INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CARNAUBA WAX	008015869					
ORAL; CAPLET		1				
ORAL; CAPSULE, SUSTAINED ACTION		2		08/12/88	600	0.364MG - 0.75MG
ORAL; TABLET		121		10/06/95	UNK	
ORAL; TABLET, COATED		46		05/17/94	600	0.02MG - 0.92MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		06/23/81	600	
ORAL; TABLET, FILM COATED		24		09/29/95	600	0.01MG - 0.4MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	0.046MG
ORAL; TABLET, SUSTAINED ACTION		29		11/14/94	UNK	0.07MG - 200.0MG
ORAL-28; TABLET		2		11/17/95	510	0.126MG - 0.157MG
CARNAUBA YELLOW WAX						
ORAL; TABLET, COATED		1				
CARRAGEENAN	009000071					
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		2		11/27/79	600	0.75%
ORAL; SYRUP		1				
TOPICAL; LOTION		2		12/17/81	600	0.5%
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
CARRAGEENAN SALT						
TOPICAL; LOTION		1				
CASSIA OIL						
ORAL; SOLUTION, ELIXIR		1				
CASTOR OIL	008001794					
INTRAMUSCULAR; INJECTION		4		02/28/79	600	
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		4		02/14/94	600	1.2MG - 1.756MG
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; TABLET		5		03/30/94	600	0.09MG - 0.15MG
ORAL; TABLET, COATED		3		01/15/70	510	0.9MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
SUBLINGUAL; TABLET		1				
TOPICAL; EMULSION, CREAM		2		01/24/80	UNK	5.0% - 12.5%
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		1				
CASTOR OIL HYDROGENATED	008001783					
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		7		08/10/92	110	1.894MG - 13.0MG
ORAL; TABLET		4		06/25/91	110	0.5MG - 37.6MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		2		01/22/87	600	111.6MG
ORAL-21; TABLET		3		04/12/88	600	0.8MG
ORAL-28; TABLET		5		02/09/89	600	0.8MG - 104.5MG
SUBLINGUAL; TABLET		1				

[illegible]

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
[illegible]						
[illegible]		1		[illegible]	[illegible]	[illegible]
[illegible]		1				
[illegible]		1				
[illegible]		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CELLULOSE						
BUCCAL/SUBLINGUAL; TABLET		1				
ORAL; CAPSULE		23		03/03/95	110	40.0MG - 405.0MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		3		04/20/88	600	
ORAL; SUSPENSION		2		08/28/81	110	1.0%
ORAL; TABLET		54		06/29/95	600	0.07GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		8		05/19/92	110	6.7MG - 20.1MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		9		08/31/81	600	5.94MG - 100.0MG
ORAL; TABLET, SUSTAINED ACTION		4		09/22/94	110	
ORAL-21; TABLET		3		05/10/82	510	20.0MG
ORAL-28; TABLET		4		11/01/84	510	20.0MG
SUBLINGUAL; TABLET		1				
CELLULOSE ACETATE	009004357					
ORAL; TABLET, SUSTAINED ACTION		5		06/01/94	110	19.0MG - 23.75MG
CELLULOSE ACETATE PHTHALATE	009004380					
ORAL; TABLET		2		06/22/89	600	37.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		07/28/88	UNK	4.75MG - 24.0MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		9		01/04/95	600	16.95MG - 70.0MG
CELLULOSE MICROCRYSTALLINE/CARBOXYMETHYLCELLULOSE SODIUM						
ORAL; CAPSULE		1				
ORAL; POWDER, FOR RECONSTITUTION		2		12/23/91	520	0.5% - 1.3125%
ORAL; SUSPENSION		1				
ORAL; TABLET		1				
CELLULOSE MICROCRYSTALLINE, AQUEOUS						
ORAL; CAPSULE		2		11/19/82	600	20.0MG
ORAL; SUSPENSION		1				
ORAL; TABLET		15		10/29/92	UNK	7.9MG - 240.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		2		03/03/95	UNK	25.0MG
CELLULOSE, MICROCRYSTALLINE	009004346					
INTRAVENOUS; INJECTION		1				
NASAL; SPRAY, METERED		3		10/19/94	UNK	1.5%
ORAL; CAPLET		1				
ORAL; CAPSULE		116		12/20/95	520	1.5MG - 363.75MG
ORAL; CAPSULE, COATED PELLETS		2		10/30/85	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		3		12/06/95	530	57.22MG - 60.0MG
ORAL; CAPSULE, SUSTAINED ACTION		6		09/11/95	110	20.7MG - 107.0MG
ORAL; GRANULE, ENTERIC COATED		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CELLULOSE, MICROCRYSTALLINE	009004346					
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		13		04/28/95	600	0.5% - 2.0%
ORAL; SUSPENSION		9		09/15/95	180	1.2%
ORAL; TABLET		1,407		12/29/95	600	0.064GM - 0.152GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		6		09/11/95	600	42.07MG - 505.0MG
ORAL; TABLET, COATED		37		06/23/95	600	1.0MG - 107.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		8		11/30/95	600	25.0MG - 125.71MG
ORAL; TABLET, DISPERSIBLE		2		04/28/95	600	
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		90		12/19/95	180	5.0MG - 160.22MG
ORAL; TABLET, SUSTAINED ACTION		14		11/18/94	UNK	23.45MG - 226.0MG
ORAL; TABLET, UNCOATED, TROCHE		1				
ORAL-21; TABLET		8		12/13/93	600	19.6MG - 20.0MG
ORAL-28; TABLET		19		11/17/95	510	7.58MG - 20.0MG
RECTAL; SUSPENSION		1				
SUBLINGUAL; TABLET		9		02/26/88	600	6.0MG - 32.0MG
