTION IN PATIENTS WITH MILD RENAL IMPAIRMENT
- U-3039 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 53 SKIPPING
- U-3040 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS WITH MILD HEPATIC IMPAIRMENT AND ARE CONCURRENTLY TAKING A STRONG OR MODERATE CYP3A INHIBITOR
- U-3041 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS WITH MODERATE TO SEVERE RENAL IMPAIRMENT
- U-3042 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 POOR METABOLIZERS WITH 84 MG ONCE DAILY OF ELIGLUSTAT (EQUIVALENT TO 100 MG OF ELIGLUSTAT TARTRATE)
- U-3043 LONG-TERM TREATMENT OF ADULTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE OR INTERMEDIATE METABOLIZERS WITH 84 MG TWICE PER DAY OF ELIGLUSTAT

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(EQUIVALENT TO 100 MG OF ELIGLUSTAT TARTRATE TWICE PER DAY)

- U-3044 AXITINIB IN COMBINATION WITH AVELUMAB FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- U-3045 TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-3046 METHOD OF ADMINISTERING VALBENAZINE WHILE AVOIDING CONCOMITANT USE OF A STRONG CYP3A4 INDUCER
- U-3047 USE IN COMBINATION WITH CAPECITABINE, FOR THE TREATMENT OF ADULT PATIENTS WITH ADVANCED OR METASTATIC HER2-POSITIVE BREAST CANCER WHO HAVE RECEIVED TWO OR MORE PRIOR ANTI-HER2 BASED REGIMENS IN THE METASTATIC SETTING
- U-3048 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 500 TO 750 MG OF ELEMENTAL IRON
- U-3049 A METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE 500 TO 750 MG OF ELEMENTAL IRON
- U-3050 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3051 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3052 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT ON STATIN THERAPY AND HAVING ATRIAL FIBRILLATION AND TRIGLYCERIDE LEVELS OF GREATER THAN 500 MG/DL
- U-3053 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF MYOCARDIAL INFARCTION IN AN ADULT PATIENT WITH ELEVATED TRIGLYCERIDE LEVELS AND ESTABLISHED CV DISEASE OR DIABETES MELLITUS AND TWO OR MORE ADDITIONAL RISK FACTORS FOR CV DISEASE
- U-3054 TREATMENT OF DRUG-INDUCED EXTRAPYRAMIDAL REACTIONS IN ADULT PATIENTS WITH PARKINSON'S DISEASE
- U-3055 A METHOD OF TREATING HUNTINGTON'S CHOREA
- U-3056 TREATMENT OF PATIENTS WITH ACTIVE LUPUS NEPHRITIS
- U-3057 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE
- U-3058 TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER AND YOUNG ADULTS WITH RELAPSED OR REFRACTORY, SYSTEMIC ANAPLASTIC LARGE CELL LYMPHOMA (ALCL) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE
- U-3059 TREATMENT OF HIV-1 INFECTION IN ADULTS TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3060 TREATMENT OF HIV INFECTION IN ADULTS
- U-3061 TREATMENT OF HIV-1 IN AN ADULT IN COMBINATION WITH RILPIVIRINE
- U-3062 REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HEART FAILURE (HF) HOSPITALIZATION FOLLOWING A HOSPITALIZATION FOR HF OR NEED FOR OUTPATIENT IV DIURETICS, IN ADULTS WITH SYMPTOMATIC CHRONIC HF AND EJECTION FRACTION LESS THAN 45%
- U-3063 RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA (MZL) WHO HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED REGIMEN
- U-3064 RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) WHO HAVE RECEIVED AT LEAST THREE PRIOR LINES OF SYSTEMIC THERAPY
- U-3065 TREATMENT OF ADULTS WITH METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-3066 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC, SURGICALLY UNRESECTABLE UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY

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- U-3067 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA THAT HAS SUSCEPTIBLE FGFR3 GENETIC ALTERATIONS AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY
- U-3068 TREATMENT IN COMBINATION WITH CABOTEGRAVIR OF HIV-1 INFECTION IN ADULTS TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3069 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN PATIENTS 6 MONTHS AND OLDER
- U-3070 REDUCING THE RISK OF MORTALITY IN HUTCHINSON-GILFORD PROGERIA SYNDROME (HGPS)
- U-3071 USE FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3072 USE FOR THE TREATMENT OF GENERALIZED SEIZURES OR FOCAL SEIZURES WITH IMPAIRMENT IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3073 USE FOR REDUCING SEIZURE FREQUENCY IN PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX
- U-3074 METHOD FOR PROVIDING SUSTAINED LOCAL ANESTHESIA FOR AT LEAST 24 HOURS
- U-3075 TREATMENT OF ADRENAL INSUFFICIENCY
- U-3076 METHOD OF TREATING TARDIVE DYSKINESIA WHILE AVOIDING CONCOMITANT USE OF A STRONG CYP3A4 INDUCER
- U-3077 TREATING A SOLID TUMOR, INCLUDING LUNG CANCER, WITH A MET ALTERATION(S), OR STABILIZING OR IMPROVING SYMPTOMS ASSOCIATED WITH HAVING A SOLID TUMOR, INCLUDING LUNG CANCER, WITH A MET ALTERATION(S), BY ADMINISTERING AN EFFECTIVE AMOUNT OF TEPOTINIB
- U-3078 TREATING A SOLID TUMOR, INCLUDING LUNG CANCER, HAVING A MET KINASE ALTERATION(S) BY ADMINISTERING AN EFFECTIVE AMOUNT OF TEPOTINIB
- U-3079 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A PLATINUM/ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3080 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A TOPOTECAN-CONTAINING REGIMEN FOR EXTENSIVE- STAGE SMALL CELL LUNG CANCER
- U-3081 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A CARBOPLATIN AND ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3082 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH OBSTRUCTIVE SLEEP APNEA (OSA) IN AN ADULT THROUGH A DOSING REGIMEN THAT INCLUDES ORAL ADMINISTRATION OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3083 METHOD OF TREATING TRANSFUSIONAL IRON OVERLOAD
- U-3084 TREATMENT OF HEART FAILURE WITH PRESERVED EJECTION FRACTION
- U-3085 DOSE MODIFICATION FOR RENAL IMPAIRMENT
- U-3086 FOR HIGH-DOSE CONDITIONING TREATMENT PRIOR TO HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- U-3087 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING BY INDUCING EXON-SKIPPING OF EXON 45
- U-3088 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING
- U-3089 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 45 SKIPPING BY RESTORING AN MRNA READING FRAME TO INDUCE DYSTROPHIN PROTEIN PRODUCTION
- U-3090 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-3091 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA IN PATIENTS WITH END-STAGE RENAL DISEASE ON HEMODIALYSIS
- U-3092 METHOD OF TREATING MOLYBDENUM COFACTOR DEFICIENCY TYPE A
- U-3093 IN COMBINATION WITH DEXAMETHASONE TO TREAT RELAPSED OR REFRACTORY MULTIPLE MYELOMA (REFRACTORY TO AT LEAST 1 PROTEASOME INHIBITOR, 1 IMMUNOMODULATORY AGENT, AND 1 ANTI-CD38 MAB) IN ADULTS WHO RECEIVED AT LEAST 4 PRIOR LINES OF THERAPY

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- U-3094 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD) WITH SERDEXMETHYLPHENIDATE AND DEXMETHYLPHENIDATE
- U-3095 TREATMENT OF HYPERLIPIDEMIA
- U-3096 TREATMENT OF ADULT PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- U-3097 EXTENDED ADJUVANT TREATMENT OF ADULT PATIENTS WITH EARLY-STAGE HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-POSITIVE BREAST CANCER, TO FOLLOW ADJUVANT TRASTUZUMAB BASED THERAPY
- U-3098 TREATMENT OF REFRACTORY EPILEPSY PATIENTS WITH FENFLURAMINE THAT REDUCES THE RISK OF CARDIOVASCULAR TOXICITY BY USING CARDIAC MONITORING AND RESTRICTED DISTRIBUTION
- U-3099 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH OBSTRUCTIVE SLEEP APNEA (OSA) IN A PATIENT WITH MODERATE RENAL IMPAIRMENT
- U-3100 A METHOD OF TREATING ADULTS WITH RELAPSED OR REFRACTORY ADVANCED RENAL CELL CARCINOMA FOLLOWING TWO OR MORE PRIOR SYSTEMIC THERAPIES BY INHIBITING THE ANGIOGENESIS OF BLOOD VESSELS WITH A VASCULAR ENDOTHELIAL GROWTH FACTOR INHIBITOR
- U-3101 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS WITH A SINGLE DOSE OF 1200MG ORITAVANCIN OR ITS SINGLE DOSE EQUIVALENT
- U-3102 REDUCTION OF CIRCULATING LYMPHOCYTES IN TREATING RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS
- U-3103 TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS), TO INCLUDE CLINICALLY ISOLATED SYNDROME, RELAPSING-REMITTING DISEASE, AND ACTIVE SECONDARY PROGRESSIVE DISEASE, IN ADULTS USING A DOSE TITRATION SCHEDULE FOLLOWED BY A MAINTENANCE DOSE
- U-3104 TREATMENT OF C. DIFFICILE-ASSOCIATED DIARRHEA
- U-3105 TREATMENT OF STAPHYLOCOCCAL ENTEROCOLITIS
- U-3106 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON TO SMOKERS OR TO PATIENTS BEING TREATED WITH A CYP1A2 INHIBITOR
- U-3107 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON TO SMOKERS OR TO PATIENTS BEING TREATED WITH A CYP1A2 INHIBITOR
- U-3108 TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS AGED 3 YEARS AND OLDER BY ADMINISTRATION OF AN EXTENDED-RELEASE SUSPENSION FORMULATION OF MIRABEGRON
- U-3109 METHOD OF USING VISMODEGIB TO TREAT BASAL CELL CARCINOMA
- U-3110 USE OF NALOXONE FOR THE EMERGENCY TREATMENT OF KNOWN OR SUSPECTED OPIOID OVERDOSE, AS MANIFESTED BY RESPIRATORY AND/OR CENTRAL NERVOUS SYSTEM DEPRESSION, FOR ADULT AND PEDIATRIC PATIENTS
- U-3111 TREATING OPIOID USE DISORDER
- U-3112 TREATING NEWLY DIAGNOSED ACUTE MYELOGENOUS LEUKEMIA (AML) CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3113 TREATING RELAPSED/REFRACTORY ACUTE MYELOGENOUS LEUKEMIA (AML) CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3114 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) BY ORALLY ADMINISTERING VENETOCLAX TO AN ADULT ACCORDING TO A DOSE RAMP-UP THAT INCLUDES A DOSE OF 50 MG PER DAY FOR 1 WEEK FOLLOWED BY 100 MG PER DAY FOR 1 WEEK
- U-3115 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING ABOUT 1 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS
- U-3116 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING ABOUT 1 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS

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U-3117 ADMINISTRATION TO THE EYE OF A PATIENT FOR TREATMENT OF DRY EYE CONDITION

U-3118 TREATMENT OF POSTSURGICAL PAIN PROVIDING ANALGESIA TO A PATIENT FOR UP TO 72 HOURS, FOR EXAMPLE, AFTER BUNIONECTOMY, OPEN INGUINAL HERNIORRHAPHY, OR TOTAL KNEE ARTHROPLASTY VIA SOFT TISSUE OR PERIARTICULAR INSTILLATION

U-3119 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 11

U-3120 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 11, AND 12

U-3121 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13 AND 23

U-3122 ADJUNCT TO DIET EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETIS BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 13, 23, AND 24

U-3123 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF COMPLEMENT INHIBITOR PEGCETACOPLAN

U-3124 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF PEGCETACOPLAN

U-3125 USE FOR LOADING DOSE IN PATIENTS WITH SYMPTOMATIC AFIB/AFL WHO ARE CURRENTLY IN SINUS RHYTHM OR FOR THE TREATMENT OF LIFE-THREATENING VENTRICULAR TACHYCARDIA

U-3126 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A SECOND AND FURTHER CARDIOVASCULAR EVENT IN AN ADULT PATIENT WITH ESTABLISHED CARDIOVASCULAR DISEASE

U-3127 REDUCTION OF THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH HEART FAILURE WITH REDUCED EJECTION FRACTION AND WITHOUT TYPE II DIABETES

U-3128 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, ENTEROBACTER CLOACAE SPECIES COMPLEX WITH MEROPENEM & VABORBACTAM AS SPECIFIED

U-3129 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS) IN PREMENOPAUSAL WOMEN

U-3130 METHOD OF POSITRON EMISSION TOMOGRAPHY (PET) IN MEN WITH PROSTATE CANCER

U-3131 USE OF ZUBSOLV FOR TREATMENT OF OPIOID DEPENDENCE

U-3132 INDICATED FOR THE TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS

U-3133 TREATING SICKLE CELL DISEASE BY ADMINISTERING 1500 MG OF VOXELOTOR ORALLY ONCE DAILY

U-3134 INCREASING HEMOGLOBIN TO TREAT SICKLE CELL DISEASE BY ADMINISTERING 1500 MG OF VOXELOTOR ORALLY ONCE DAILY

U-3135 TREATING SCHIZOPHRENIA

U-3136 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCED ANTIPSYCHOTIC INDUCED WEIGHT GAIN

U-3137 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCED ANTIPSYCHOTIC INDUCED WEIGHT GAIN

U-3138 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCTION OF THE ADVERSE METABOLIC PROFILE

U-3139 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN, WITH REDUCTION OF THE ADVERSE METABOLIC PROFILE

U-3140 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN

U-3141 METHOD OF TREATING BIPOLAR DISORDER BY ADMINISTERING THE ATYPICAL ANTIPSYCHOTIC OLANZAPINE AND SAMIDORPHAN

U-3142 PREVENTIVE TREATMENT OF EPISODIC MIGRAINE IN ADULTS

U-3143 FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE AND OLDER

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PATENT USE

- U-3144 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3145 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3146 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3147 FOR THE TREATMENT OF NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN ADULTS AND PEDIATRIC PATIENTS 1 YEAR AND OLDER
- U-3148 METHOD OF TREATING RELAPSING FORMS OF MULTIPLE SCLEROSIS BEFORE AND AFTER ADMINISTERING AN INACTIVE VACCINE
- U-3149 METHOD OF RECONSTITUTING A LYOPHILIZED LIPOSOMAL COMPOSITION FOR ADMINISTERING CYTARABINE AND DAUNORUBICIN TO TREAT NEWLY-DIAGNOSED THERAPY-RELATED AML (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PATIENTS 1 YEAR AND OLDER
- U-3150 METHOD OF ADMINISTERING A RECONSTITUTED LIPOSOMAL COMPOSITION CONTAINING CYTARABINE AND DAUNORUBICIN TO TREAT NEWLY-DIAGNOSED THERAPY-RELATED AML (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC) IN PATIENTS 1 YEAR AND OLDER
- U-3151 TO IMPROVE WAKEFULNESS IN ADULT PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY OR OBSTRUCTIVE SLEEP APNEA (OSA) WITH A DOSING REGIMEN THAT INCLUDES A DOSE OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3152 USE BY FEMALES OF REPRODUCTIVE POTENTIAL TO PREVENT PREGNANCY
- U-3153 TREATMENT OF ADVANCED GASTROINTESTINAL STROMAL TUMOR IN PATIENTS HAVING PROGRESSED FROM A FIRST LINE ADMINISTRATION OF IMATINIB, A SECOND LINE ADMINISTRATION OF SUNITINIB, AND A THIRD LINE ADMINISTRATION OF REGORAFENIB
- U-3154 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH A COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR; AND ANOTHER COMPOSITION COMPRISING IVACAFTOR
- U-3155 TREATMENT OF CF IN PATIENTS 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3156 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACAFTOR, TEZACAFTOR, AND IVACAFTOR
- U-3157 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-3158 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3159 TREATMENT OF ADULT AND POST-MENARCHAL PEDIATRIC FEMALES WITH VULVOVAGINAL CANDIDIASIS (VVC)
- U-3160 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WHO ARE SEVERELY IMMUNOCOMPROMISED
- U-3161 METHOD FOR WEIGHT MANAGEMENT ACCORDING TO A DOSE ESCALATION SCHEDULE
- U-3162 METHOD FOR WEIGHT MANAGEMENT
- U-3163 TREATMENT OF ACTINIC KERATOSES OF UPPER EXTREMITIES BY PHOTODYNAMIC THERAPY
- U-3164 GASTROINTESTINAL TABLETS INDICATED FOR CLEANSING THE COLON IN PREPARATION FOR COLONOSCOPY
- U-3165 METHOD OF TREATING HUMAN SMALLPOX DISEASE