VAGINAL; TABLET		2		10/17/85	600	320.0MG - 390.0MG
CELLULOSE, OXIDIZED						
ORAL; TABLET		5		10/03/74	600	0.03GM
CELLULOSIC POLYMERS						
ORAL; TABLET		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
CERESIN	008001750					
TOPICAL; OINTMENT		1				
CETEARETH-12						
TOPICAL; EMULSION, CREAM		1				
CETEARETH-15						
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		7		01/29/93	600	1.0% - 7.5%
CETEARETH-20						
TOPICAL; EMULSION, CREAM		11		12/18/90	UNK	0.75% - 10.0%
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		1				
CETEARETH-30						
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		3		07/10/84	UNK	2.25% - 7.2%
CETEARYL ALCOHOL						
ORAL; TABLET, SUSTAINED ACTION		6		11/08/93	UNK	20.0MG - 40.0MG
TOPICAL; EMULSION, CREAM		37		09/13/95	UNK	1.8% - 11.5%
TOPICAL; LOTION		2		05/31/89	UNK	2.5% - 3.7%
TOPICAL; OINTMENT		2		10/10/85	600	1.2%
TOPICAL; SUPPOSITORY		1				
VAGINAL; EMULSION, CREAM		5		12/04/95	600	3.21%

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CETEARYL ALCOHOL						
VAGINAL; SUPPOSITORY		1				
CETEARYL OCTANOATE						
TOPICAL; EMULSION, CREAM		1				
CETETH-10						
TOPICAL; LOTION		1				
CETETH-2						
TOPICAL; EMULSION, CREAM		1				
CETETH-20						
TOPICAL; EMULSION, CREAM		13		06/17/94	UNK	0.91% - 4.0%
VAGINAL; TAMPON		1				
CETYL ALCOHOL	000124298					
OPHTHALMIC; SUSPENSION		2		05/11/88	600	0.5%
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, SUSTAINED ACTION		5		01/04/95	600	0.58MG - 2.0MG
OTIC; SUSPENSION		3		09/29/87	600	0.2% - 1.0%
RECTAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; AEROSOL		1				
TOPICAL; EMULSION, AEROSOL FOAM		2		12/19/79	600	3.226%
TOPICAL; EMULSION, CREAM		77		06/13/95	600	0.2% - 10.0%
TOPICAL; LOTION		31		09/30/92	UNK	0.0014% - 5.0%
TOPICAL; OINTMENT		6		07/24/78	600	1.5% - 7.0%
VAGINAL; EMULSION, CREAM		13		12/21/95	520	1.5% - 4.0%
CETYL ESTERS						
ORAL; POWDER, FOR RECONSTITUTION		1				
TOPICAL; EMULSION, CREAM		12		10/26/94	600	2.0% - 5.0%
TOPICAL; LOTION		1				
TOPICAL; SUPPOSITORY		1				
VAGINAL; EMULSION, CREAM		5		12/04/95	600	
VAGINAL; SUPPOSITORY		1				
CETYL PALMITATE	000540103					
TOPICAL; EMULSION, CREAM		4		09/13/95	UNK	0.3144%
TOPICAL; OINTMENT		1				
VAGINAL; EMULSION, CREAM		1				
CETYLPYRIDINIUM CHLORIDE	000123035					
INHALATION; AEROSOL, METERED		2		07/10/62	UNK	
ORAL; AEROSOL SPRAY		1				
ORAL; CAPSULE		5		05/05/94	110	0.0043MG
ORAL; CAPSULE, SUSTAINED ACTION		6		01/06/81	UNK	0.02MG
CHERRY						
ORAL; TABLET		1				
CHERRY JUICE	008012995					
ORAL; CONCENTRATE		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CHLOROBUTANOL	000057158					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		4		03/18/75	600	0.5%
IM - SC; INJECTION		2		06/24/75	600	0.35% - 0.5%
INHALATION; SOLUTION		1				
INTRAMUSCULAR; INJECTION		20		03/13/86	600	0.25% - 5.0%
INTRAVENOUS; SOLUTION, INJECTION		1				
IV(INFUSION); INJECTION		5		11/19/80	510	0.5%
NASAL; SOLUTION		3		02/21/78	510	0.05% - 0.5%
NASAL; SPRAY, METERED		2		03/07/94	510	
NERVE BLOCK; INJECTION		2		11/15/79	600	0.25% - 0.5%
OPHTHALMIC; OINTMENT		5		12/01/89	UNK	0.5%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		4		08/31/81	600	0.2% - 0.5%
OTIC; SOLUTION		1				
SUBCUTANEOUS; INJECTION		1				
TOPICAL; SOLUTION		1				
CHLOROBUTANOL HEMIHYDRATE	006001645					
INTRAMUSCULAR; INJECTION		1				
CHLOROBUTANOL, ANHYDROUS	001320667					
OPHTHALMIC; SOLUTION		1				
CHLOROCRESOL	000059507					
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		21		08/03/94	600	0.075% - 0.75%
CHLOROXYLENOL	000088040					
TOPICAL; EMULSION, CREAM		1				
CHOLESTEROL	000057885					
INTRAVENOUS; SUSPENSION, INJECTION		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		3		05/18/65	600	0.9% - 5.0%
VAGINAL; EMULSION, CREAM		3		06/09/86	600	0.3% - 0.5%
CHOLETH						
VAGINAL; EMULSION, CREAM		2		07/23/82	600	0.4%
CINNAMALDEHYDE	000104552					
ORAL; SUSPENSION		1				
ORAL; TABLET		1				
CINNAMON						
ORAL; SOLUTION, ELIXIR		1				
CINNAMON OIL	008007805					
ORAL; PASTILLE		1				
ORAL; SOLUTION, ELIXIR		9		12/16/83	600	0.011%
ORAL; SUSPENSION		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CITRIC ACID	000077929					
CAUDAL BLOCK; INJECTION		1				
EPIDURAL; INJECTION		5		02/26/93	600	0.02%
IM - IV - SC; INJECTION		10		08/23/95	600	0.063% - 1.26%
IM - IV; INJECTION		21		08/31/90	600	0.01% - 0.8%
IM - IV; SOLUTION, INJECTION		1				
IM - SC; INJECTION		1				
INHALATION; SOLUTION		4		02/19/92	UNK	0.44042% - 1.0%
INTRA-ARTERIAL; SOLUTION, INJECTION		1				
INTRA-ARTICULAR; INJECTION		8		04/09/86	600	0.205%
INTRABURSAL; INJECTION		4		02/13/74	600	1.0%
INTRACARDIAC; INJECTION		1				
INTRALESIONAL; INJECTION		2		02/17/84	600	
INTRAMUSCULAR; INJECTION		17		04/03/87	600	0.02% - 2.0%
INTRAPLEURAL; INJECTION		1				
INTRASYNOVIAL; INJECTION		2		02/17/84	600	1.0%
INTRATHECAL; INJECTION		1				
INTRAVASCULAR; SOLUTION, INJECTION		1				
INTRAVENOUS; INJECTION		28		12/15/95	600	0.0175% - 0.86%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		3		07/12/88	600	
INTRAVENOUS; SOLUTION, INJECTION		1				
IONTOPHORESIS; SOLUTION		1				
IV(INFUSION); INJECTION		32		08/30/95	600	0.0229% - 1.0%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		2		08/31/95	600	
NASAL; SOLUTION		1				
NASAL; SPRAY, METERED		3		03/08/95	UNK	0.1%
NERVE BLOCK; INJECTION		15		02/26/93	600	0.02%
OPHTHALMIC; SOLUTION		5		01/04/95	600	0.035% - 0.05%
OPHTHALMIC; SUSPENSION		4		05/09/89	600	
ORAL; CAPSULE		6		08/18/95	600	5.0MG - 12.8MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SOFT GELATIN		2		12/30/86	150	0.015MG - 1.0MG
ORAL; CONCENTRATE		10		01/30/92	600	0.05% - 0.24%
ORAL; DROPS		3		05/25/95	UNK	0.18%
ORAL; GRANULE		4		06/15/88	600	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER		2		12/05/88	510	0.0005%
ORAL; POWDER, FOR RECONSTITUTION		28		12/23/93	530	
ORAL; SOLUTION		32		11/17/95	530	0.0121% - 0.65%
ORAL; SOLUTION, ELIXIR		15		10/27/92	600	0.0299% - 0.15%
ORAL; SUSPENSION		30		06/16/95	UNK	0.1% - 0.2%
ORAL; SYRUP		79		09/25/95	600	0.1% - 7.3519%
ORAL; TABLET		33		03/20/92	600	0.06MG - 20.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		5		11/16/94	UNK	0.2MG - 2.13MG
ORAL; TABLET, COATED		2		04/08/81	600	
ORAL; TABLET, FILM COATED		9		12/23/91	510	1.25MG - 20.0MG
ORAL; TABLET, SUSTAINED ACTION		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CITRIC ACID	000077929					
ORAL-28; TABLET		1				
OTIC; SOLUTION		4		10/31/94	600	0.05% - 0.2%
OTIC; SOLUTION, DROPS		1				
SOFT TISSUE; INJECTION		2		02/27/79	600	0.205%
SUBLINGUAL; TABLET		2		08/11/81	600	
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		32		06/13/95	600	0.001% - 0.85%
TOPICAL; GEL		1				
TOPICAL; LOTION		9		01/24/92	600	0.21% - 0.85%
TOPICAL; OINTMENT		3		09/30/83	UNK	0.001%
TOPICAL; POWDER, FOR RECONSTITUTION		1				
TOPICAL; SHAMPOO		2		08/27/90	UNK	0.075%
TOPICAL; SOLUTION		24		02/27/95	600	0.005% - 0.4%
TOPICAL; SUSPENSION, SHAMPOO		3		01/10/91	600	0.77758%
TOPICAL; SWAB		2		07/23/87	600	0.04%
VAGINAL; EMULSION, CREAM		1				
CITRIC ACID MONOHYDRATE	005949291					
IM - IV - SC; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRAVENOUS; INJECTION		3		08/18/95	110	0.075%
IV(INFUSION); INJECTION		6		08/18/95	110	0.05% - 0.075%
NASAL; SOLUTION		1				
NERVE BLOCK; INJECTION		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		2		06/10/87	UNK	
TOPICAL; EMULSION, CREAM		2		02/16/94	600	0.05%
TOPICAL; GEL		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
VAGINAL; EMULSION, CREAM		1				
CITRIC ACID, ANHYDROUS						
IM - IV; INJECTION		1				
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
TOPICAL; EMULSION, CREAM		1				
CITRIC ACID, HYDROUS	015686654					
INTRAVENOUS; INJECTION		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
TOPICAL; EMULSION, CREAM		4		06/17/94	UNK	0.02% - 0.05%
CLOVE OIL	008000348					
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		8		12/16/83	600	0.008%
ORAL; SUSPENSION		2		12/27/91	600	0.001%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
COCAMIDE DIETHANOLAMINE	061791319					
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		4		04/15/91	600	2.0% - 4.0%
TOPICAL; SPONGE		2		02/28/91	600	2.0MG
COCAMIDE ETHER SULFATE						
TOPICAL; SHAMPOO		1				
COCAMINE OXIDE	061788907					
TOPICAL; SHAMPOO		1				
COCOA BUTTER						
RECTAL; SUPPOSITORY		4		10/04/83	600	1.868GM - 2.0265GM
COCOA BUTTER (POND'S TYPE 520A)						
RECTAL; SUPPOSITORY		1				
TOPICAL; LOTION		1				
COCOAMPHOCARBOXYGLYCINATE						
TOPICAL; SUSPENSION, SHAMPOO		1				
COCOGLYCERIDES						
TOPICAL; EMULSION, CREAM		1				
COCONUT OIL	008001318					
TOPICAL; EMULSION, CREAM		1				
TOPICAL; OINTMENT		2		09/30/94	600	
COCONUT OIL, HYDROGENATED						
RECTAL; SUPPOSITORY		3		11/24/93	600	
COLORING SUSPENSION						
ORAL; TABLET		1				
TOPICAL; EMULSION, CREAM		1				
CONFECTIONERS GLAZE						
ORAL; CAPSULE, SUSTAINED ACTION		1				
CORIANDER OIL	008008524					
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		8		12/16/83	600	0.003%
CORN GLYCERIDES						
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; SOLUTION		1				
CORN OIL	008001307					
ORAL; CAPSULE		6		06/29/93	600	205.0MG - 918.0MG
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; TABLET		15		03/21/77	600	0.02GM
ORAL; TABLET, COATED		1				
CORN OIL PEG-6 ESTERS						
ORAL; CAPSULE, SOFT GELATIN		1				
CORN SYRUP						
ORAL; SYRUP		7		10/28/94	600	
ORAL; TABLET		3		12/03/73	600	8.0MG - 14.065MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CORN SYRUP						