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- U-3166 OTC USE: ALLERGY SYMPTOM RELIEVER; TEMPORARY RELIEF OF THESE SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES: NASAL CONGESTION, RUNNY NOSE, SNEEZING AND ITCHY NOSE
- U-3167 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, OR 6 IN PEDIATRIC PATIENTS 3 TO LESS THAN 12 YEARS OF AGE OR WEIGHING LESS THAN 45 KG
- U-3168 TREATMENT OF ADVANCED SYSTEMIC MASTOCYTOSIS, INCLUDING PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH AN ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), AND MAST CELL LEUKEMIA (MCL)
- U-3169 TREATMENT OF TRICHOMONIASIS IN ADULTS
- U-3170 TREATING CHRONIC HEART FAILURE WITH REDUCED EJECTION FRACTION IN PATIENTS NOT TAKING AN ACE INHIBITOR OR AN ARB OR PREVIOUSLY TAKING LOW DOSES OF THESE AGENTS, BY TITRATING UP FROM HALF THE USUALLY RECOMMENDED STARTING DOSE
- U-3171 TREATMENT OF INVASIVE ASPERGILLOSIS IN ADULTS AND PEDIATRIC PATIENTS 13 YEARS OF AGE AND OLDER
- U-3172 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN
- U-3173 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN TWICE WEEKLY
- U-3174 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF 1080 MG OF PEGCETACOPLAN EVERY THREE DAYS
- U-3175 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS BY ADMINISTERING THE FORMULATION OF DAPTOMYCIN AS RECITED IN CLAIM 18
- U-3176 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND S. AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS BY RECONSTITUTING AND ADMINISTERING THE FORMULATION AS RECITED IN CLAIM 12
- U-3177 TREATMENT OF VENOUS THROMBOTIC DISEASE
- U-3178 REDUCE THE RISK OF RECURRENCE OF VENOUS THROMBOTIC DISEASE
- U-3179 METHOD OF TREATING FOLLICULAR LYMPHOMA
- U-3180 DECREASING OFF TIME IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-3181 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE TABLET ACCORDING TO CLAIM 1 OF U.S. PATENT NO. 11,052,075, WHERE THE TABLET FURTHER COMPRISES IVACAFTOR
- U-3182 METHOD OF PROVIDING POSTSURGICAL PAIN MANAGEMENT, FOR EXAMPLE, VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3183 USE OF EPHEDRINE SULFATE FOR TREATING HYPOTENSION
- U-3184 MEKINIST(R) IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-3185 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-3186 METHOD OF TREATING PRURITUS IN PATIENTS 3 MONTHS OR OLDER SUFFERING FROM PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- U-3187 METHOD OF REDUCING SERUM BILE ACIDS IN PATIENTS 3 MONTHS OR OLDER SUFFERING FROM PROGRESSIVE FAMILIAL INTRAHEPATIC CHOLESTASIS (PFIC)
- U-3188 IMPROVING GLYCEMIC CONTROL IN PATIENTS 10 YEARS OF AGE AND OLDER WITH TYPE 2 DIABETES MELLITUS BY ADMINISTERING A SUSTAINED-RELEASE EXENATIDE FORMULATION AS AN ADJUNCT TO DIET AND EXERCISE
- U-3189 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS 10 TO 17 YEARS OF AGE WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH INSULIN ALONE OR INSULIN PLUS ONE OTHER ORAL ANTIDIABETIC MEDICATION

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- U-3190 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS 10 TO 17 YEARS OF AGE WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN AND/OR SULFONYLUREA
- U-3191 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (30 ML/MIN/1.73 M² ≤ EGFR < 60 ML/MIN/1.73 M²) BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-3192 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (EGFR < 60 ML/MIN/1.73 M²) BY INITIATION OF EMPAGLIFLOZIN, LINAGLIPTIN AND METFORMIN HCL IF EGFR ≥ 45 ML/MIN/1.73 M² AND DISCONTINUATION IF EGFR < 30 ML/MIN/1.73 M²
- U-3193 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (EGFR < 60 ML/MIN/1.73 M²) BY INITIATION OF EMPAGLIFLOZIN AND METFORMIN HCL IF EGFR ≥ 45 ML/MIN/1.73 M² AND DISCONTINUATION IF EGFR < 30 ML/MIN/1.73 M²
- U-3194 TOPICAL TREATMENT OF ACNE VULGARIS IN ADULTS AND PEDIATRIC PATIENTS 9 YEARS OF AGE AND OLDER
- U-3195 TREATMENT OF ADULTS WITH REFRACTORY, MODERATE-TO-SEVERE ATOPIC DERMATITIS NOT ADEQUATELY CONTROLLED WITH OTHER SYSTEMIC DRUG PRODUCTS OR WHEN USE OF THOSE THERAPIES IS INADVISABLE
- U-3196 TREATMENT OF ADULTS WITH LOCALLY ADVANCED OR METASTATIC UROTHELIAL CARCINOMA WITH SUSCEPTIBLE FGFR3 OR FGFR2 GENETIC ALTERATIONS, AND PROGRESSED DURING OR FOLLOWING PRIOR PLATINUM-CONTAINING CHEMOTHERAPY, WITH DOSING BASED ON SERUM PHOSPHATE LEVELS
- U-3197 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE AN OPIOID ANALGESIC AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
- U-3198 METHOD OF TREATING PATIENTS WITH IDIOPATHIC HYPERSOMNIA WITH SODIUM OXYBATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3199 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH PLUS HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS, HEART FAILURE AND REDUCED EJECTION FRACTION BY ONCE DAILY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3200 INCREASING SURVIVAL IN MCRPC PATIENTS PREVIOUSLY TREATED WITH DOCETAXEL BY ADMINISTERING CABAZITAXEL IN COMBINATION WITH PREDNISONE OR PREDNISOLONE AFTER A PREMEDICATION REGIMEN THAT INCLUDES AN ANTIHISTAMINE, A CORTICOSTEROID, AND AN H₂-ANTAGONIST
- U-3201 TREATMENT OF ADULT PATIENTS WITH VON HIPPEL-LINDAU DISEASE WHO REQUIRE THERAPY FOR ASSOCIATED RENAL CELL CARCINOMA, CENTRAL NERVOUS SYSTEM HEMANGIOBLASTOMAS, OR PANCREATIC NEUROENDOCRINE TUMORS, NOT REQUIRING IMMEDIATE SURGERY
- U-3202 MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) COMPRISING THE ONCE PER DAY ADMINISTRATION OF TRELEGY ELLIPTA, 100 MCG FLUTICASONE FUROATE/62.5 MCG UMECLIDINIUM/25 MCG VILANTEROL
- U-3203 MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-3204 TREATMENT OF MODERATE-TO-SEVERE PRURITUS ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD-AP) IN ADULTS UNDERGOING HEMODIALYSIS (HD)
- U-3205 REDUCTION OF RISK OF MAJOR CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION AND STROKE) IN PATIENTS WITH CAD
- U-3206 REDUCTION OF RISK OF MAJOR THROMBOTIC VASCULAR EVENTS (MYOCARDIAL INFARCTION, ISCHEMIC STROKE, ACUTE LIMB ISCHEMIA, AND MAJOR AMPUTATION OF VASCULAR ETIOLOGY) IN PATIENTS WITH PAD
- U-3207 REDUCTION OF RISK OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH CAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-3208 REDUCTION OF RISK OF MYOCARDIAL INFARCTION AND ISCHEMIC STROKE IN PATIENTS WITH PAD BY ADMINISTERING CLINICALLY PROVEN EFFECTIVE AMOUNTS THAT ARE 2.5 MG RIVAROXABAN TWICE DAILY AND 75-100 MG ASPIRIN DAILY
- U-3209 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK MYOCARDIAL INFARCTION IN AN ADULT PATIENT HAVING ATRIAL FIBRILLATION OR ATRIAL FLUTTER AND ELEVATED TRIGLYCERIDE LEVELS
- U-3210 ONCE DAILY TREATMENT OF ANXIETY DISORDER IN ADULTS
- U-3211 TREATING DISTRIBUTIVE SHOCK WITH ANGIOTENSIN II

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U-3212 TREATING SEPTIC SHOCK WITH ANGIOTENSIN II

U-3213 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHEREIN THE CANCER IS PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA

U-3214 A METHOD OF TREATING PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL

U-3215 A METHOD OF TREATING PREVIOUSLY TREATED, LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA CHARACTERIZED BY AN IDH1 MUTATION

U-3216 A METHOD FOR TREATING A BCRABL POSITIVE LEUKEMIA IN A SUBJECT THAT IS RESISTANT TO IMATINIB COMPRISING ADMINISTERING TO THE SUBJECT A THERAPEUTICALLY EFFECTIVE AMOUNT OF BOSUTINIB, WHEREIN THE SUBJECT HAS A MUTATION IN THE BCRABL PROTEIN AT 949T>C

U-3217 A METHOD FOR TREATING A BCRABL POSITIVE LEUKEMIA IN A SUBJECT THAT IS RESISTANT TO IMATINIB COMPRISING ADMINISTERING TO THE SUBJECT A THERAPEUTICALLY EFFECTIVE AMOUNT OF BOSUTINIB, WHEREIN THE SUBJECT HAS A MUTATION IN THE BCRABL PROTEIN AT F317L

U-3218 NASAL ADMINISTRATION OF DIHYDROERGOTAMINE MESYLATE BY METERED SPRAY FOR THE ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA

U-3219 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR

U-3220 TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 20 EXON INSERTION MUTATIONS WHOSE DISEASE HAS PROGRESSED ON OR AFTER PLATINUM-BASED CHEMOTHERAPY

U-3221 USE OF VASCEPA AS AN ADJUNCT TO STATIN THERAPY TO REDUCE THE RISK OF A CARDIOVASCULAR EVENT IN A PATIENT WITH PRIOR PERCUTANEOUS CORONARY INTERVENTION

U-3222 TREATMENT OF ACUTE MYELOID LEUKEMIA (AML) BY ORALLY ADMINISTERING VENETOCLAX WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE IN ADULTS 75 YEARS OR OLDER OR HAVING CERTAIN COMORBIDITIES ACCORDING TO A DOSE RAMP-UP INCLUDING A 100 MG PER DAY DOSE

U-3223 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) BY ORALLY ADMINISTERING VENETOCLAX TO AN ADULT ACCORDING TO A DOSE RAMP-UP INCLUDING A 100 MG PER DAY DOSE

U-3224 A METHOD OF TREATING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE BY DECREASING THE LEVEL OF LDL-C USING A FIXED DOSE COMBINATION OF 180 MG BEMPEDOIC ACID AND 10 MG EZETIMIBE

U-3225 TREATMENT OF DIFFERENTIATED THYROID CANCER THAT HAS PROGRESSED FOLLOWING PRIOR VEGFR-TARGETED THERAPY

U-3226 FOR TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR TWO LINES OF SYSTEMIC THERAPY

U-3227 FOR TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS (MF), INCLUDING PRIMARY MF, POST-POLYCYTHEMIA VERA MF AND POST-ESSENTIAL THROMBOCYTHEMIA MF

U-3228 FOR TREATMENT OF POLYCYTHEMIA VERA (PV) IN PATIENTS WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA

U-3229 FOR TOPICAL SHORT-TERM, NON-CONTINUOUS CHRONIC TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS IN NON-IMMUNOCOMPROMISED PATIENTS WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH TOPICAL PRESCRIPTION THERAPIES OR WHEN THOSE THERAPIES ARE NOT ADVISABLE

U-3230 FOR TREATMENT OF STEROID-REFRACTORY ACUTE GRAFT-VERSUS-HOST DISEASE (AGVHD)

U-3231 TREATMENT OF PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML) BY ADMINISTERING NILOTINIB DISPERSED IN A FRUIT PREPARATION

U-3232 USE OF ORAL OCTREOTIDE FOR LONG-TERM MAINTENANCE TREATMENT IN ACROMEGALY PATIENTS WHILE AVOIDING CONCOMITANT ADMINISTRATION OF LEVONORGESTREL

U-3233 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX IN PATIENTS TAKING EVEROLIMUS

U-3234 TREATMENT OF MACULAR EDEMA ASSOCIATED WITH UVEITIS

U-3235 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME

U-3236 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME

U-3237 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 IN

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- ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER OR WEIGHING AT LEAST 45 KG
- U-3238 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6 IN PEDIATRIC PATIENTS 3 TO LESS THAN 12 YEARS OF AGE OR WEIGHING LESS THAN 45 KG
- U-3239 TREATMENT OF ADVANCED RENAL CELL CARCINOMA (RCC) IN PATIENTS WHO HAVE RECEIVED PRIOR ANTI-ANGIOGENIC THERAPY
- U-3240 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT HAVING TRIGLYCERIDE LEVELS OF AT LEAST ABOUT 500 MG/DL, ON ANTICOAGULANT/ANTIPLATELET/THROMBOLYTIC THERAPY, AND HAVING ATRIAL FIBRILLATION AND/OR ATRIAL FLUTTER
- U-3241 IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE, NODE-POSITIVE, EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE AND A KI-67 SCORE $\geq 20\%$
- U-3242 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY
- U-3243 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN, AND MEN WITH HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3244 A METHOD FOR TREATMENT OF PAIN IN ADULTS USING TRAMADOL HYDROCHLORIDE AND CELECOXIB
- U-3245 MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR I DISORDER
- U-3246 FOR THE TREATMENT OF CHRONIC GRAFT VERSUS HOST DISEASE
- U-3247 FOR THE TREATMENT OF CHRONIC GRAFT VERSUS HOLD DISEASE
- U-3248 TREATING SECONDARY HYPERPARATHYROIDISM IN STAGE 3/4 CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL WHILE AVOIDING PTH OVERSUPPRESSION
- U-3249 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (AT LEAST 12 YEARS OF AGE AND 40 KG)
- U-3250 METHOD OF TREATING PAIN, FOR EXAMPLE, TREATING POSTSURGICAL PAIN VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3251 ADMINISTRATION OF FERROUS BISGLYCINATE TABLETS
- U-3252 USE OF VUITY FOR THE TREATMENT OF PRESBYOPIA IN ADULTS
- U-3253 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO AT LEAST ONE OF CLAIMS 1-9 OF US11179367
- U-3254 USE, IN COMBINATION WITH LOW-DOSE CYTARABINE, FOR THE TREATMENT OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULT PATIENTS WHO ARE ≥ 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- U-3255 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE RHEUMATOID ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3256 USE TO INCREASE LINEAR GROWTH IN PEDIATRIC PATIENTS WITH ACHONDROPLASIA WHO ARE 5 YEARS OF AGE AND OLDER WITH OPEN EPIPHYSES
- U-3257 TREATMENT OF TRD IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE IN A MAINTENANCE PHASE WEEKLY OR 1X EVERY TWO WEEKS TO ADULTS WHO HAVE BEEN ADMINISTERED ESKETAMINE IN A INDUCTION PHASE FOR ABOUT 4 WEEKS
- U-3258 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA) WITH A DOSE BETWEEN ABOUT 56 MG/M2 AND ABOUT 100 MG/M2 ADMINISTERED ON DAYS 1 AND 8 OF A 21-DAY CYCLE
- U-3259 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC MALIGNANT PERIVASCULAR EPITHELIOID CELL TUMOR (PECOMA)
- U-3260 METHOD OF TREATING SPASTICITY
- U-3261 FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY
- U-3262 TREATING HYPOTENSION WITH ANGIOTENSIN II IN A PATIENT RECEIVING AN ANGIOTENSIN CONVERTING ENZYME INHIBITOR

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- U-3263 METHOD FOR TREATING SPASTICITY
- U-3264 AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3265 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3266 IN COMBINATION WITH FULVESTRANT AS INITIAL ENDOCRINE-BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY IN POSTMENOPAUSAL WOMEN OR IN MEN, FOR THE TREATMENT OF HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-3267 USE OF LASTACFT TO TEMPORARY RELIEVE ITCHY EYES DUE TO POLLEN, RAGWEED, GRASS, ANIMAL HAIR AND DANDER
- U-3268 TREATMENT OF MULTIPLE SCLEROSIS IN PEDIATRIC PATIENTS 10 YEARS OF AGE AND OLDER AND WEIGHING LESS THAN OR EQUAL TO 40 KG
- U-3269 TREATMENT OF PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) IN ADULTS AT RISK OF RAPID DISEASE PROGRESSION
- U-3270 TREATMENT OF BACTERIAL VAGINOSIS IN FEMALE PATIENTS 12 YEARS OF AGE AND OLDER
- U-3271 TREATMENT OF BIPOLAR DEPRESSION
- U-3272 AS AN ADJUNCT TO DIET AND MAXIMALLY TOLERATED STATIN THERAPY FOR THE TREATMENT OF ADULTS WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH) OR CLINICAL ATHEROSCLEROTIC CARDIOVASCULAR DISEASE (ASCVD), BY INHIBITING EXPRESSION OF THE PCSK9 GENE
- U-3274 TREATMENT OF BIPOLAR I DISORDER, BIPOLAR II DISORDER, OR BIPOLAR DEPRESSION
- U-3275 TREATMENT OF ADULTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3276 TREATMENT OF PATIENTS WITH PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY
- U-3277 USE FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH TUBEROUS SCLEROSIS COMPLEX
- U-3278 A METHOD OF REDUCING POST-SURGICAL PAIN FOLLOWING OCULAR SURGERY
- U-3279 A METHOD OF TREATING POSTOPERATIVE INFLAMMATION FOLLOWING OCULAR SURGERY
- U-3280 METHOD OF TREATING ACNE VULGARIS WITH TOPICALLY APPLIED CORTEXOLONE 17ALPHA-PROPIONATE
- U-3281 TREATMENT OF SCHIZOPHRENIA IN ADULTS AND PEDIATRIC PATIENTS AGES 13 YEARS AND OLDER
- U-3282 DURING LEVOKETOCONAZOLE DOSAGE TITRATION FOR THE TREATMENT OF CUSHING'S SYNDROME IN PATIENTS WHO CONCOMITANTLY USE METFORMIN, MONITORING GLYCEMIA, KIDNEY FUNCTION AND VITAMIN B-12 AND ADJUSTING DOSAGE OF METFORMIN AS NEEDED
- U-3283 TREATMENT OF ENDOGENOUS HYPERCORTISOLEMIA IN PATIENTS WITH CUSHING'S SYNDROME FOR WHOM SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- U-3284 PROPHYLAXIS OF THROMBOEMBOLIC DISEASES IN PEDIATRIC PATIENTS AGED 2 YEARS AND OLDER WITH CONGENITAL HEART DISEASE WHO HAVE UNDERGONE THE FONTAN PROCEDURE AND A BODY WEIGHT OF ≥ 50 KG
- U-3285 TREATMENT OF DVT AND/OR PE AND REDUCTION IN THE RISK OF RECURRENT DVT AND/OR PE IN PEDIATRIC PATIENTS (≥ 50 KG) ONCE DAILY WITH RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS AFTER AT LEAST 5 DAYS PARENTERAL ANTICOAGULANT TREATMENT
- U-3286 TREATMENT OF VENOUS THROMBOEMBOLISM (VTE) AND THE REDUCTION IN THE RISK OF RECURRENT VTE IN PEDIATRIC PATIENTS FROM BIRTH TO LESS THAN 18 YEARS WITH A BODY WEIGHT OF 30 KG TO 49.9 KG AFTER AT LEAST 5 DAYS OF INITIAL PARENTERAL ANTICOAGULANT TREATMENT
- U-3287 TREATMENT OF VENOUS THROMBOEMBOLISM (VTE) AND THE REDUCTION IN THE RISK OF RECURRENT VTE IN PEDIATRIC PATIENTS FROM BIRTH TO LESS THAN 18 YEARS WITH A BODY WEIGHT OF ≥ 50 KG AFTER AT LEAST 5 DAYS OF INITIAL PARENTERAL ANTICOAGULANT TREATMENT
- U-3288 PROPHYLAXIS OF PE, DVT AND/OR STROKE IN PEDIATRIC PATIENTS (≥ 50 KG) AGED 2 YEARS AND OLDER WITH CONGENITAL HEART DISEASE AFTER FONTAN PROCEDURE WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS

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- U-3289 TREATMENT OF DVT AND/OR PE AND REDUCTION IN RISK OF RECURRENT DVT AND/OR PE IN PEDIATRIC PATIENTS (30-49.9 KG) ONCE DAILY WITH RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS AFTER AT LEAST 5 DAYS PARENTERAL ANTICOAGULANT TREATMENT
- U-3290 TREATMENT OF CHOLESTATIC PRURITUS IN PATIENTS WITH ALAGILLE SYNDROME (ALGS)
- U-3291 CYTALUX IS AN OPTICAL IMAGING AGENT INDICATED IN ADULT PATIENTS WITH OVARIAN CANCER AS AN ADJUNCT FOR INTRAOPERATIVE IDENTIFICATION OF MALIGNANT LESIONS
- U-3292 USE OF VASCEPA TO REDUCE TRIGLYCERIDE LEVELS IN AN ADULT PATIENT ON STATIN THERAPY AND HAVING ATRIAL FIBRILLATION OR ATRIAL FLUTTER AND TRIGLYCERIDE LEVELS OF ABOUT 500 MG/DL TO ABOUT 2,000 MG/DL
- U-3293 METHOD OF TREATING BACTERIAL VAGINOSIS BY SINGLE DOSE ADMINISTRATION OF A CLINDAMYCIN PHARMACEUTICAL GEL FORMULATION
- U-3294 METHOD OF TREATING A BACTERIAL INFECTION BY ADMINISTERING A RECONSTITUTED SOLID FORMULATION OF DAPTOMYCIN CONTAINING 31.0 TO 59.4% WT TOTAL MANNITOL AND SORBITOL
- U-3295 METHOD OF DELIVERING A COMBINATION OF OLOPATADINE HYDROCHLORIDE AND MOMETASONE FUROATE TO A NASAL AIRWAY
- U-3296 TREATMENT OF SEASONAL ALLERGIC RHINITIS BY NASALY ADMINISTERING A COMBINATION OF OLOPATADINE HYDROCHLORIDE AND MOMETASONE FUROATE TO A PEDIATRIC PATIENT
- U-3297 TREATMENT OF SEASONAL ALLERGIC RHINITIS BY NASALY ADMINISTERING A COMBINATION OF OLOPATADINE HYDROCHLORIDE AND MOMETASONE FUROATE
- U-3298 TREATMENT OF ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH REFRACTORY, MODERATE TO SEVERE ATOPIC DERMATITIS WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH OTHER SYSTEMIC DRUG PRODUCTS, OR WHEN USE OF THOSE THERAPIES ARE INADVISABLE
- U-3299 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN PEDIATRIC PATIENTS
- U-3300 PROPHYLAXIS TO PREVENT ATTACKS OF HEREDITARY ANGIOEDEMA (HAE) IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3301 TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH HEPATIC VENO-OCCLUSIVE DISEASE (VOD), ALSO KNOWN AS SINUSOIDAL OBSTRUCTION SYNDROME (SOS), WITH RENAL OR PULMONARY DYSFUNCTION FOLLOWING HEMATOPOIETIC STEM-CELL TRANSPLANTATION (HSCT)
- U-3302 TREATMENT OF TRICHOMONIASIS IN PATIENTS 12 YEARS OF AGE AND OLDER
- U-3303 TOPICAL LESION-DIRECTED AND FIELD-DIRECTED TREATMENT OF ACTINIC KERATOSIS OF THE FACE AND SCALP WITH PHOTODYNAMIC THERAPY BY POSITIONING AN ILLUMINATION DEVICE IN AN APPROPRIATE DISTANCE AND ILLUMINATING THE TREATMENT AREA WITH NARROWBAND RED LIGHT
- U-3304 METHOD OF TREATING PARKINSON'S DISEASE BY ORALLY ADMINISTERING SEGMENTS OF A FUNCTIONALLY MULTISCORED, BILAYERED TABLET HAVING CARBIDOPA-25 MG/LEVODOPA-100 MG, EACH SEGMENT HAVING CARBIDOPA-6.25 MG/LEVODOPA-25 MG
- U-3305 METHOD OF TREATING PARKINSON'S DISEASE BY ORALLY ADMINISTERING A FUNCTIONALLY MULTISCORED, BILAYERED TABLET HAVING CARBIDOPA-25 MG/LEVODOPA-100 MG
- U-3306 TREATMENT OF ADULT PATIENTS WITH KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), AS DETERMINED BY AN FDA-APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-3307 TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 35KG WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED
- U-3308 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS TO TREAT HIV-1 INFECTION IN PEDIATRIC PATIENTS WEIGHING AT LEAST 35KG WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR TO REPLACE THE CURRENT ANTIRETROVIRAL REGIMEN IF VIROLOGICALLY SUPPRESSED
- U-3309 NORLIQVA IS INDICATED FOR THE TREATMENT OF HYPERTENSION, TO LOWER BLOOD PRESSURE IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-3310 NORLIQVA IS INDICATED FOR THE SYMPTOMATIC TREATMENT OF CHRONIC STABLE ANGINA
- U-3311 NORLIQVA IS INDICATED FOR THE TREATMENT OF CONFIRMED OR SUSPECTED VASOSPASTIC ANGINA
- U-3312 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 40 KG BY

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- INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-3313 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 GRAMS OF ELEMENTAL IRON
- U-3314 METHOD OF TREATING IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER HAVING INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON ASSOCIATED WITH HEAVY UTERINE BLEEDING OR GASTROINTESTINAL DISORDER BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE
- U-3315 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 40 KG BY ADMINISTERING IV AT LEAST ABOUT 0.6 G OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MIN OR LESS
- U-3316 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING AT LEAST ABOUT 0.6 GRAMS OF IRON AS FERRIC CARBOXYMALTOSE IN ABOUT 15 MINUTES OR LESS
- U-3317 POSITRON EMISSION TOMOGRAPHY (PET) OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA) POSITIVE LESIONS IN MEN WITH PROSTATE CANCER
- U-3318 MANAGEMENT OF MODERATE-TO-SEVERE PAIN BY INJECTION
- U-3319 METHOD OF USING A PYRUVATE KINASE ACTIVATOR FOR THE TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY
- U-3320 METHOD FOR INCREASING THE LIFETIME OF RED BLOOD CELLS (RBCS) FOR THE TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY
- U-3321 METHOD OF USING A PYRUVATE KINASE ACTIVATOR FOR THE TREATMENT OF HEMOLYTIC ANEMIA IN ADULTS WITH PYRUVATE KINASE (PK) DEFICIENCY BY ADMINISTERING A DAILY DOSE OF 10MG TO 100MG
- U-3322 USE FOR DETECTING NEUTRALIZING ANTIBODIES
- U-3323 METHOD OF REDUCING ADVERSE EFFECTS IN PATIENTS WHO ARE CONCOMITANTLY ADMINISTERED A SALT OF GAMMA-HYDROXYBUTYRATE AND DIVALPROEX SODIUM
- U-3324 METHOD OF TREATING PATIENTS WITH A SALT OF GAMMA-HYDROXYBUTYRATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED
- U-3325 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND HEART FAILURE BY ONCE DAILY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3326 A METHOD OF TREATING ANKYLOSING SPONDYLITIS BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 8, 15-20, 27-31, 34-43, 45, 47, 49, 50, 54, 59, 63, 68-71, 73, 77, 82-84, AND 87-98
- U-3327 A METHOD OF TREATING PSORIATIC ARTHRITIS BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 8, 15-20, 27-31, 34-44, 46, 48, 50, 53, 59, 62, 68-71, 73, 76, 82-84, AND 87-98
- U-3328 A METHOD OF TREATING RHEUMATOID ARTHRITIS BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 2, 8, 9, 15-21, 27-31, 34-43, 50, 51, 59, 60, 68-71, 73, 74, 82-84 AND 87-98
- U-3329 A METHOD OF TREATING ULCERATIVE COLITIS, BY ADMINISTERING THE FORMULATION OF TOFACITINIB OF CLAIMS 1, 3, 8, 10, 15-20, 22, 27-31, 34-43, 50, 52, 59, 61, 68-71, 73, 75, 82-84 AND 87-98
- U-3330 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM PALMER-PLANTER ERYTHRODYSESTHESIA SYNDROME
- U-3331 TREATMENT OF MYELOFIBROSIS WITH PACRITINIB
- U-3332 USE OF PACRITINIB FOR INHIBITING JANUS ASSOCIATED KINASE 2 (JAK2)
- U-3333 ADJUVANT TREATMENT OF PATIENTS WITH GBRCA-MUTATED HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE HIGH RISK EARLY BREAST CANCER WHO HAVE BEEN TREATED WITH NEOADJUVANT OR ADJUVANT CHEMOTHERAPY
- U-3334 A METHOD OF TRANSDERMAL DELIVERY OF DONEPEZIL FOR TREATING MILD, MODERATE AND SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE
- U-3335 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA (HABP)
- U-3336 TREATMENT OF VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (VABP)
- U-3337 ADMINISTERING DAILY A UNIT DOSAGE OF AN IRREVERSIBLE EGFR INHIBITOR COVALENTLY