ORAL; TABLET, UNCOATED, TROCHE		1				
COTTONSEED OIL	008001294					
INTRAMUSCULAR; INJECTION		7		03/13/86	600	56.0% - 91.66%
COTTONSEED OIL, HYDROGENATED						
ORAL; CAPSULE		3		08/04/86	600	2.5MG - 4.25MG
ORAL; TABLET		46		08/30/91	600	0.002MG - 26.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		5		08/14/87	600	32.4MG - 402.0MG
CREAM BASE						
TOPICAL; EMULSION, CREAM		2		05/15/85	600	
CREATINE	000057001					
INTRA-ARTICULAR; INJECTION		1				
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
CREATININE	000060275					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		4		07/01/82	600	0.8%
INTRA-ARTICULAR; INJECTION		4		05/24/82	600	0.5% - 0.8%
INTRABURSAL; INJECTION		1				
INTRADERMAL; INJECTION, SUSTAINED ACTION		1				
INTRALESIONAL; INJECTION		3		06/19/80	600	0.8%
INTRAMUSCULAR; INJECTION		1				
INTRASYNOVIAL; INJECTION		2		03/01/77	UNK	0.8%
OPHTHALMIC; SOLUTION		1				
OTIC; SOLUTION		1				
SOFT TISSUE; INJECTION		3		05/24/82	600	0.5% - 0.8%
TOPICAL; EMULSION, CREAM		1				
CRESOL, M-	000108394					
IM - SC; INJECTION		2		01/23/85	510	0.158% - 0.25%
INTRADERMAL; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
SUBCUTANEOUS; INJECTION		10		03/31/94	510	0.15% - 0.31%
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
SUBCUTANEOUS; SUSPENSION, INJECTION		1				
CROSCARMELLOSE SODIUM						
ORAL; CAPSULE		59		07/03/95	600	0.4MG - 21.0MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; TABLET		450		11/09/95	600	0.013GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		3		12/30/92	110	4.75MG - 33.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		8		11/30/95	600	2.0MG - 32.44MG
ORAL; TABLET, FILM COATED		22		12/12/95	120	1.5MG - 40.0MG
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CROSCARMELLOSE SODIUM						
SUBLINGUAL; TABLET		5		09/18/86	600	1.8MG - 6.5MG
CROSPOVIDONE	009003398					
IMPLANTATION; PELLET		1				
ORAL; CAPSULE		6		03/03/95	110	1.5MG - 20.0MG
ORAL; CAPSULE, SUSTAINED ACTION		12		04/25/85	UNK	0.5MG - 10.71MG
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; SUSPENSION		1				
ORAL; TABLET		131		12/22/95	600	1.3MG - 180.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		14		02/28/95	600	1.64MG - 28.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		14		09/29/95	600	2.0MG - 55.3MG
ORAL; TABLET, SUSTAINED ACTION		7		12/15/89	110	3.0MG - 144.0MG
ORAL-21; TABLET		1				
TOPICAL; LOTION		1				
VAGINAL; SUPPOSITORY		1				
VAGINAL; TABLET		1				
CUPRIC SULFATE	007758998					
OTIC; SOLUTION		1				
OTIC; SUSPENSION		1				
CUPRIC SULFATE, ANHYDROUS	007758987					
OTIC; SOLUTION		1				
CYCLOMETHICONE						
ORAL; POWDER, FOR RECONSTITUTION		1				
TOPICAL; CREAM, AUGMENTED		1				
CYSTEINE						
IM - SC; INJECTION, SUSTAINED ACTION		1				
CYSTEINE HYDROCHLORIDE	007048046					
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
ORAL; CAPSULE		1				
DC ANTIFOAM AF TRITURATION 1% ON SUCROSE						
ORAL; POWDER, FOR RECONSTITUTION		1				
DEHYDROACETIC ACID	000520456					
TOPICAL; LOTION		2		11/26/85	600	0.1%
DEHYMULS E						
TOPICAL; OINTMENT		2		12/17/90	UNK	5.0% - 7.5%
DENATONIUM BENZOATE	003734336					
TOPICAL; GEL		1				
DEOXYCHOLIC ACID	000083443					
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
DEXTRATES						
ORAL; TABLET		2		11/30/95	600	54.0MG - 86.5MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DEXTRIN	009004539					
ORAL; TABLET		3		11/13/84	600	1.56MG - 1.8MG
TOPICAL; EMULSION, CREAM		3		09/28/92	600	
TOPICAL; SHAMPOO		1				
DEXTRINS MODIFIED						
ORAL-28; TABLET		2		04/30/73	510	
DEXTROSE	005996101					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		3		07/23/86	520	3.6% - 5.0%
IM - IV; POWDER, FOR INJECTION SOLUTION		5		12/27/91	600	
IM - SC; INJECTION		2		07/25/83	600	3.75%
INTERSTITIAL; INJECTION		1				
INTRACAVITARY; INJECTION		1				
INTRAMUSCULAR; INJECTION		3		05/24/79	600	4.4% - 5.0%
INTRAPERITONEAL; INJECTION		1				
INTRAPLEURAL; INJECTION		1				
INTRAVENOUS; INJECTION		13		07/30/93	520	2.0% - 5.0%
INTRAVENOUS; SOLUTION, INJECTION		2		01/22/92	UNK	5.0%
IV(INFUSION); INJECTION		39		01/31/95	180	1.4% - 5.0%
IV(INFUSION); SOLUTION, INJECTION		3		12/29/92	520	2.2% - 3.8%
NASAL; SPRAY, METERED		3		10/19/94	UNK	5.0%
ORAL; CONCENTRATE		2		10/16/87	600	5.0% - 14.32708%
ORAL; PASTILLE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SYRUP		1				
ORAL; TABLET		6		09/27/93	600	72.459MG - 150.0MG
ORAL; TABLET, UNCOATED, TROCHE		1				
SPINAL; INJECTION		6		12/11/87	600	7.5% - 8.25%
SUBLINGUAL; TABLET		1				
DEXTROSE SOLUTION	008012246					
INTRAVENOUS; INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); SOLUTION, INJECTION		1				
ORAL; SYRUP		1				
DEXTROSE, ANHYDROUS	000050997					
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; SOLUTION, INJECTION		1				