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- BINDING AS CLAIMED FOR 1ST LINE TREATMENT OF GEFITINIB OF ERLOTINIB RESISTANT METASTATIC NSCLC WITH EGFR EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION
- U-3338 ADMINISTERING DAILY A UNIT DOSAGE OF AN IRREVERSIBLE EGFR INHIBITOR COVALENTLY BINDING AS CLAIMED FOR 1ST LINE TREATMENT OF GEFITINIB OR ERLOTINIB RESISTANT METASTATIC NSCLC WITH EGFR EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION WITH T790M MUTATION
- U-3339 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11147770
- U-3340 A METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER BY ADMINISTERING A TRANSDERMAL COMPOSITION CONTAINING AMPHETAMINE
- U-3341 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3342 TREATMENT OF NIGHTTIME SLEEP DISTURBANCES IN SMITH-MAGENIS SYNDROME BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH BETA-ADRENERGIC RECEPTOR ANTAGONISTS
- U-3343 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH BETA-ADRENERGIC RECEPTOR ANTAGONISTS
- U-3344 A METHOD OF TREATING ADULTS WITH MULTIPLE MYELOMA USING DEXAMETHASONE IN COMBINATION WITH AN ANTI-MYELOMA PRODUCT
- U-3345 FOR TREATMENT OF ADULT PATIENTS WITH PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA)-POSITIVE METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC) WHO HAVE BEEN TREATED WITH ANDROGEN RECEPTOR (AR) PATHWAY INHIBITION AND TAXANE-BASED CHEMOTHERAPY
- U-3346 METHOD OF PROVIDING LOCAL OR REGIONAL ANALGESIA VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK OR FEMORAL NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3347 USE IN COMBINATION WITH CANNABIDIOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME
- U-3348 TREATMENT OF HIV-1 INFECTION IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 35 KG
- U-3349 REINITIATION OF SCHIZOPHRENIA TREATMENT WHEREIN MORE THAN 6 MONTHS 3 WEEKS BUT LESS THAN 8 MONTHS HAVE ELAPSED SINCE THE LAST DOSE
- U-3350 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER BY SUBLINGUAL ADMINISTRATION
- U-3351 METHOD OF TREATING IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSATE
- U-3352 METHOD TO TREAT IRON DEFICIENCY ANEMIA IN ADULTS & PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSATE
- U-3353 TREATMENT IN COMBINATION WITH CABOTEGRAVIR OF HIV-1 INFECTION IN ADULTS AND ADOLESCENTS 12 AND OLDER TO REPLACE CURRENT REGIMEN IN THOSE WHO ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN WITH NO HISTORY OF TREATMENT FAILURE
- U-3354 TREATMENT OF ADULT PATIENTS WITH PAROXYSMAL NOCTURNAL HEMOGLOBINURIA (PNH) BY ADMINISTRATION OF PEGCETACOPLAN SO AS TO REDUCE THE SENSITIVITY OF CELLS TO COMPLEMENT-DEPENDENT DAMAGE
- U-3355 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-3356 TOPICAL TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA IN ADULTS 18 YEARS OF AGE AND OLDER
- U-3357 TOPICAL TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA IN ADULTS 65 YEARS OF AGE AND OLDER
- U-3358 USE OF VASCEPA TO REDUCE THE INCIDENCE OF MI IN AN ADULT PATIENT ON STATIN THERAPY AND WITH ELEVATED TRIGLYCERIDE LEVELS (>150 MG/DL), WHEREIN THE PATIENT EXPERIENCES ATRIAL FIBRILLATION AND/OR FLUTTER INSTEAD OF AN INCIDENCE OF MI
- U-3359 TREATMENT OF SCHIZOPHRENIA BY ADMINISTERING A DOSE UP TO TWO WEEKS BEFORE OR THREE WEEKS AFTER THE SCHEDULED SIX-MONTH DOSE
- U-3360 COMPLICATED INTRA-ABDOMINAL INFECTIONS (CIAI), USED IN COMBINATION WITH

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- METRONIDAZOLE, IN ADULT AND PEDIATRIC PATIENTS (BIRTH TO LESS THAN 18 YEARS OF AGE)
- U-3361 COMPLICATED URINARY TRACT INFECTIONS (CUTI), INCLUDING PYELONEPHRITIS, IN ADULT AND PEDIATRIC PATIENTS (BIRTH TO LESS THAN 18 YEARS OLD)
- U-3362 TREATMENT OF BIPOLAR DEPRESSION MEDIATED BY THE 5-HT2A RECEPTOR, SEROTONIN TRANSPORTER, AND/OR DOPAMINE D1/D2 SIGNALING PATHWAYS
- U-3363 TREATMENT OF SCHIZOPHRENIA MEDIATED BY THE 5-HT2A RECEPTOR, SEROTONIN TRANSPORTER, AND/OR DOPAMINE D1/D2 SIGNALING PATHWAYS
- U-3364 TREATMENT OF BIPOLAR DISORDER I, BIPOLAR DISORDER II, OR BIPOLAR DEPRESSION
- U-3365 THE PRODUCT COMPOSITION (NATROBA) IS FOR THE TOPICAL TREATMENT OF HUMAN SCABIES MITE INFESTATIONS BY MELTING AND DELIVERING THE ACTIVE INGREDIENT, SPINOSAD, TO THE STRATUM CORNEUM WHERE SCABIES MITES LIVE AND BREED
- U-3366 VIVJOA IS INDICATED TO REDUCE THE INCIDENCE OF RECURRENT VULVOVAGINAL CANDIDIASIS (RVVC) IN FEMALES WITH A HISTORY OF RVVC WHO ARE NOT OF REPRODUCTIVE POTENTIAL
- U-3367 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG) REQUIRING HOSPITALIZATION
- U-3368 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG)
- U-3369 TREATMENT OF ADULTS AND PEDIATRIC PATIENTS 12 AND OLDER WITH SCLERODERMATOUS FORM OF CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF AT LEAST TWO PRIOR LINES OF SYSTEMIC THERAPY
- U-3370 A METHOD FOR THE TREATMENT OF ADULT PATIENTS WITH STABLE WILSON'S DISEASE WHO ARE DE-COPPERED AND TOLERANT TO PENICILLAMINE
- U-3371 TREATMENT OF ADULTS WITH ACTIVE ANKYLOSING SPONDYLITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3372 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN HOSPITALIZED ADULTS REQUIRING SUPPLEMENTAL OXYGEN, NON-INVASIVE OR INVASIVE MECHANICAL VENTILATION, OR EXTRACORPOREAL MEMBRANE OXYGENATION (ECMO)
- U-3373 TREATMENT OF ADULTS WITH SYMPTOMATIC NEW YORK HEART ASSOCIATION (NYHA) CLASS II-III OBSTRUCTIVE HYPERTROPHIC CARDIOMYOPATHY (HCM) TO IMPROVE FUNCTIONAL CAPACITY AND SYMPTOMS
- U-3374 TREATMENT OF SEIZURES ASSOCIATED WITH CYCLIN-DEPENDENT KINASE-LIKE 5 (CDKL5) DEFICIENCY DISORDER (CDD) IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-3375 USE FOR THE TREATMENT OF ABSENCE SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-3376 USE FOR THE TREATMENT OF ABSENCE SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME
- U-3377 TPOXX IS INDICATED FOR THE TREATMENT OF HUMAN SMALLPOX DISEASE IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 3 KG
- U-3378 MOUNJARO IS INDICATED AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-3379 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF EPINEPHRINE
- U-3380 METHOD OF TREATING PAIN, FOR EXAMPLE, VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK FOR REGIONAL ANALGESIA
- U-3381 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM HYPERTENSION
- U-3382 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM A GRADE 3 ADVERSE REACTION WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3383 TREATING NEWLY DIAGNOSED AML CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE OF IVOSIDENIB TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL, IN COMBINATION WITH AZACITIDINE
- U-3384 A METHOD FOR TREATING AML BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING IVOSIDENIB WHEREIN THE AML IS NEWLY DIAGNOSED AND CHARACTERIZED BY A MUTANT IDH1 AND THE COMPOSITION IS ADMINISTERED IN COMBINATION WITH AZACITIDINE
- U-3385 A METHOD FOR TREATING NEWLY DIAGNOSED AML WITH IVOSIDENIB AND AZACITIDINE WHEREIN THE AML HAS AN IDH1 MUTATION CAPABLE OF CONVERTING ALPHA-KETOGLUTARATE TO 2-HYDROXYGLUTARATE (2HG)
- U-3386 A METHOD FOR TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WITH IVOSIDENIB

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- IN COMBINATION WITH AZACITIDINE WHEREIN THE CANCER IS NEWLY DIAGNOSED AML
- U-3387 A METHOD FOR TREATING NEWLY DIAGNOSED AML CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 SELECTED FROM R132H, R132C, R132L, R132V, R132S AND R132GF BY ADMINISTERING IVOSIDENIB AND AZACITIDINE
- U-3388 USE OF ELAGOLIX 200 MG BID FOR 6 MONTHS TO MANAGE MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS IN PREMENOPAUSAL WOMEN TO REDUCE DYSMENORRHEA AND NON-MENSTRUAL PELVIC PAIN
- U-3389 USE OF ELAGOLIX 200 MG BID FOR 6 MONTHS TO MANAGE MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS IN PREMENOPAUSAL WOMEN HAVING DYSpareunia ASSOCIATED WITH ENDOMETRIOSIS
- U-3390 SODIUM THIOSULFATE INJECTION IS ADMINISTERED BY INTRAVENOUS INJECTION
- U-3391 A METHOD OF TITRATING AN OPIOID TO MANAGE PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
- U-3392 A METHOD OF TITRATING AN OPIOID TO MANAGE NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE
- U-3393 METHOD OF TREATING ACUTE PAIN DUE TO MINOR STRAINS, SPRAINS, AND CONTUSIONS USING A DICLOFENAC PATCH CONTAINING HEPARIN FOR ONCE DAILY ADMINISTRATION WHERE HEPARIN IS NOT RELEASED
- U-3394 SODIUM NITRITE INJECTION IS ADMINISTERED BY INTRAVENOUS INJECTION
- U-3395 SODIUM NITRITE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM THIOSULFATE FOR THE TREATMENT OF ACUTE CYANIDE POISONING
- U-3396 AMVUTTRA IS INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- U-3397 TADLIQ IS INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) TO IMPROVE EXERCISE ABILITY
- U-3398 FOR CHRONIC WEIGHT MANAGEMENT IN ADULTS WITH BMI ≥ 30 KG/M² OR BMI ≥ 27 KG/M² WITH A WEIGHT-RELATED COMORBIDITY, AND PATIENTS AGE 12-17 WITH BMI ≥ 25 KG/M² IN THE 95TH PERCENTILE OR GREATER (STANDARDIZED FOR AGE AND SEX)
- U-3399 FOR CHRONIC WEIGHT MANAGEMENT IN ADULTS WITH BMI ≥ 30 KG/M², AND PATIENTS AGE 12-17 WITH BMI ≥ 30 KG/M² AND IN THE 95TH PERCENTILE OR GREATER (STANDARDIZED FOR AGE AND SEX), EACH HAVING A WEIGHT-RELATED COMORBIDITY
- U-3400 FOR USE AFTER RADIOLABELING WITH GALLIUM-68, FOR POSITRON EMISSION TOMOGRAPHY OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA)-POSITIVE LESIONS IN MEN WITH PROSTATE CANCER
- U-3401 CAROSPIR IS INDICATED AS AN ADD-ON THERAPY FOR THE TREATMENT OF HYPERTENSION, TO LOWER BLOOD PRESSURE IN ADULT PATIENTS WHO ARE NOT ADEQUATELY CONTROLLED ON OTHER AGENTS
- U-3402 CAROSPIR IS INDICATED FOR THE MANAGEMENT OF EDEMA IN ADULT CIRRHOTIC PATIENTS WHEN EDEMA IS NOT RESPONSIVE TO FLUID AND SODIUM RESTRICTION
- U-3403 METHOD OF REVERSING OR INHIBITING THE PROGRESS OF UNRESECTABLE, RECURRENT, OR REFRACTORY INFLAMMATORY MYOFIBROBLASTIC TUMOR (IMT) THAT IS ALK-POSITIVE IN ADULT AND PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER
- U-3404 FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3405 TREATMENT OF HIV-1 INFECTION IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER AND WEIGHING AT LEAST 35 KG BY MONTHLY ADMINISTRATION OF RILPIVIRINE SUSPENSION AS PART OF COMBINATION THERAPY
- U-3406 USE OF FENFLURAMINE AT REDUCED AMOUNTS WITH STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME
- U-3407 USE OF FENFLURAMINE AT REDUCED AMOUNTS WITH STIRIPENTOL FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX GASTAUT SYNDROME
- U-3408 TOPICAL TREATMENT OF PLAQUE PSORIASIS, INCLUDING INTERTRIGINOUS AREAS, IN PATIENTS 12 YEARS OF AGE AND OLDER. (1)
- U-3409 TREATING ADULT PATIENTS WITH INTERMEDIATE-2 OR HIGH-RISK PRIMARY OR SECONDARY MYELOFIBROSIS, MONITORING THIAMINE LEVELS AND ADMINISTERING THIAMINE OR A THIAMINE EQUIVALENT

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- U-3410 A METHOD OF INJECTING AN IMPLANT
- U-3411 TREATING NON-EARLY SPMS BY ADMINISTERING ORAL CLADRIBINE AT A FIXED DOSE PER PATIENT, PER BODY WEIGHT AND PER TREATMENT YEAR, WHICH FIXED DOSE IS 1.75 +/- 0.2 MG/KG, TO BE ADMINISTERED WITHIN MONTHS 1 AND 2 IN EACH OF 2 ADJACENT TREATMENT YEARS
- U-3412 TREATMENT OF ADULTS WITH RELAPSED, REFRACTORY OR PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA INCLUDING A DOSE RAMP-UP AND IN COMBINATION WITH OBINUTUZUMAB IN MULTIPLE 28-DAY DOSING CYCLES FOLLOWED BY ADMINISTRATION IN ABSENCE OF OBINUTUZUMAB
- U-3413 INDICATED AS ADJUNCTIVE THERAPY FOR THE TREATMENT OF PARTIAL-ONSET SEIZURES, PRIMARY GENERALIZED TONIC-CLONIC SEIZURES, AND SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-3414 INDICATED AS INITIAL MONOTHERAPY FOR THE TREATMENT OF PARTIAL-ONSET OR PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-3415 INDICATED FOR THE PREVENTIVE TREATMENT OF MIGRAINE IN PATIENTS 12 YEARS AND OLDER
- U-3416 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS BEING TREATED CONCURRENTLY WITH A CYP3A4 INHIBITOR
- U-3417 TREATMENT OF POSTSURGICAL PAIN PROVIDING ANALGESIA TO A PATIENT FOR UP TO 72 HOURS, FOR EXAMPLE, AFTER FOOT AND ANKLE, SMALL-TO-MEDIUM OPEN ABDOMINAL, AND LOWER EXTREMITY TOTAL JOINT ARTHROPLASTY SURGICAL PROCEDURES VIA INSTILLATION
- U-3418 A METHOD OF LOADING MEDICATION INTO A SYRINGE AND DELIVERING THE MEDICATION TO A TREATMENT SITE
- U-3419 DEXTROMETHORPHAN AND BUPROPION IN COMBINATION TO TREAT MAJOR DEPRESSIVE DISORDER
- U-3420 A METHOD OF TREATING TESTOSTERONE DEFICIENCY IN MEN
- U-3421 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS (CUTI) INCLUDING PYELONEPHRITIS CAUSED BY THE FOLLOWING SUSCEPTIBLE MICROORGANISMS: ESCHERICHIA COLI, KLEBSIELLA PNEUMONIA, ENTEROBACTER CLOACAE SPECIES COMPLEX WITH MEROPENEM & VABORBACTAM AS SPECIFIED
- U-3422 TREATMENT OF PEDIATRIC PATIENTS AGE 1 YEAR AND OLDER WITH CHRONIC GRAFT-VERSUS-HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- U-3423 METHOD OF TREATING GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM GRADE 2 OR GRADE 3 MYALGIA WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3424 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND IVACAFTOR
- U-3425 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOUND OF CLAIM 1 OR COMPOSITION OF CLAIM 29 OF US11426407
- U-3426 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 8716338 AND IVACAFTOR
- U-3427 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR FORM I AND IVACAFTOR
- U-3428 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AND IVACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 9192606
- U-3429 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-3430 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 1-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING A PHARMACEUTICAL COMPOSITION ACCORDING TO CLAIM 2 OF U.S. PATENT NO. 10,597,384, FURTHER COMPRISING IVACAFTOR
- U-3431 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS WHO AWAKEN AT LEAST 2 TIMES PER NIGHT TO VOID BY INDUCING AN ANTIDIURETIC EFFECT BY INTRANASALLY ADMINISTERING A PLUME OF DROPLETS COMPRISING A DOSE OF ABOUT 0.05-5 MCG DESMOPRESSIN
- U-3432 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS (EDS) IN PATIENTS 7 YEARS OF AGE AND OLDER WITH NARCOLEPSY WITH A MIXTURE OF SODIUM, POTASSIUM, MAGNESIUM, AND CALCIUM SALTS OF GHB ADMINISTERED BETWEEN 2 AND 4 HOURS AFTER EATING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3433 INDICATED FOR THE TREATMENT OF SPASTICITY RESULTING FROM MULTIPLE SCLEROSIS
- U-3434 TREATMENT OF MODERATE-TO-SEVERE PLAQUE PSORIASIS IN ADULTS WHO ARE CANDIDATES FOR SYSTEMIC THERAPY OR PHOTOTHERAPY
- U-3435 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 46.7 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN 15 MINUTES
- U-3436 METHOD OF TREATING IDA IN ADULT PATIENTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 46.7 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN LESS THAN 15 MINUTES
- U-3437 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YEAR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & HEAVY UTERINE BLEEDING OR GI DISORDER BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO GIVE AT LEAST 0.7 G OF IRON IN 15 MINUTES
- U-3438 METHOD TO TREAT IDA IN ADULTS & PEDIATRIC PATIENTS 1 YR & OLDER WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 46.7 KG BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN < 15 MINUTES
- U-3439 METHOD OF TREATING PAIN, FOR EXAMPLE, VIA INFILTRATION FOR LOCAL ANALGESIA OR VIA NERVE BLOCK, FOR EXAMPLE, NTERSCALENE BRACHIAL PLEXUS FOR REGIONAL ANALGESIA
- U-3440 A METHOD OF ADMINISTERING APREPITANT FOR PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING
- U-3441 A METHOD OF TREATING CYSTINURIA BY ORALLY ADMINISTERING TIOPRONIN WITH FOOD TO PREVENT CYSTINE STONE FORMATION IN ADULTS AND PEDIATRIC PATIENTS WITH SEVERE HOMOZYGOUS CYSTINURIA
- U-3442 TREATMENT OF ADVANCED GASTROINTESTINAL STROMAL TUMOR IN PATIENTS PREVIOUSLY ADMINISTERED AT LEAST THREE TYROSINE KINASE INHIBITORS, WHERE ONE OF THE KINASE INHIBITORS IS IMATINIB
- U-3443 A METHOD OF REDUCING OTOTOXICITY IN A HUMAN PEDIATRIC PATIENT ABOUT 5 YEARS OF AGE OR UNDER WITH LOCALIZED MEDULLOBLASTOMA COMPRISING ADMINISTERING SODIUM THIOSULFATE ABOUT SIX HOURS AFTER ADMINISTRATION OF CISPLATIN
- U-3444 TREATMENT OF DEPRESSIVE SYMPTOMS IN ADULTS WITH MDD WITH ACUTE SUICIDAL IDEATION OR BEHAVIOR BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE AS A PHARMACEUTICAL COMPOSITION TWICE PER WEEK IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3445 TREATMENT OF TREATMENT-RESISTANT DEPRESSION IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE TWICE PER WEEK AS A PHARMACEUTICAL COMPOSITION IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3446 TREATMENT OF TREATMENT-RESISTANT DEPRESSION IN ADULTS BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE AS A PHARMACEUTICAL COMPOSITION IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT
- U-3447 A METHOD OF TREATING CORONARY ARTERY DISEASE
- U-3448 A METHOD OF TREATING HYPERTENSION
- U-3449 USE IN COMBINATION WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-3450 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH A REARRANGED DURING TRANSFECTION (RET) GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST
- U-3451 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC MEDULLARY THYROID CANCER (MTC) WITH A RET MUTATION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY
- U-3452 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER WITH ADVANCED OR METASTATIC THYROID CANCER WITH A RET GENE FUSION, AS DETECTED BY AN FDA-APPROVED TEST, WHO REQUIRE SYSTEMIC THERAPY AND WHO ARE RADIOACTIVE IODINE-REFRACTORY
- U-3453 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC SOLID TUMORS WITH A RET GENE FUSION THAT HAVE PROGRESSED ON OR FOLLOWING PRIOR SYSTEMIC TREATMENT OR WHO HAVE NO SATISFACTORY ALTERNATIVE TREATMENT OPTIONS
- U-3454 METHOD OF TREATING OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION IN PATIENTS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-3455 TREATMENT OF OCULAR INFLAMMATION AND PAIN FOLLOWING OPHTHALMIC SURGERY

U-3456 METHOD OF TREATING INTRAHEPATIC CHOLANGIOCARCINOMA

U-3457 METHOD OF INDUCING OCULAR ANESTHESIA

U-3458 A METHOD OF TREATING SEIZURES

U-3459 TREATING SICKLE CELL DISEASE BY ADMINISTERING VOXELOTOR AND ANOTHER ACTIVE AGENT

U-3460 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS

U-3461 INDICATED TO RAISE BLOOD PRESSURE IN ADULT PATIENTS WITH SEVERE, ACUTE HYPOTENSION

U-3462 USE OF A LIQUID FORMULATION COMPRISING FUROSEMIDE TO TREAT CONGESTION DUE TO FLUID OVERLOAD (EDEMA) IN ADULTS WITH NYHA CLASS II/III CHRONIC HEART FAILURE

U-3463 USE OF TRINTELLIX FOR THE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD) IN ADULTS

U-3464 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING ONCE DAILY A TABLET CONTAINING ABOUT 0.5 MG TO ABOUT 10 MG OF PEMIGATINIB

U-3465 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB QD FOR 14 DAYS THEN NOT ADMINISTERING PEMIGATINIB FOR 7 DAYS IN A 21-DAY CYCLE

U-3466 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB IN A DAILY DOSE OF ABOUT 5 MG TO ABOUT 20 MG

U-3467 PREVENTION AND TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING

U-3468 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS

U-3469 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-3470 METHOD OF TREATING COMPLICATED URINARY TRACT INFECTIONS (CUTI), INCLUDING PYELONEPHRITIS, COMPRISING ADMINISTERING CEFIDEROCOL SULFATE TOSYLATE

U-3471 METHOD OF TREATING HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) COMPRISING ADMINISTERING CEFIDEROCOL SULFATE TOSYLATE

U-3472 METHOD TO TREAT IDA IN ADULTS WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & HEAVY UTERINE BLEEDING OR GI DISORDER WEIGHING AT LEAST 40 KG BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO GIVE AT LEAST 0.6 G OF IRON IN 15 MINUTES OR LESS

U-3473 METHOD TO TREAT IRON DEFICIENCY ANEMIA IN ADULTS WITH INTOLERANCE OR UNSATISFACTORY RESPONSE TO ORAL IRON & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.6 G OF ELEMENTAL IRON IN 15 MINUTES OR LESS

U-3474 METHOD TO TREAT IRON DEFICIENCY ANEMIA IN ADULTS WITH NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE & WEIGHING AT LEAST 40 KG BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.6 G OF ELEMENTAL IRON IN 15 MINUTES OR LESS

U-3475 REDUCTION OF THE FREQUENCY OF ABDOMINAL PAIN AND DIARRHEA, IN AN INFLAMMATORY BOWEL DISEASE WITH DIARRHEA (IBS-D) PATIENT, WITH ELUXADOLINE TWICE DAILY WITH FOOD

U-3476 TREATMENT OF ADULTS WITH MODERATE HEPATIC IMPAIRMENT AND RELAPSED OR REFRACTORY ADVANCED RENAL CELL CARCINOMA FOLLOWING TWO OR MORE PRIOR SYSTEMIC ANTI-CANCER THERAPIES WITH 1MG TIVOZANIB HCL ORALLY FOR 21 DAYS FOLLOWED BY NO DRUG FOR 7 DAYS

U-3477 ACCELERATE THE TRANSIT OF A BARIUM MEAL THROUGH THE SMALL BOWEL, THEREBY DECREASING THE TIME AND EXTENT OF RADIATION ASSOCIATED WITH FLUOROSCOPY AND X-RAY EXAMINATION OF THE INTESTINAL TRACT

U-3478 STIMULATE GALLBLADDER CONTRACTION, AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS

U-3479 STIMULATE PANCREATIC SECRETION IN COMBINATION WITH SECRETIN PRIOR TO OBTAINING A DUODENAL ASPIRATE FOR ANALYSIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3480 STIMULATE GALLBLADDER CONTRACTION. AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS
- U-3481 DIAGNOSIS OF GALL BLADDER DISORDERS OR OTHER DIAGNOSTIC IMAGING BY STIMULATING GALLBLADDER CONTRACTION, AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS
- U-3482 DIAGNOSIS OF PANCREATIC DISORDERS BY STIMULATING PANCREATIC SECRETION IN COMBINATION WITH SECRETIN PRIOR TO OBTAINING A DUODENAL ASPIRATE FOR ANALYSIS
- U-3483 DIAGNOSTIC IMAGING BY ACCELERATING THE TRAN IT OFA BARIUM MEAL THROUGH THE SMALL BOWEL, THEREBY DECREASING THE TIME AND EXTENT OF RADIATION ASSOCIATED WITH FLUOROSCOPY AND X-RAY EXAMINATION OF THE INTESTINAL TRACT
- U-3484 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG) REQUIRING HOSPITALIZATION AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALE THEREOF, IS NOT RECOMMENDED
- U-3485 TREATMENT OF CORONAVIRUS DISEASE 2019 (COVID-19) IN NON-HOSPITALIZED ADULTS AND PEDIATRIC PATIENTS (AT LEAST 28 DAYS OF AGE AND 3 KG) AND FOR WHOM CONCOMITANT USE OF CHLOROQUINE, OR AN ANALOG OR SALT THEREOF, IS NOT RECOMMENDED
- U-3486 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA
- U-3487 TREATMENT OF ADULTS WITH ACTIVE NON-RADIOGRAPHIC AXIAL SPONDYLOARTHRITIS WITH OBJECTIVE SIGNS OF INFLAMMATION WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3488 TREATMENT OF SPASTICITY RESULTING FROM MULTIPLE SCLEROSIS, PARTICULARLY FOR THE RELIEF OF FLEXOR SPASMS AND CONCOMITANT PAIN, CLONUS, AND MUSCULAR RIGIDITY
- U-3489 TREATMENT OF SPASTICITY RESULTING FROM SPINAL CORD INJURIES AND OTHER SPINAL CORD DISEASES
- U-3490 TREATMENT OF ADULT PATIENTS WITH KRAS G12C-MUTATED LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), AS DETERMINED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-3491 METHOD OF TREATING LYMPHOMA
- U-3492 METHOD OF TREATING SARCOMA
- U-3493 THE TREATMENT OF RELAPSING FORMS OF MULTIPLE SCLEROSIS (MS) IN PATIENTS 10 YEARS OF AGE AND OLDER
- U-3494 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN FEMALE ADULTS
- U-3495 A METHOD OF TREATING ACUTE MYELOID LEUKEMIA (AML) IN PATIENTS WITH AN ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION
- U-3496 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS ACUTE MYELOID LEUKEMIA (AML)
- U-3497 A METHOD OF TREATING A CANCER WHERE THE CANCER IS ACUTE MYELOID LEUKEMIA (AML)
- U-3498 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA BY ADMINISTERING DAILY ELX (200 MG OR 100 MG); TEZ; AND IVA
- U-3499 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSSI) IN ADULT AND PEDIATRIC PATIENTS USING A TWO-DOSE REGIMEN OF DALBAVANCIN
- U-3500 TREATMENT OF ADULT PATIENTS WITH SEVERE ALOPECIA AREATA
- U-3501 PALBOCICLIB FOR HR-POS. HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER IN COMBO WITH AN AROMATASE INHIBITOR IN PTS AS INITIAL ENDOCRINE-BASED THERAPY OR WITH FULVESTRANT WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-3502 TREATMENT OF A UREA CYCLE DISORDER INVOLVING DEFICIENCIES OF CARBAMYLPHOSPHATE SYNTHETASE, ORNITHINE TRANSCARBAMYLASE, OR ARGININOSUCCINIC ACID SYNTHETASE
- U-3503 ADJUNCTIVE THERAPY TO ANTIDEPRESSANTS FOR THE TREATMENT OF MAJOR DEPRESSIVE DISORDER
- U-3504 A METHOD TO DECREASE THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A PLATINUM/ETOPOSIDE-CONTAINING REGIMEN THAT INCLUDES AN IMMUNE CHECKPOINT INHIBITOR FOR EXTENSIVE-STAGE SMALL CELL CANCER