IV(INFUSION); INJECTION		5		08/09/94	110	4.5% - 4.94%
IV(INFUSION); SOLUTION, INJECTION		1				
DI-PAC (97% SUCROSE-3% MODIFIED DEXTRINS						
ORAL; TABLET		5		03/18/80	600	51.0MG - 322.35MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		10/04/76	600	162.674MG - 260.12MG
ORAL; TABLET, COATED		3		03/07/77	600	24.8MG - 105.4MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DIACETYLATED MONOGLYCERIDES	008029923					
ORAL; TABLET		48		11/05/92	600	0.00003ML - 0.00006ML
ORAL; TABLET, COATED		2		04/08/81	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		6		05/31/91	600	0.04MG - 1.1MG
DIATOMACEOUS EARTH						
ORAL; TABLET, SUSTAINED ACTION		1				
DIATRIZOIC ACID	000117964					
INTRA-ARTERIAL; INJECTION		2		08/29/74	160	
INTRA-ARTICULAR; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRADISCAL; INJECTION		1				
INTRAUTERINE; SOLUTION		1				
INTRAVENOUS; INJECTION		2		08/29/74	160	
IV(INFUSION); INJECTION		1				
PERIARTICULAR; INJECTION		1				
URETERAL; SOLUTION		1				
DIAZOLYDINYLUREA						
TOPICAL; EMULSION, CREAM		2		12/27/90	UNK	0.2%
TOPICAL; LOTION		1				
DIBUTYL PHTHALATE	000084742					
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
DIBUTYL SEBACATE	000109433					
ORAL; CAPSULE, SUSTAINED ACTION		2		08/19/88	600	
ORAL; GRANULE, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		2		10/03/90	600	
ORAL; TABLET, SUSTAINED ACTION		1				
DICHLORODIFLUOROMETHANE	000075718					
INHALATION; AEROSOL, METERED		24		12/28/95	600	11.25% - 68.5611%
NASAL; AEROSOL SPRAY		1				
NASAL; AEROSOL, METERED		4		02/14/94	UNK	99.789%
ORAL; AEROSOL		1				
RECTAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; AEROSOL		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
DICHLOROFLUOROMETHANE	000075434					
ORAL; AEROSOL SPRAY		1				
DICHLOROTETRAFLUOROETHANE	000076142					
INHALATION; AEROSOL, METERED		15		12/30/92	UNK	21.857% - 51.12%
NASAL; AEROSOL SPRAY		1				
NASAL; AEROSOL, METERED		1				
ORAL; AEROSOL		1				
RECTAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; AEROSOL		2		05/17/88	600	50.0%
TOPICAL; EMULSION, AEROSOL FOAM		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DICYCLOHEXYL-CARBODIIMIDE						
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
DIETHANOLAMINE	000111422					
IV(INFUSION); INJECTION		8		09/11/92	600	0.3% - 0.35%
DIETHYL PHTHALATE	000084662					
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, SUSTAINED ACTION		7		01/04/95	600	8.0MG - 12.0MG
DIETHYL SEBACATE	000110407					
TOPICAL; EMULSION, CREAM		1				
DIETHYLAMINE						
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
DIGLYCERIDES						
TOPICAL; LOTION		1				
DIGLYCOL STEARATE	000106116					
VAGINAL; EMULSION, CREAM		1				
DIHYDROXYALUMINUM SODIUM CARBONATE	000539684					
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
DIISOPROPANOLAMINE	000110974					
TOPICAL; GEL		1				
TOPICAL; SOLUTION		1				
DIISOPROPYL ADIPATE	006938949					
TOPICAL; LOTION		2		06/05/78	600	
TOPICAL; SOLUTION		4		02/27/95	600	10.0%
DIISOPROPYLBENZOTHIAZYL-2-SULFENAMIDE						
ORAL; TABLET		1				
DIMETHICONE	009006659					
ORAL; CAPSULE		4		08/29/95	600	2.5MG - 8.2MG
ORAL; TABLET		3		08/17/88	600	0.2MG
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; EMULSION, CREAM		2		08/26/59	UNK	
TOPICAL; LOTION		2		12/07/92	UNK	0.4%
DIMETHICONE 350	009006659					
ORAL; CAPSULE		1				
TOPICAL; EMULSION, CREAM		3		01/24/80	UNK	1.0%
DIMETHICONE 360						
TOPICAL; EMULSION, CREAM		1				
DIMETHYLDIOCTADECYLAMMONIUM BENTONITE						
RECTAL; SUSPENSION		1				
DIMYRISTOYL LECITHIN	018194246					
IV(INFUSION); SUSPENSION, INJECTION		1				
DIMYRISTOYL PHOSPHATIDYLGLYCEROL, L-	057618287					
IV(INFUSION); SUSPENSION, INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DIOCTYLPHTHALATE						
OPHTHALMIC; DRUG DELIVERY SYSTEM		1				
OPHTHALMIC; SUPPOSITORY, INSERT, CONTROLLED RELEASE		1				
DIPROPYLENE GLYCOL	000110985					
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
DISODIUM EDISYLATE						
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
ORAL; TABLET		1				
DISODIUM MONOOLEAMIDE SULFASUCCINATE						
TOPICAL; SHAMPOO		1				
DISOFENIN	065717977					
IV(INFUSION); INJECTION		1				
DOCUSATE	010041197					
ORAL; TABLET		1				
DOCUSATE SODIUM	000577117					
INTRAMUSCULAR; INJECTION		1				
ORAL; CAPSULE		11		08/19/91	600	0.4MG - 2.0MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; GRANULE FOR RECONSTITUTION, CR		1				
ORAL; SUSPENSION		2		01/26/84	600	0.04%
ORAL; TABLET		47		12/29/95	600	0.002MG - 11.0MG
TOPICAL; GEL		2		10/26/84	UNK	
TOPICAL; SHAMPOO		1				
DOCUSATE SODIUM/SODIUM BENZOATE						
ORAL; CAPSULE		4		06/03/87	600	0.425MG - 4.0MG
ORAL; TABLET		16		10/20/95	600	0.2MG - 6.0MG