PATENT AND EXCLUSIVITY TERMS

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PATENT USE

- U-3505 FOR TOPICAL TREATMENT OF MODERATE AD IN NON-IMMUNOCOMPROMISED PATIENTS, WITH BASELINE BSA OF 3-20% AND ITCH NRS SCORE OF ≥ 4 , WHOSE DISEASE IS NOT ADEQUATELY CONTROLLED WITH TOPICAL PRESCRIPTION THERAPIES OR WHEN THOSE THERAPIES ARE NOT ADVISABLE
- U-3506 TREATMENT OF INDOLENT SYSTEMIC MASTOCYTOSIS (ISM)
- U-3507 IN COMBINATION WITH OTHER ANTIRETROVIRAL(S), FOR THE TREATMENT OF HIV-1 INFECTION IN HEAVILY-TREATMENT EXPERIENCED ADULTS WITH MULTIDRUG RESISTANT HIV-1 INFECTION
- U-3508 REDUCTION IN THE INCIDENCE OF RECURRENT VULVOVAGINAL CANDIDIASIS (RVVC) IN ADULT AND POST-MENARCHAL PEDIATRIC FEMALES
- U-3509 AS-NEEDED TREATMENT OR PREVENTION OF BRONCHOCONSTRICTION AND REDUCTION OF THE RISK OF EXACERBATIONS IN PATIENTS WITH ASTHMA 18 YEARS OF AGE AND OLDER
- U-3510 COMBINATION TREATMENT OF COLORECTAL CANCER INCLUDING RAS WILD-TYPE HER2 (ERBB2)-POSITIVE OR -OVEREXPRESSING UNRESECTABLE OR METASTATIC COLORECTAL CANCER
- U-3511 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULT FEMALE SUBJECTS AND SYMPTOMS THEREOF
- U-3512 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULT FEMALE SUBJECTS
- U-3513 TREATMENT OF SCHIZOPHRENIA IN ADULTS
- U-3514 INCREASING BLOOD PRESSURE IN A PATIENT HAVING DISTRIBUTIVE SHOCK
- U-3515 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF AN OPIOID ANTAGONIST COMPRISING AN EMERGENCY SYRINGE DEVICE
- U-3516 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF AN OPIOID ANTAGONIST COMPRISING AN EMERGENCY SYRINGE DEVICE INCLUDING A NEEDLE GUARD
- U-3517 A METHOD FOR ADMINISTERING A THERAPEUTIC DOSE OF AN OPIOID ANTAGONIST COMPRISING AN EMERGENCY SYRINGE DEVICE INCLUDING A WINDOW CONFIGURED TO ALLOW THE USER TO VIEW THE OPIOID ANTAGONIST IN THE SYRINGE
- U-3518 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MANTLE CELL LYMPHOMA (MCL) AFTER AT LEAST TWO LINES OF SYSTEMIC THERAPY, INCLUDING A BTK INHIBITOR
- U-3519 TREATMENT WITH LENVIMA BY ADMINISTERING LENVIMA AS A SUSPENSION
- U-3520 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING A COMPOSITION COMPRISING SOLRIAMFETOL HYDROCHLORIDE AND 2-CHLOROPROPANE, WHEREIN THE COMPOSITION COMPRISES LESS THAN ABOUT 5 PPM 2CHLOROPROPANE
- U-3521 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING MILD, MODERATE, OR SEVERE RENAL IMPAIRMENT
- U-3522 METHOD OF TREATING EMESIS
- U-3523 TREATMENT OF AN ER-POSITIVE BREAST CANCER
- U-3524 TREATMENT OF AN ER-POSITIVE BREAST CANCER FOLLOWING AT LEAST ONE LINE OF ENDOCRINE THERAPY
- U-3525 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA BY ADMINISTERING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-3526 TREATMENT OF CF IN A PATIENT AGE 1 TO <6 YEARS AND WEIGHING 7 KG OR MORE WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3527 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO IS HOMOZYGOUS FOR F508DEL OR HAS AT LEAST ONE CFTR GENE MUTATION RESPONSIVE TO TEZ/IVA BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE USING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-3528 TREATMENT OF CF IN A PATIENT AGE 4 MONTHS TO <6 YEARS WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3529 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3530 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO HAS ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-3531 TREATMENT OF A TYPE 2 DIABETES PATIENT WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN THERAPY USING A COMPOSITION COMPRISING AN EXTENDED RELEASE CORE COMPRISING METFORMIN AND AN OUTER COATING COMPRISING EMPAGLIFLOZIN AND LINAGLIPTIN
- U-3532 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN ADULTS WITH TYPE 2 DIABETES MELLITUS BY ONCE DAILY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3533 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND CARDIOVASCULAR DISEASE BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-3534 PREVENTIVE TREATMENT OF MIGRAINE IN ADULTS
- U-3535 A METHOD OF TREATING ANEMIA
- U-3536 TREATMENT OF DEPRESSION IN ADULTS WITH MOD AND ACUTE SUICIDAL IDEATION OR BEHAVIOR IN CONJUNCTION WITH AN ORAL ANTIDEPRESSANT BY NASALLY ADMINISTERING 56MG OR 84MG OF ESKETAMINE IN A MAINTENANCE PHASE WEEKLY OR LX EVERY 2 WEEKS AFTER INDUCTION PHASE
- U-3537 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS SUFFERING FROM GRADE 2 OR GRADE 3 ARTHRALGIA WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3538 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-3539 TREATING ACQUIRED, GENERALIZED HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD) IN A PREMENOPAUSAL FEMALE PATIENT WITH CONTROLLED HYPERTENSION BY INJECTING BREMELANOTIDE MORE THAN ONCE WITH AT LEAST 24 HOURS BETWEEN DOSES AND NO MORE THAN 8 DOSES PER MONTH
- U-3540 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY INTRAVITREAL ADMINISTRATION OF PEGCETACOPLAN
- U-3541 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY ADMINISTERING COMPLEMENT INHIBITOR PEGCETACOPLAN
- U-3542 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY INTRAVITREAL ADMINISTRATION OF PEGCETACOPLAN AND ALSO ADMINISTERING AN ANTI-VEGF AGENT
- U-3543 TREATMENT TO INCREASE BONE DENSITY IN MEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE
- U-3544 TREATMENT OF CF IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA BY ADMINISTERING ELEXACAFTOR, IVACAFTOR, AND A SOLID DISPERSION OF TEZACAFTOR AND A POLYMER
- U-3545 TREATMENT OF CF IN A PATIENT AGE 6 YEARS AND OLDER WHO IS HOMOZYGOUS FOR F508DEL OR HAS AT LEAST ONE CFTR GENE MUTATION RESPONSIVE TO TEZ/IVA BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE USING THE COMPOSITION RECITED IN US 11578062 CLAIM 6 OR 13
- U-3546 IN COMBINATION WITH ENDOCRINE THERAPY (TAMOXIFEN OR AN AROMATASE INHIBITOR) FOR THE ADJUVANT TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE, NODE POSITIVE, EARLY BREAST CANCER AT HIGH RISK OF RECURRENCE
- U-3547 INTRAVENOUS SOTALOL DOSING REGIMEN FOR ACHIEVING STEADY STATE EXPOSURE IN A FACILITY THAT CAN PROVIDE ELECTROCARDIOGRAPHIC MONITORING
- U-3548 TREATMENT OF AML BY ORALLY ADMINISTERING VENETOCLAX WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE TO ADULTS 75 YEARS OR OLDER OR HAVING CERTAIN COMORBIDITIES PER A DOSE RAMP-UP INCLUDING AN INITIAL 100 MG OR A FINAL 400 MG PER DAY DOSE
- U-3549 INTRAVENOUS SOTALOL DOSING REGIMEN FOR USE IN A FACILITY THAT CAN PROVIDE ELECTROCARDIOGRAPHIC MONITORING
- U-3550 FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER, IN THE ABSENCE OF LASER OR PHOTOTHERAPY, WHEREIN THE VITILIGO AFFECTS AT LEAST ONE OF THE LOWER EXTREMITIES, TRUNK, AND FEET OF THE PATIENT
- U-3551 FOR THE TOPICAL TREATMENT OF NONSEGMENTAL VITILIGO IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER, IN THE ABSENCE OF PHOTOTHERAPY, WHEREIN THE VITILIGO AFFECTS AT LEAST ONE OF THE LOWER EXTREMITIES, TRUNK, AND FEET OF THE PATIENT
- U-3552 TREATMENT OF FRIEDREICH'S ATAXIA IN ADULTS AND ADOLESCENTS AGED 16 YEARS AND

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- U-3553 A METHOD OF TREATING MILD TO MODERATE ACUTE PAIN IN ADULTS BY ADMINISTERING 975-1000 MG OF ACETAMINOPHEN AND 292.5-300 MG OF IBUPROFEN IN A SINGLE ADMINISTRATION
- U-3554 AS A SINGLE AGENT FOR THE TREATMENT OF ADULT PATIENTS WITH HISTIOCYTIC NEOPLASMS
- U-3555 ADMINISTRATION OF ZAVEGEPANT FOR ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-3556 TREATMENT OF RETT SYNDROME OR A SYMPTOM THEREOF
- U-3557 TREATMENT OF PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND SYMPTOMATIC PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-3558 AN ADJUNCTIVE TREATMENT OF ADULT PATIENTS WITH TAVNEOS (AVACOPAN) WITH SEVERE ACTIVE ANCA-ASSOCIATED VASCULITIS (GPA AND MPA) IN COMBINATION WITH STANDARD THERAPY INCLUDING GLUCOCORTICOIDS
- U-3559 A METHOD OF TARGETING RELEASE OF A NON-STEROIDAL ANTI-INFLAMMATORY DRUG (NSAID) TO THE SMALL INTESTINE OF THE SUBJECT WHEN ADMINISTERED ORALLY
- U-3560 FOR THE TREATMENT OF VERNAL KERATOCONJUNCTIVITIS IN CHILDREN AND ADULTS
- U-3561 TREATMENT OF PRESBYOPIA IN ADULTS BY ADMINISTRATION OF PILOCARPINE HCl FORMULATION ONCE DAILY
- U-3562 TREATMENT OF PRESBYOPIA IN ADULTS BY ADMINISTRATION OF PILOCARPINE HCl FORMULATION TWICE DAILY
- U-3563 DEXTROMETHORPHAN AND BUPROPION IN COMBINATION TO INCREASE DEXTROMETHORPHAN PLASMA LEVELS
- U-3564 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA (LGG) WITH A BRAF V600E MUTATION WHO REQUIRE SYSTEMIC THERAPY
- U-3565 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH LOW-GRADE GLIOMA (LGG) WITH A BRAF V600E MUTATION WHO REQUIRE SYSTEMIC THERAPY
- U-3566 TREATMENT OF CANDIDEMIA AND INVASIVE CANDIDIASIS WITH REZAFUNGIN BY INTRAVENOUS ADMINISTRATION
- U-3567 TO INCREASE MEAN ARTERIAL BLOOD PRESSURE IN ADULT PATIENTS WITH HYPOTENSION ASSOCIATED WITH SEPTIC SHOCK
- U-3568 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INHIBITORS
- U-3569 TREATMENT OF PREVIOUSLY TREATED, UNRESECTABLE LOCALLY ADVANCED OR METASTATIC CHOLANGIOCARCINOMA WITH A FGFR2 FUSION OR OTHER REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INDUCERS
- U-3570 TREATMENT OF RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS WITH FGFR1 REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INDUCERS
- U-3571 TREATMENT OF RELAPSED OR REFRACTORY MYELOID/LYMPHOID NEOPLASMS WITH FGFR1 REARRANGEMENT BY ADMINISTERING PEMIGATINIB WHILE AVOIDING THE CONCOMITANT USE OF STRONG AND MODERATE CYP3A INHIBITORS
- U-3572 RAISE FOLATE LEVELS IN WOMEN WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION FOR THE PURPOSE OF REDUCING THE RISK OF A NEURAL TUBE DEFECT IN A PREGNANCY
- U-3573 TREATMENT OF MODERATE ACNE VULGARIS IN WOMEN AT LEAST 14 YEARS OF AGE IF THE PATIENT DESIRES AN ORAL CONTRACEPTIVE FOR BIRTH CONTROL
- U-3574 TREATMENT OF MOOD CHANGES AND/OR ANXIETY AS SYMPTOMS OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) IN WOMEN WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
- U-3575 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS) IN ADULTS WHO HAVE A MUTATION IN THE SUPEROXIDE DISMUTASE 1 (SOD1) GENE
- U-3576 TREATMENT OF NARCOLEPSY-RELATED CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS USING A ONCE-DAILY PHARMACEUTICAL FORMULATION COMPRISING AN OXYBATE

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- U-3577 TREATMENT OF NARCOLEPSY-RELATED CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS USING A SINGLE DAILY, BEDTIME DOSE OF A GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3578 TREATMENT OF NARCOLEPSY, CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS USING A ONCE-NIGHTLY GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3579 TREATMENT OF NARCOLEPSY AND ASSOCIATED DISORDERS AND SYMPTOMS USING A COMPOSITION COMPRISING GAMMA-HYDROXYBUTYRATE ONCE DAILY
- U-3580 TREATMENT OF NARCOLEPSY USING A DOSE PROPORTIONAL, ORAL PHARMACEUTICAL COMPOSITION COMPRISING GAMMA-HYDROXYBUTYRATE ONCE DAILY
- U-3581 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS IN ADULTS WITH NARCOLEPSY
- U-3582 LIQREV IS INDICATED FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP 1) IN ADULTS TO IMPROVE EXERCISE ABILITY AND DELAY CLINICAL WORSENING
- U-3583 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3584 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA BY ADMINISTERING ELEXACAFOTOR, IVACAFOTOR, AND A SOLID DISPERSION OF TEZACAFOTOR AND A POLYMER
- U-3585 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING THE COMPOSITION RECITED IN US 11564916 CLAIM 1
- U-3586 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA BY ADMINISTERING DAILY ELX (100 MG OR 80 MG); TEZ; AND IVA
- U-3587 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 6 YEARS AND OLDER WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A RESPONSIVE MUTATION BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3588 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS AND OLDER WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3589 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3590 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH AN EFFECTIVE AMOUNT OF ELX, TEZ, AND IVA
- U-3591 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA USING A SOLID COMPOSITION COMPRISING ELX, TEZ, AMORPHOUS IVA, AND < ~30% CRYSTALLINE IVA
- U-3592 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH ELX, TEZ, AND IVA
- U-3593 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE A R117H MUTATION IN THE CFTR GENE WITH ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3594 TREATMENT OF IRON DEFICIENCY ANEMIA (DIA) IN ADULTS PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON, WHO HAVE NON-HEMODIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE, BY ADMINISTERING FERRIC DERISOMALTOSE
- U-3595 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOUND OF CLAIM 1 OR COMPOSITION OF CLAIM 29 OF US 11426407
- U-3596 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE WITH AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL COMPOSITION COMPRISING ELEXACAFOTOR, TEZACAFOTOR, AND IVACAFOTOR
- U-3597 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE IN THE CFTR GENE AT LEAST ONE F508DEL MUTATION OR A MUTATION THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO ANY ONE OF CLAIMS 1-3 AND 7-9 OF

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- U-3598 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 11147770
- U-3599 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 10272046
- U-3600 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION IN THE CFTR GENE OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA WITH A COMPOSITION ACCORDING TO CLAIM 1 OF US 10081621
- U-3601 TREATMENT OF A DISORDER TREATABLE WITH GAMMA-HYDROXYBUTYRATE USING A SINGLE, DAILY DOSE OF A GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3602 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS WITH A DOSING REGIMEN THAT INCLUDES ORAL ADMINISTRATION OF 75 MG ONCE DAILY FOR AT LEAST 3 DAYS FOLLOWED BY 150 MG ONCE DAILY
- U-3603 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11564916
- U-3604 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11147770
- U-3605 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 10272046
- U-3606 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-3607 TREATMENT OF A MODERATE TO MILD CLINICAL PHENOTYPE OF CF USING IVACAFTOR IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE CFTR MUTATION RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-3608 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS A R117H MUTATION IN THE CFTR GENE
- U-3609 TREATMENT OF CYSTIC FIBROSIS USING IVACAFTOR IN A PATIENT AGE 1 MONTH TO <4 MONTHS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- U-3610 CO-ADMINISTRATION OF CENOBAMATE WITH PHENOBARBITAL AND/OR PHENYTOIN FOR THE TREATMENT OF PARTIAL ONSET SEIZURES
- U-3611 TREATMENT OF PEDIATRIC PATIENTS WITH CENTRAL PRECOCIOUS PUBERTY
- U-3612 AS AN ADJUNCT TO DIET TO REDUCE LOW-DENSITY LIPOPROTEIN CHOLESTEROL IN ADULTS AND PEDIATRIC PATIENTS AGED 10 YEARS AND OLDER WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-3613 AS AN ADJUNCT TO OTHER LOW-DENSITY LIPOPROTEIN CHOLESTEROL (LDL-C) LOWERING THERAPIES, OR ALONE IF SUCH TREATMENTS ARE UNAVAILABLE, TO REDUCE LDL-C IN ADULTS AND PEDIATRIC PATIENTS AGED 10 YEARS AND OLDER WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-3614 POSITRON EMISSION TOMOGRAPHY (PET) OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA) POSITIVE LESIONS IN MEN WITH PROSTATE CANCER WITH SUSPECTED METASTASIS WHO ARE CANDIDATES FOR INITIAL DEFINITIVE THERAPY
- U-3615 POSITRON EMISSION TOMOGRAPHY (PET) OF PROSTATE-SPECIFIC MEMBRANE ANTIGEN (PSMA) POSITIVE LESIONS IN MEN WITH PROSTATE CANCER WITH SUSPECTED RECURRENCE BASED ON ELEVATED SERUM PROSTATE-SPECIFIC ANTIGEN (PSA) LEVEL
- U-3616 A METHOD OF ADMINISTERING AN OPIOID MAINTENANCE TREATMENT COMPRISING BUPRENORPHINE
- U-3617 A METHOD OF ADMINISTERING AN OPIOID MAINTENANCE TREATMENT COMPRISING BUPRENORPHINE. A METHOD OF TREATING OPIOID WITHDRAWAL USING AN OPIOID

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- U-3618 A METHOD OF SUSTAINED DELIVERY OF BUPRENORPHINE TO A HUMAN OR NON-HUMAN ANIMAL BODY. A METHOD FOR TREATMENT FOR OPIOID MAINTENANCE THERAPY
- U-3619 A METHOD OF DELIVERY OF A BIOACTIVE AGENT BY SUBCUTANEOUS INJECTION. A METHOD OF TREATMENT OF A HUMAN FOR ADDICTION
- U-3620 A METHOD OF DELIVERY OF AN OPIOID BIOACTIVE AGENT. A METHOD OF TREATMENT OR PROPHYLAXIS OF A HUMAN OR NON-HUMAN ANIMAL FOR THE TREATMENT OF OPIOID ADDICTION AND/OR THE SYMPTOMS OF OPIOID WITHDRAWAL
- U-3621 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS DUE WITH MENOPAUSE
- U-3622 TREATMENT OF MODERATE TO SEVERE VASOMETER SYMPTOMS DUE TO MENOPAUSE
- U-3623 MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS AGED 5 YEARS AND OLDER. RECOMMENDED DOSAGES: BREO 100/25 OR 200/25 AGES 18 YEARS AND OLDER; BREO 100/25 AGES 12-17 YEARS, AND BREO 50/25, AGES 5-11 YEARS
- U-3624 TREATMENT OF ADULTS WITH MODERATELY TO SEVERELY ACTIVE CROHN'S DISEASE WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO ONE OR MORE TNF BLOCKERS
- U-3626 TREATMENT OF SCHIZOPHRENIA BY ADMINISTRATION OF A SIX-MONTH PALIPERIDONE PALMITATE INJECTABLE SUSPENSION FILLED SYRINGE THAT HAS BEEN SHIPPED AND STORED IN A HORIZONTAL POSITION
- U-3627 TREATMENT OF DRY EYE DISEASE (DED)
- U-3628 REDUCTION OF RISK OF CARDIOVASCULAR DEATH, HOSPITALIZATION FOR HEART FAILURE, AND URGENT HEART FAILURE IN ADULTS WITH HEART FAILURE OR TYPE 2 DIABETES MELLITUS, CHRONIC KIDNEY DISEASE, AND OTHER CARDIOVASCULAR RISK FACTORS
- U-3629 TREATMENT OF MILD-TO-MODERATE CORONAVIRUS DISEASE 2019 (COVID-19) IN ADULTS WHO ARE AT HIGH RISK FOR PROGRESSION TO SEVERE COVID-19, INCLUDING HOSPITALIZATION OR DEATH
- U-3630 TREATING OPIOID OVERDOSE
- U-3631 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS BRCA-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER IN COMBINATION WITH ABIRATERONE AND PREDNISONE OR PREDNISOLONE
- U-3632 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA
- U-3633 TREATMENT OF ADULT PATIENTS WITH MULTIPLE MYELOMA
- U-3634 METHOD OF TREATING IDA IN ADULT PATIENTS WEIGHING AT LEAST 46.7 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST 0.7 G OF ELEMENTAL IRON IN 15 MINUTES
- U-3635 METHOD TO TREAT IDA IN ADULTS WEIGHING AT LEAST 40 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON IN ABOUT ≤ 15 MIN
- U-3636 METHOD TO TREAT IDA IN ADULTS WEIGHING AT LEAST 40 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY INTRAVENOUSLY ADMINISTERING FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-3637 METHOD TO TREAT IRON DEFICIENCY IN ADULTS WEIGHING AT LEAST 40 KG WITH HEART FAILURE & NY HEART ASSOCIATION CLASS II/III TO IMPROVE EXERCISE CAPACITY BY ADMINISTERING IV FERRIC CARBOXYMALTOSE TO PROVIDE AT LEAST ABOUT 0.6 G OF ELEMENTAL IRON
- U-3638 A METHOD FOR TREATING AND/OR REDUCING THE RISK OF ACUTE MYOCARDIAL INFARCTION
- U-3639 A METHOD FOR TREATING AND/OR REDUCING THE RISK OF A CARDIOVASCULAR EVENT
- U-3640 A METHOD OF TREATING AND/OR REDUCING THE RISK OF INFLAMMATION, ATHEROSCLEROTIC VASCULAR DISEASE, AND CHOLESTEROL CRYSTAL INDUCED INFLAMMATION WITHIN ATHEROSCLEROTIC PLAQUES
- U-3641 A METHOD OF TREATING AND/OR REDUCING THE RISK OF A CARDIOVASCULAR EVENT; ACUTE CORONARY SYNDROME, OUT-OF-HOSPITAL CARDIAC ARREST, AND/OR NONCARDIOEMBOLIC ISCHEMIC STROKE
- U-3642 A METHOD OF TREATING CARDIOVASCULAR DISEASE
- U-3643 METHOD OF TREATING AND/OR REDUCING THE RISK OF A CARDIOVASCULAR EVENT

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- U-3644 TREATMENT OF FUNCTIONAL CONSTIPATION IN PEDIATRIC PATIENTS 6 TO 17 YEARS OF AGE
- U-3645 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER BY BUCCAL ADMINISTRATION
- U-3646 A METHOD OF TREATMENT OF OVARIAN CANCER OR FALLOPIAN TUBE CANCER
- U-3647 A METHOD OF TREATMENT OF RECURRENT OVARIAN CANCER OR FALLOPIAN TUBE CANCER ASSOCIATED WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATION
- U-3648 METHOD OF TREATING CHOLESTATIC PRURITUS IN PATIENTS 12 MONTHS OR OLDER SUFFERING FROM ALAGILLE SYNDROME (ALGS)
- U-3649 METHOD OF REDUCING SERUM BILE ACIDS IN PATIENTS 12 MONTHS OR OLDER SUFFERING FROM ALAGILLE SYNDROME (ALGS)
- U-3650 TREATMENT OF PATIENTS WITH POST-TRANSPLANT CYTOMEGALOVIRUS (CMV) INFECTION/DISEASE REFRACTORY TO TREATMENT WITH GANCICLOVIR, VALGANCICLOVIR, CIDOFOVIR, OR FOSCARNET, WHERE THE PATIENT IS A STEM CELL, KIDNEY, OR LIVER TRANSPLANT RECIPIENT
- U-3651 TREATMENT OF ADULT PATIENTS WITH HRR GENE-MUTATED METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (MCRPC) IN COMBINATION WITH ENZALUTAMIDE
- U-3652 AS AN ADJUNCT TO DIET AND STATIN THERAPY FOR THE TREATMENT OF ADULTS WITH PRIMARY HYPERLIPIDEMIA, INCLUDING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH), BY INHIBITING EXPRESSION OF THE PCSK9 GENE
- U-3653 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS IN A PATIENT WITH MODERATE HEPATIC IMPAIRMENT
- U-3654 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150 MG OR 200 MG ELAGOLIX WHILE CO-ADMINISTERING OMEPRAZOLE
- U-3655 MANAGEMENT OF HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS) USING 300 MG ELAGOLIX WHILE CO-ADMINISTERING OMEPRAZOLE
- U-3656 TREATMENT OF METASTATIC COLORECTAL CANCER ALONE OR WITH BEVACIZUMAB IN PATIENTS PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY
- U-3657 TREATMENT OF METASTATIC COLORECTAL CANCER ALONE OR WITH BEVACIZUMAB IN SEVERELY RENALLY IMPAIRED PATIENTS PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN THERAPY, AN ANTI-VEGF BIOLOGIC, AND IF RAS WILD-TYPE, ANTI-EGFR THERAPY
- U-3658 TREATMENT OF METASTATIC GASTRIC OR GJA IN SEVERELY RENALLY IMPAIRED PATIENTS TREATED WITH AT LEAST TWO LINES OF CHEMOTHERAPY THAT INCLUDED A FLUOROPYRIMIDINE, A PLATINUM, A TAXANE OR IRINOTECAN, AND IF APPROPRIATE, HER2/NEU-TARGETED THERAPY
- U-3659 TREATMENT OF METASTATIC COLORECTAL CANCER WITH BEVACIZUMAB IN PATIENTS PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY
- U-3660 TREATMENT OF PARTIAL-ONSET SEIZURES IN ADULTS AND IN PEDIATRIC PATIENTS WEIGHING AT LEAST 50 KG
- U-3661 COMBINATION WITH STANDARD CYTARABINE AND ANTHRACYCLINE INDUCTION AND CYTARABINE CONSOLIDATION, AND AS MAINTENANCE MONOTHERAPY FOLLOWING CONSOLIDATION CHEMOTHERAPY, FOR ADULT PATIENTS WITH NEWLY DIAGNOSED FLT3-ITD POSITIVE ACUTE MYELOID LEUKEMIA
- U-3662 DEXTROMETHORPHAN AND BUPROPRION IN COMBINATION TO TREAT MAJOR DEPRESSIVE DISORDER
- U-3663 TOPICAL TREATMENT OF MOLLUSCUM CONTAGIOSUM IN ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- U-3664 TOPICAL TREATMENT OF SKIN LESIONS CAUSED BY AN INFECTION WITH MOLLUSCUM CONTAGIOSUM IN ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- U-3665 TOPICAL DELIVERY OF A CANTHARIDIN FORMULATION TO ADULT AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER FOR TREATING MOLLUSCUM CONTAGIOSUM
- U-3666 USE IN ADULT AND PEDIATRIC PATIENTS WITH SUBOPTIMAL ECHOCARDIOGRAMS TO OPACIFY THE LEFT VENTRICULAR CHAMBER AND TO IMPROVE THE DELINEATION OF THE LEFT VENTRICULAR ENDOCARDIAL BORDER
- U-3667 USE IN ULTRASONOGRAPHY OF THE URINARY TRACT IN PEDIATRIC PATIENTS FOR THE

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- EVALUATION OF SUSPECTED OR KNOWN VESICoureTERAL REFLUX
- U-3668 USE WITH ULTRASOUND OF THE LIVER IN ADULT AND PEDIATRIC PATIENTS TO CHARACTERIZE FOCAL LIVER LESIONS
- U-3669 USE OF VASCEPA TO REDUCE THE INCIDENCE OF STROKE IN AN ADULT PATIENT ON STATIN THERAPY AND WITH ELEVATED TRIGLYCERIDE LEVELS (>150 MG/DL), WHEREIN THE PATIENT EXPERIENCES ATRIAL FIBRILLATION AND/OR FLUTTER INSTEAD OF AN INCIDENCE OF STROKE
- U-3670 ADMINISTRATION OF AN EXTENDED RELEASE TABLET FOR THE TREATMENT OF NEUROGENIC DETRUSOR OVERACTIVITY (NDO) IN PEDIATRIC PATIENTS AGED 3 YEARS AND OLDER AND WEIGHING 35 KG OR MORE
- U-3671 USE OF INTRANASAL NALOXONE FOR THE TREATMENT OF OPIOID OVERDOSE
- U-3672 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS USING 150 MG ELAGOLIX FOR UP TO 24 MONTHS
- U-3673 TREATMENT OF GEOGRAPHIC ATROPHY SECONDARY TO AGE-RELATED MACULAR DEGENERATION BY ADMINISTERING AVACINCAPTAD PEGOL TO THE EYE
- U-3674 TREATMENT OF DEMODEX BLEPHARITIS VIA TOPICAL ADMINISTRATION TO AN OCULAR SURFACE
- U-3675 A PERCUTANEOUS HEPATIC PERFUSION PROCEDURE FOR TREATING A PATIENT WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES
- U-3676 REDUCTION OF HETEROTOPIC OSSIFICATION IN PATIENTS WITH FIBRODYSPLASIA (MYOSITIS) OSSIFICANS PROGRESSIVA
- U-3677 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA IN A PATIENT WITH SEVERE HEPATIC IMPAIRMENT
- U-3678 A METHOD FOR REDUCING THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A PLATINUM/ETOPOSIDE-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3679 A METHOD FOR REDUCING THE INCIDENCE OF CHEMOTHERAPY-INDUCED MYELOSUPPRESSION IN ADULT PATIENTS WHEN ADMINISTERED PRIOR TO A TOPOTECAN-CONTAINING REGIMEN FOR EXTENSIVE-STAGE SMALL CELL LUNG CANCER
- U-3680 A METHOD FOR TREATING A SUBJECT WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES
- U-3681 TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE SERIOUS OR LIFE-THREATENING
- U-3682 SODIUM THIOSULFATE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM NITRITE FOR THE TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE SERIOUS OR LIFE-THREATENING
- U-3683 A METHOD OF TREATING A PATIENT WITH UVEAL MELANOMA WITH UNRESECTABLE HEPATIC METASTASES
- U-3684 A METHOD OF TREATING AN ADULT PATIENT WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA HAVING A SUSCEPTIBLE IDH1 MUTATION
- U-3685 METHOD OF ADMINISTERING AN EFFECTIVE DOSE OF TROPICAMIDE AND PHENYLEPHRINE HYDROCHLORIDE TO AN EYE
- U-3686 TREATMENT OF EXACERBATIONS OF MULTIPLE SCLEROSIS IN ADULTS WITH CORTICOTROPIN BY PROMOTING NEW VESSEL FORMATION WHEREIN VCAM-1 EXPRESSION AND ANGIOPOETIN-2 EXPRESSION IS INCREASED AFTER THE ADMINISTERING
- U-3687 TREATMENT OF INFANTILE SPASMS WITH CORTICOTROPIN BY PROMOTING NEW VESSEL FORMATION WHEREIN VCAM-1 EXPRESSION AND ANGIOPOETIN-2 EXPRESSION IS INCREASED AFTER THE ADMINISTERING
- U-3688 TREATMENT OF OPHTHALMIC DISEASES WITH CORTICOTROPIN BY PROMOTING NEW VESSEL FORMATION WHEREIN VCAM-1 EXPRESSION AND ANGIOPOETIN-2 EXPRESSION IS INCREASED AFTER THE ADMINISTERING
- U-3689 DIAGNOSIS OF GALL BLADDER DISORDERS BY STIMULATING GALLBLADDER CONTRACTION, AS MAY BE ASSESSED BY VARIOUS METHODS OF DIAGNOSTIC IMAGING, OR TO OBTAIN BY DUODENAL ASPIRATION A SAMPLE OF CONCENTRATED BILE FOR ANALYSIS
- U-3690 A METHOD FOR PREVENTING OF POST-OPERATIVE NAUSEA AND VOMITING
- U-3691 TREATMENT AND REDUCTION OF RISK BY ADMINISTRATION OF EMPAGLIFLOZIN TO ADULTS WITH CHRONIC KIDNEY DISEASE AT RISK OF PROGRESSION
- U-3692 A METHOD OF TREATING HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA OR ESTABLISHED ATHEROSCLEROTIC CARDIOVASCULAR DISEASE BY DECREASING THE LEVEL OF LDL-C USING 180 MG BEMPEDOIC ACID AND 10 MG EZETIMIBE