DRI KLEAR 042						
ORAL; TABLET		1				
ORAL; TABLET, COATED		1				
DRY FLO						
ORAL; TABLET		1				
DURO-TAK 280-2516						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
DURO-TAK 80-1196						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
DUSTING POWDER						
ORAL; TABLET		3		04/14/83	600	
ORAL; TABLET, COATED		1				
DYE BEIGE P-1437						
ORAL; TABLET		1				
DYE BLACK						
ORAL; CAPSULE		1				
ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE BLACK LB-442						
ORAL; TABLET		1				
DYE BLUE						
ORAL; CAPSULE		2		09/10/80	600	0.014MG
ORAL; TABLET		1				
RECTAL; SUPPOSITORY		1				
DYE BLUE #1						
ORAL; CAPSULE		3		05/31/90	520	
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		1				
DYE BLUE #2						
ORAL; TABLET, SUSTAINED ACTION		1				
DYE BROWN LAKE						
ORAL; TABLET		1				
DYE BROWN LB-292						
ORAL; TABLET		1				
DYE BROWN LB-464						
ORAL; TABLET		1				
DYE CARAMEL						
ORAL; SYRUP		4		07/26/88	UNK	0.025% - 0.05%
DYE CARAMEL ACID PROOF 100						
ORAL; SYRUP		1				
DYE DC BLUE #2 LAKE	000130201					
ORAL; CAPSULE		2		04/21/87	600	
ORAL; TABLET, COATED		1				
DYE DC BLUE #6	000482893					
ORAL; CAPSULE		2		09/21/77	600	
DYE DC GREEN #3 LAKE						
ORAL; CAPSULE		1				
ORAL; TABLET, COATED		1				
DYE DC GREEN #5	004403901					
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		2		04/10/78	600	0.0009%
ORAL; TABLET, COATED		1				
ORAL-21; TABLET		2		04/13/84	510	0.0024MG
ORAL-28; TABLET		3		04/13/84	510	0.0024MG
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
DYE DC RED #19	000081889					
ORAL; SUSPENSION		1		--		
DYE DC RED #21 LAKE						
ORAL; CAPSULE		1				
DYE DC RED #22	000548265					
ORAL; CAPSULE		9		09/11/92	530	
ORAL; CAPSULE, SUSTAINED ACTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE DC RED #27	002134158					
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		8		01/02/87	600	0.5UGM
ORAL; TABLET, COATED		1				
DYE DC RED #27 ALUMINUM LAKE						
ORAL; TABLET		4		03/11/88	600	
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		2		11/21/77	600	
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-28; TABLET		1				
DYE DC RED #28	004618239					
ORAL; CAPSULE		81		07/03/95	600	0.002MG
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		3		10/05/95	180	
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		9		09/11/95	110	
ORAL; POWDER, FOR RECONSTITUTION		2		04/22/68	520	0.014%
ORAL; TABLET		2		08/31/90	520	
DYE DC RED #3 LAKE						
ORAL; TABLET		2		08/06/79	600	0.2MG
ORAL; TABLET, FILM COATED		1				
DYE DC RED #30	002379740					
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		16		04/27/95	600	0.0105MG - 0.93MG
ORAL; TABLET, COATED		3		12/20/82	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		4		12/29/92	120	
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-21; TABLET		2		12/30/91	600	
ORAL-28; TABLET		5		12/30/91	600	
DYE DC RED #30 ALUMINUM LAKE						
ORAL; TABLET		26		01/31/95	600	0.0324MG - 3.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		5		08/30/83	600	0.29GM
ORAL-21; TABLET		1				
DYE DC RED #30 LAKE						
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; TABLET		6		10/30/92	600	0.04MG - 0.4MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, SUSTAINED ACTION		2		11/05/81	600	
ORAL-28; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE DC RED #33	003567666					
ORAL; CAPSULE		28		03/03/96	110	0.0001MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		4		07/03/85	110	0.0013MG
ORAL; CONCENTRATE		1				
ORAL; SOLUTION		2		04/27/88	510	
ORAL; SOLUTION, ELIXIR		7		12/16/83	600	
ORAL; SUSPENSION		2		12/18/80	600	
ORAL; SYRUP		13		03/07/85	600	0.001848%
ORAL; TABLET		8		06/16/88	110	0.007MG - 0.15MG
TOPICAL; SHAMPOO		1				
DYE DC RED #33 LAKE						
ORAL; CAPSULE		3		02/25/88	600	1.79MG
ORAL; SYRUP		3		11/22/85	600	
DYE DC RED #36						
ORAL; TABLET		3		09/07/77	110	0.01MG - 48.75MG
TOPICAL; EMULSION, CREAM		1				
TOPICAL; OINTMENT		1				
DYE DC RED #39	006371557					
TOPICAL; LOTION		1				
DYE DC RED #4 LAKE						
ORAL; CAPSULE		1				
DYE DC RED #40 LAKE						
ORAL; CAPSULE		4		03/02/92	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; TABLET		4		01/31/95	600	
ORAL; TABLET, COATED		1				
RECTAL; SUPPOSITORY		1				
DYE DC RED #6	005858811					
ORAL; TABLET, COATED		1				
DYE DC RED #6 LAKE						
ORAL; TABLET		2		03/01/90	180	0.4MG
DYE DC RED #7	005281049					
ORAL; TABLET		10		12/21/93	600	0.16MG - 0.6MG
ORAL; TABLET, COATED		2		01/26/84	510	
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
DYE DC RED #7 CALCIUM LAKE						
ORAL; CAPSULE		2		12/20/90	110	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		14		12/21/93	600	
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE DC RED #7 CALCIUM LAKE						
ORAL; TABLET, SUSTAINED ACTION		2		01/16/85	600	
DYE DC RED #7 LAKE						
ORAL; TABLET		4		01/25/88	600	0.02MG - 0.6MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
DYE DC RED LAKE						
ORAL; TABLET		6		09/25/95	600	0.009MG - 2.4MG
DYE DC VIOLET #2 LAKE						
ORAL; TABLET		1				
DYE DC YELLOW						
ORAL; CAPSULE		3		07/31/92	600	
DYE DC YELLOW #10	008004920					
BUCCAL; GUM, CHEWING		1				
DENTAL; GEL		1				
DENTAL; PASTE		1				
ORAL; CAPLET		1				
ORAL; CAPSULE		169		08/31/95	600	2.0UGM
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		2		01/29/93	600	
ORAL; CAPSULE, COATED PELLETS		2		06/30/92	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		3		03/27/95	600	
ORAL; CAPSULE, SOFT GELATIN		2		07/26/82	110	0.111MG
ORAL; CAPSULE, SUSTAINED ACTION		16		05/10/93	180	0.25MG
ORAL; CONCENTRATE		3		01/30/92	600	0.0008% - 0.0014%
ORAL; POWDER		2		12/05/88	510	0.00003%
ORAL; SOLUTION		7		06/30/93	600	0.0005% - 0.005%
ORAL; SOLUTION, ELIXIR		3		08/28/92	600	
ORAL; SUSPENSION		9		06/16/95	UNK	0.00027% - 0.0015%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; SYRUP		13		08/17/90	UNK	0.0028% - 0.0036%
ORAL; TABLET		131		06/29/95	600	0.0035MG - 5.0MG
ORAL; TABLET, COATED		8		04/26/78	600	2.5MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		19		04/28/95	110	0.006MG - 0.068MG
ORAL; TABLET, SUSTAINED ACTION		8		11/08/93	UNK	0.071MG - 2.01MG
ORAL-21; TABLET		6		12/30/91	600	0.02MG
ORAL-28; TABLET		7		12/30/91	600	0.02MG - 0.06MG
RECTAL; SUPPOSITORY		1				
SUBLINGUAL; TABLET		3		02/26/88	600	0.028MG - 0.23MG
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
TOPICAL; SHAMPOO		1				
DYE DC YELLOW #10 ALUMINUM LAKE						
ORAL; CAPLET		1				
ORAL; CAPSULE		7		08/15/94	180	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; TABLET		72		10/17/95	600	0.008MG - 2.5MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE DC YELLOW #10 ALUMINUM LAKE						
ORAL; TABLET, COATED		8		03/07/77	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		7		04/14/95	110	0.0075MG - 0.045MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	
ORAL; TABLET, SUSTAINED ACTION		6		07/31/92	600	
ORAL-21; TABLET		2		01/29/87	600	0.096MG - 0.415MG
ORAL-28; TABLET		12		07/03/92	510	0.096MG - 0.415MG
DYE DC YELLOW #10 HT LAKE						
ORAL; TABLET		5		06/13/88	600	0.1MG - 1.4MG
DYE DC YELLOW #10 LAKE						
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET		74		08/29/95	600	0.000289MG - 5.71MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		3		04/20/76	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		5		05/20/83	600	0.12GM
ORAL; TABLET, SUSTAINED ACTION		3		07/29/88	110	0.18MG - 2.33MG
SUBLINGUAL; TABLET		1				
DYE DC YELLOW #5 LAKE						
ORAL; TABLET		8		08/11/80	600	0.016MG - 2.69MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
DYE DC YELLOW #6						
ORAL; CAPSULE		3		12/28/90	600	
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET		3		01/12/73	600	0.005MG
ORAL-21; TABLET		1				
DYE DC YELLOW #6 LAKE						
ORAL; CAPSULE		1				
ORAL; SOLUTION		1				
ORAL; TABLET		15		11/02/87	600	0.27UGM
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		1				
DYE FDC BLUE #1	002650182					
DENTAL; PASTE		1				
DENTAL; SOLUTION		4		12/28/95	600	0.00005% - 0.0015%
ORAL; CAPSULE		237		10/18/95	600	5.0UGM
ORAL; CAPSULE, COATED PELLETS		2		06/30/92	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		4		10/05/95	180	
ORAL; CAPSULE, HARD GELATIN		6		03/27/95	600	3.708MG
ORAL; CAPSULE, SOFT GELATIN		3		03/08/94	180	
ORAL; CAPSULE, SUSTAINED ACTION		28		09/11/95	110	0.019MG - 0.9MG
ORAL; POWDER, FOR RECONSTITUTION		2		04/23/79	600	
ORAL; SOLUTION		5		05/28/93	600	0.000075% - 0.75%
ORAL; SOLUTION, ELIXIR		2		04/29/92	600	
ORAL; SUSPENSION		2		02/12/86	520	0.005%
ORAL; SYRUP		16		12/23/88	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE FDC BLUE #1	002650182					
ORAL; TABLET		129		06/29/95	600	5.0UGM - 11.4UGM
ORAL; TABLET, COATED		9		02/11/82	120	0.0052MG - 0.0085MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		10		10/31/91	180	0.51UGM
ORAL; TABLET, SUSTAINED ACTION		6		11/08/93	UNK	0.03MG
ORAL-21; TABLET		3		07/01/88	600	0.01MG - 0.05MG
ORAL-28; TABLET		6		07/01/88	600	0.01MG - 0.05MG
RECTAL; SOLUTION		3		05/28/93	600	0.000075%
SUBLINGUAL; TABLET		1				
TOPICAL; SHAMPOO		2		08/27/90	UNK	0.0036%
TOPICAL; SUSPENSION, SHAMPOO		1				
TOPICAL; SWAB		1				
DYE FDC BLUE #1 ALUMINUM LAKE						
ORAL; CAPSULE		12		08/15/94	180	
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SYRUP		1				
ORAL; TABLET		114		12/29/95	600	0.001MG - 2.5MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		2		04/20/76	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		05/15/90	600	