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- U-3693 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS IN A BREAST-FEEDING PATIENT WHILE REDUCING INFANT EXPOSURE TO SOLRIAMFETOL
- U-3694 METHOD OF REDUCING OR AMELIORATING SEIZURES IN A PATIENT BY ADMINISTERING A LIQUID FORMULATION OF FENFLURAMINE WITH STIRIPENTOL THEREBY MODULATING DOWN THE FORMATION OF NORFENFLURAMINE AND RESULTING IN HIGHER LEVELS OF FENFLURAMINE
- U-3695 MAINTENANCE TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE OR SOMATIC BRCA-MUTED RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-3696 TREATMENT OF CF IN PATIENTS AGED 2 TO <6 YEARS WHO HAVE AT LEAST ONE F508DEL MUTATION OR A MUTATION IN THE CFTR GENE THAT IS RESPONSIVE BASED ON IN VITRO DATA COMPRISING ADMINISTERING A COMPOSITION ACCORDING TO CLAIM 1 OF US 11752106
- U-3697 TREATMENT OF CF IN A PATIENT AGE 1 MONTH TO <6 YEARS WHO HAS AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA USING THE COMPOSITION RECITED IN CLAIM 1 OF US 11752106
- U-3698 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA BY SUBLINGUAL OR BUCCAL ADMINISTRATION
- U-3699 TREATMENT OF SUBJECTS WITH MAJOR DEPRESSION WITH SEXUAL DYSFUNCTION CAUSED BY EITHER MAJOR DEPRESSION OR PRIOR TREATMENTS, OR TREATMENT OF SUBJECTS WITH MAJOR DEPRESSION WITHOUT THE RISK OF SEXUAL DYSFUNCTION ADVERSE REACTIONS
- U-3700 TREATMENT OF ADULTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3701 TREATMENT OF ADULTS WITH MYCOSIS FUNGOIDES WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3702 TREATMENT OF ADULTS WITH RELAPSED OR REFRACTORY NON-HODGKIN LYMPHOMAS WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3703 TREATMENT OF ADULTS WITH RHEUMATOID ARTHRITIS WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3704 TREATMENT OF ADULTS WITH SEVERE PSORIASIS WITH AN ORAL SOLUTION OF METHOTREXATE
- U-3705 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY WHO HAVE AN APNEA/HYPOPNEA INDEX ≤ 15 WITH A ONCE-NIGHTLY FORMULATION OF GAMMA-HYDROXYBUTYRATE
- U-3706 TREATMENT OF BLOATING ASSOCIATED WITH DIARRHEA-PREDOMINANT IRRITABLE BOWEL SYNDROME (IBS-D) IN ADULT FEMALE SUBJECTS
- U-3707 A METHOD OF TREATING PATIENTS 1 YEAR OF AGE AND OLDER WITH CHRONIC PHASE PH+ CML, NEWLY-DIAGNOSED OR RESISTANT OR INTOLERANT TO PRIOR THERAPY
- U-3708 A METHOD OF TREATING PATIENTS WITH ACCELERATED, OR BLAST PHASE PH+ CML WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY
- U-3709 METHOD OF TREATING PRIMARY HYPEROXALURIA TYPE 1 (PH1)
- U-3710 TREATMENT OF ADULT PATIENTS WITH PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA BY ORALLY ADMINISTERING 100 MG OF ACALABRUTINIB TWICE DAILY IN COMBINATION WITH INTRAVENOUS ADMINISTRATION OF OBINUTUZUMAB
- U-3711 A METHOD TO IMPROVE KIDNEY FUNCTION IN ADULTS WITH HEPATORENAL SYNDROME WITH RAPID REDUCTION IN KIDNEY FUNCTION
- U-3712 TOPICAL TREATMENT OF PLAQUE PSORIASIS, INCLUDING INTERTRIGINOUS AREAS, IN PATIENTS 6 YEARS OF AGE OR OLDER
- U-3713 TOPICAL TREATMENT OF ACNE VULGARIS IN ADULT AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-3714 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS IN PATIENTS HAVING PRE-EXISTING SEVERE HEPATIC IMPAIRMENT AND SUFFERING FROM AN ADVERSE EVENT WHILE BEING ADMINISTERED RIPRETINIB DAILY
- U-3715 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) RECEIVING A MODERATE CYP3A INDUCER
- U-3716 TREATMENT OF ADULT PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA (WM) RECEIVING A MODERATE CYP3A INDUCER
- U-3717 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL) RECEIVING A MODERATE CYP3A INDUCER
- U-3718 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY MARGINAL ZONE LYMPHOMA

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- (MZL) RECEIVING A MODERATE CYP3A INDUCER, WHO HAVE RECEIVED AT LEAST ONE ANTI-CD20-BASED REGIMEN
- U-3719 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) RECEIVING A MODERATE CYP3A INDUCER, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3720 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) RECEIVING A MODERATE CYP3A INDUCER
- U-3721 TREATMENT OF SECONDARY HYPERPARATHYROIDISM WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN CHRONIC KIDNEY DISEASE PATIENTS RECEIVING CHOLESTYRAMINE
- U-3722 TREATMENT OF SECONDARY HYPERPARATHYROIDISM WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN CHRONIC KIDNEY DISEASE PATIENTS RECEIVING PHENOBARBITAL OR OTHER ANTICONVULSANTS
- U-3723 TREATMENT OF SECONDARY HYPERPARATHYROIDISM WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN CHRONIC KIDNEY DISEASE PATIENTS RECEIVING CYP3A INHIBITORS
- U-3724 TO PRODUCE POST-SURGICAL ANALGESIA
- U-3725 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I OR II DISORDER BY SUBLINGUAL OR BUCCAL ADMINISTRATION IN PATIENTS WITH SEVERE HEPATIC IMPAIRMENT
- U-3726 THE TREATMENT OF POMPE PATIENTS
- U-3727 TREATMENT OF ADULT PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3728 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3729 TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-3730 A METHOD FOR TREATING ULCERATIVE COLITIS BY ADMINISTERING ESTRASIMOD ARGININE IN AN AMOUNT EQUIVALENT TO ABOUT 2.0 MG OF ESTRASIMOD
- U-3731 A METHOD FOR TREATING ULCERATIVE COLITIS BY ADMINISTERING A THERAPEUTICALLY EFFECTIVE AMOUNT OF THE FORM OF ESTRASIMOD ARGININE AS CLAIMED
- U-3732 A METHOD FOR TREATING ULCERATIVE COLITIS BY ADMINISTERING A THERAPEUTICALLY EFFECTIVE AMOUNT OF ESTRASIMOD ARGININE
- U-3733 METHOD OF TREATING SEIZURES IN A PATIENT BY ADMINISTERING A LIQUID FORMULATION OF FENFLURAMINE OR ITS SALTS PLUS STIRIPENTOL THEREBY REDUCING NORFENFLURAMINE FORMATION WHILE INCREASING THE FENFLURAMINE LEVEL. PATIENTS CAN HAVE E.G. DRAVET OR LGS
- U-3734 METHOD OF TREATING SCHIZOPHRENIA IN A PATIENT WHO HAS PREVIOUSLY EXPERIENCED SIGNIFICANT WEIGHT GAIN INDUCED BY OLANZAPINE ALONE BY ADMINISTERING A COMPOSITION COMPRISING OLANZAPINE AND SAMIDORPHAN
- U-3735 TREATMENT OF GENERALIZED MYASTHENIA GRAVIS (GMG) IN AN ADULT PATIENT WHO IS ANTI-ACETYLCHOLINE RECEPTOR (ACHR) ANTIBODY POSITIVE BY SUBCUTANEOUS ADMINISTRATION OF C5 COMPLEMENT INHIBITOR ZILUCOPLAN
- U-3736 REDUCTION OF SERUM PHOSPHORUS IN ADULTS
- U-3737 MEKTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH ENCORAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON SMALL CELL LUNG CANCER WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- U-3738 BRAFTOVI IS A KINASE INHIBITOR INDICATED IN COMBINATION WITH BINIMETINIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON SMALL CELL LUNG CANCER WITH A BRAF V600E MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- U-3739 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE ADMINISTRATION OF TASIMELTEON WITH FOOD
- U-3740 TREATMENT OF MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC) IN ADULTS
- U-3741 TREATMENT OF PRESBYOPIA
- U-3742 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHEREIN THE CANCER IS RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES
- U-3743 A METHOD OF TREATING RELAPSED OR REFRACTORY MYELODYSPLASTIC SYNDROMES CHARACTERIZED BY THE PRESENCE OF A MUTANT ALLELE OF IDH1 BY ADMINISTERING A ONCE DAILY 500 MG ORAL DOSE TO A SUBJECT THAT HAS NOT INGESTED A HIGH-FAT MEAL
- U-3744 METHOD OF TREATING MILD TO MODERATE PAIN IN ADULTS

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- U-3745 METHOD OF TREATING MODERATE TO SEVERE PAIN IN ADULTS AS AN ADJUNCT TO OPIOID ANALGESICS
- U-3746 METHODS OF MAKING AQUEOUS COMPOSITION AND TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ATERIOSIS WITH AQUEOUS COMPOSITION
- U-3747 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY
- U-3748 TOPICAL TREATMENT OF PLAQUE PSORIASIS, INCLUDING INTERTRIGINOUS AREAS, IN PATIENTS 6 YEARS OF AGE AND OLDER
- U-3749 METHOD OF TREATING PULMONARY HYPERTENSION ASSOCIATED WITH INTERSTITIAL LUNG DISEASE BY ADMINISTERING TREPROSTINIL OR A SALT THEREOF BY INHALATION USING A DEVICE
- U-3750 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR WITH A WILD TYPE KIT MUTATION IN PATIENTS PREVIOUSLY ADMINISTERED THREE OR MORE KINASE INHIBITORS, WHERE ONE OF THE KINASE INHIBITORS IS IMATINIB
- U-3751 TREATMENT OF CATAPLEXY OR EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY WITH A ONCE-NIGHTLY GAMMA-HYDROXYBUTYRATE FORMULATION
- U-3752 METHOD OF TREATING DIABETIC HYPOGLYCEMIA
- U-3753 TREATMENT OF PATIENTS WITH METASTATIC COLORECTAL CANCER WHO HAVE BEEN PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN-, AND IRINOTECAN-BASED CHEMOTHERAPY, ANTI-VEGF THERAPY, AND, IF RAS WILD-TYPE AND MEDICALLY APPROPRIATE, ANTI-EGFR THERAPY
- U-3754 TREATMENT OF ADULT PATIENTS WITH PROGRESSING DESMOID TUMORS
- U-3755 TREATMENT OF ADULT PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ROS1-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)
- U-3756 ACUTE TREATMENT OF AGITATION ASSOCIATED WITH BIPOLAR I OR II DISORDER BY SUBLINGUAL OR BUCCAL ADMINISTRATION
- U-3757 TREATMENT OF HYPOTHYROIDISM BY ORAL ADMINISTRATION OF L-THYROXINE TO A PATIENT ON AN EMPTY STOMACH 15 MINUTES BEFORE BREAKFAST
- U-3758 TREATMENT OF PITUITARY THYROTROPIN SUPPRESSION BY ORAL ADMINISTRATION OF L-THYROXINE TO A PATIENT ON AN EMPTY STOMACH 15 MINUTES BEFORE BREAKFAST
- U-3759 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE IN PATIENTS WITH HEART FAILURE AND TYPE 2 DIABETES MELLITUS BY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3760 METHOD FOR REDUCING THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION IN PATIENTS WITH HEART FAILURE BY ADMINISTRATION OF EMPAGLIFLOZIN
- U-3761 ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA (CLL/SLL) WHO HAVE RECEIVED AT LEAST TWO PRIOR LINES OF THERAPY, INCLUDING A BTK INHIBITOR AND A BCL-2 INHIBITOR
- U-3762 TREATMENT WITH FULVESTRANT OF HR-POS. HER2-NEG. LOCALLY ADVANCED OR METASTATIC BREAST CANCER WITH PIK3CA/AKT1/PTEN-ALTERATION(S) FOLLOWING PROGRESSION ON ENDOCRINE THERAPY IN THE METASTATIC SETTING OR RECURRENCE ON OR WITHIN 12 MONTHS OF ADJUVANT THERAPY
- U-3763 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (NMCSPC) WITH BIOCHEMICAL RECURRENCE (BCR) AT HIGH RISK FOR METASTASIS
- U-3764 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING OBSTRUCTIVE SLEEP APNEA (OSA) AND NO, MILD, MODERATE, OR SEVERE RENAL IMPAIRMENT
- U-3765 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING MODERATE OR SEVERE RENAL IMPAIRMENT
- U-3766 REDUCE THE RISK OF CARDIOVASCULAR DEATH AND HOSPITALIZATION FOR HEART FAILURE AND URGENT HEART FAILURE VISITS IN ADULTS WITH HEART FAILURE WITH PRESERVED EJECTION FRACTION AND WITHOUT TYPE II DIABETES
- U-3767 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH CHRONIC, ACCELERATED, OR MYELOID OR LYMPHOID BLAST PHASE PH+ CML WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB, WHEN ADMINISTERED AT ELEVATED GASTRIC PH
- U-3768 USE OF DASATINIB FOR TREATMENT OF NEWLY DIAGNOSED ADULTS WITH PH+ CML IN CHRONIC PHASE, WHEN ADMINISTERED AT ELEVATED GASTRIC PH
- U-3769 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH PH+ ALL WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY, WHEN ADMINISTERED AT ELEVATED GASTRIC PH

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- U-3770 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH CHRONIC, ACCELERATED, OR MYELOID OR LYMPHOID BLAST PHASE PH+ CML WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB, WHEN COADMINISTERED WITH A GASTRIC ACID REDUCING AGENT
- U-3771 USE OF DASATINIB FOR TREATMENT OF ADULTS WITH PH+ ALL WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY, WHEN COADMINISTERED WITH A GASTRIC ACID REDUCING AGENT
- U-3772 USE OF DASATINIB FOR TREATMENT OF NEWLY DIAGNOSED ADULTS WITH PH+ CML IN CHRONIC PHASE, WHEN COADMINISTERED WITH A GASTRIC ACID REDUCING AGENT
- U-3773 TOPICAL TREATMENT OF SEBORRHEIC DERMATITIS IN PATIENTS 9 YEARS OF AGE AND OLDER
- U-3774 A CATHETER LOCK SOLUTION TO REDUCE CATHETER-RELATED BLOODSTREAM INFECTIONS IN ADULT PATIENTS RECEIVING HEMODIALYSIS THROUGH A CENTRAL VENOUS CATHETER
- U-3775 TREATMENT OF EXCESSIVE DAYTIME SLEEPINESS BY ADMINISTERING SOLRIAMFETOL TO A SUBJECT HAVING NO, MILD, MODERATE, OR SEVERE RENAL IMPAIRMENT
- U-3776 TREATING TYPE 2 DIABETES MELLITUS BY ASSESSING RENAL FUNCTION AND ORALLY ADMINISTERING EMPAGLIFLOZIN IN A DAILY AMOUNT OF 10 MG OR 25 MG IF THE EGFR IS ≥ 30 ML/MIN/1.73 M² AND < 60 ML/MIN/1.73 M², WHEREIN THE TREATMENT IMPROVES GLYCEMIC CONTROL
- U-3777 TREATING TYPE 2 DIABETES MELLITUS BY ASSESSING RENAL FUNCTION AND ORALLY ADMINISTERING EMPAGLIFLOZIN IN A DAILY AMOUNT OF 10 MG OR 25 MG IF THE EGFR ≥ 45 ML/MIN/1.73 M² AND < 60 ML/MIN/1.73 M², WHEREIN THE TREATMENT IMPROVES GLYCEMIC CONTROL
- U-3778 TREATMENT OF MAJOR DEPRESSIVE DISORDER BY ADMINISTERING DEXTROMETHORPHAN AND BUPROPION TO A SUBJECT HAVING MODERATE HEPATIC IMPAIRMENT
- U-3779 METHOD OF TREATING NEONATAL SEIZURES
- U-3780 TREATMENT OF ADULT PATIENTS WITH ADVANCED RENAL CELL CARCINOMA FOLLOWING A PROGRAMMED DEATH RECEPTOR-1 OR PROGRAMMED DEATH-LIGAND INHIBITOR AND A VASCULAR ENDOTHELIAL GROWTH FACTOR TYROSINE KINASE INHIBITOR
- U-3781 REDUCTION IN LOSS OF KIDNEY FUNCTION IN ADULTS WITH PRIMARY IMMUNOGLOBULIN A NEPHROPATHY (IGAN) WHO ARE AT RISK OF DISEASE PROGRESSION