ORAL; TABLET, FILM COATED		4		05/30/95	600	0.1MG - 0.2MG
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		4		07/29/88	110	0.009MG - 0.29MG
ORAL-28; TABLET		1				
DYE FDC BLUE #1 H.T. ALUMINUM LAKE						
ORAL; TABLET		11		08/31/88	600	0.012MG - 0.3MG
DYE FDC BLUE #1 LAKE						
ORAL; CAPSULE		1				
ORAL; TABLET		35		08/29/95	600	0.00364MG - 2.675MG
ORAL; TABLET, COATED		2		04/08/81	600	
ORAL; TABLET, FILM COATED		1				
ORAL-28; TABLET		2		12/30/94	510	0.045MG
DYE FDC BLUE #10						
ORAL; TABLET		1				
DYE FDC BLUE #2	000860220					
BUCCAL; TABLET		1				
ORAL; CAPSULE		15		10/18/95	600	0.012MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		2		05/03/95	550	
ORAL; CAPSULE, SUSTAINED ACTION		3		02/21/92	110	2.1UGM
ORAL; TABLET		75		06/29/95	600	0.0015MG - 20.016MG
ORAL; TABLET, COATED		10		02/25/92	600	24.12MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		11		12/29/92	120	0.7MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE FDC BLUE #2	000860220					
ORAL; TABLET, SUSTAINED ACTION		5		09/22/94	110	
DYE FDC BLUE #2 LAKE	012227859					
ORAL; CAPSULE		11		08/15/96	180	
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		77		12/29/95	600	0.035MG - 6.0MG
ORAL; TABLET, COATED		14		12/20/82	600	0.095MG - 0.1425MG
ORAL; TABLET, FILM COATED		9		06/23/95	530	0.015MG - 0.3MG
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		7		01/09/87	600	0.02MG
ORAL; TABLET, UNCOATED, TROCHE		1				
ORAL-21; TABLET		3		07/03/92	510	0.1MG - 0.208MG
ORAL-28; TABLET		12		11/17/95	510	0.0649MG - 0.208MG
DYE FDC GREEN #3	002353459					
DENTAL; GEL		1				
ORAL; CAPSULE		45		05/03/95	520	0.003MG - 0.066MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		6		09/11/95	110	0.015MG
ORAL; SOLUTION, ELIXIR		1				
ORAL; SYRUP		4		08/17/90	UNK	0.00032%
ORAL; TABLET		6		09/15/83	600	0.0004MG - 10.0MG
ORAL; TABLET, COATED		2		03/22/60	110	0.005MG
RECTAL; SUPPOSITORY		1				
DYE FDC GREEN #6						
ORAL; CAPSULE		1				
DYE FDC RED #27 LAKE						
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		3		06/19/85	600	0.06MG
DYE FDC RED #28						
ORAL; CAPSULE		13		03/29/91	600	
ORAL; POWDER, FOR RECONSTITUTION		1				
DYE FDC RED #3	016423680					
ORAL; CAPSULE		84		10/18/95	600	0.00036MG - 0.333MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		2		03/28/67	UNK	
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		11		01/26/89	600	0.037MG - 0.3MG
ORAL; DROPS		2		12/18/80	600	
ORAL; GRANULE		2		12/18/80	600	
ORAL; POWDER, FOR RECONSTITUTION		12		12/23/91	520	0.004% - 0.525%
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		2		09/23/74	520	
ORAL; TABLET		44		08/31/90	520	2.5UGM
ORAL; TABLET, COATED		4		08/16/85	120	0.005MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		2		09/28/77	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE FDC RED #3	016423680					
ORAL; TABLET, SUSTAINED ACTION		2		08/30/84	110	
ORAL-28; TABLET		1				
TOPICAL; SHAMPOO		1				
DYE FDC RED #3 LAKE						
ORAL; TABLET		13		11/10/88	600	0.006MG - 1.2MG
ORAL; TABLET, SUSTAINED ACTION		1				
DYE FDC RED #3-ALUMINUM LAKE	012227780					
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; GRANULE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET		32		05/11/90	600	0.02MG - 8.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		10/21/80	520	0.126MG - 4.25MG
ORAL; TABLET, FILM COATED		1				
TOPICAL; SOLUTION		1				
DYE FDC RED #30 LAKE						
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		4		12/21/87	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
DYE FDC RED #33						
ORAL; CAPSULE		4		03/06/87	600	262.0MG
ORAL; SYRUP		5		09/17/93	530	0.002%
DYE FDC RED #40						
BUCCAL; TABLET		1				
ORAL; CAPSULE		128		10/18/95	600	0.0074MG - 73.2MG
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		10/05/95	180	
ORAL; CAPSULE, HARD GELATIN		2		01/28/91	110	
ORAL; CAPSULE, SOFT GELATIN		2		07/26/82	110	0.074MG
ORAL; CAPSULE, SUSTAINED ACTION		17		09/11/95	110	0.029MG - 0.129MG
ORAL; CONCENTRATE		3		11/30/94	600	0.0025% - 0.0075%
ORAL; DROPS		2		05/25/95	UNK	0.001%
ORAL; POWDER		1				
ORAL; POWDER, FOR RECONSTITUTION		31		10/19/95	520	0.00001% - 0.032%
ORAL; SOLUTION		13		12/22/94	600	0.0004% - 0.0007%
ORAL; SOLUTION, ELIXIR		14		10/27/92	600	0.0012% - 0.005%
ORAL; SUSPENSION		21		09/15/95	180	0.004% - 0.9%
ORAL; SYRUP		36		01/13/95	600	0.002% - 0.04%
ORAL; TABLET		60		10/17/95	600	0.55UGM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		10/04/76	600	0.2MG
ORAL; TABLET, COATED		6		08/16/77	180	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		10/26/84	120	0.043MG
ORAL; TABLET, FILM COATED		8		02/25/94	600	0.028MG
ORAL; TABLET, SUSTAINED ACTION		3		05/14/85	UNK	
SUBLINGUAL; TABLET		2		07/29/88	110	0.003MG
TOPICAL; SOLUTION		3		01/28/92	600	0.6%