INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT / ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
GLYCERIN	000056815					
ORAL; CAPSULE, SOFT GELATIN		12		11/22/95	150	3.055MG - 111.0MG
ORAL; CAPSULE, SUSTAINED ACTION		3		04/25/95	UNK	0.1MG
ORAL; CONCENTRATE		18		11/30/94	600	5.0% - 86.6%
ORAL; DROPS		1				
ORAL; SOLUTION		37		11/1/95	600	2.5% - 20.0%
ORAL; SOLUTION, ELIXIR		15		04/29/93	600	2.5% - 5.0%
ORAL; SUSPENSION		33		06/16/95	UNK	1.0% - 34.3%
ORAL; SYRUP		54		07/30/93	600	5.0% - 50.0%
ORAL; TABLET		9		02/21/95	600	0.04ML
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		07/29/92	600	0.75MG - 1.0MG
ORAL; TABLET, FILM COATED		2		11/18/83	600	
ORAL; TABLET, SUSTAINED ACTION		2		08/19/91	UNK	1.2MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
OTIC; SOLUTION		5		12/29/95	600	52.62%
OTIC; SUSPENSION		2		05/25/75	600	0.05%
PERFUSION, BILIARY; LIQUID		1				
RECTAL; SUPPOSITORY		5		11/24/93	600	
SUBCUTANEOUS; INJECTION		13		03/31/94	510	1.6% - 32.5%
SUBCUTANEOUS; SUSPENSION, INJECTION		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		41		09/20/95	UNK	0.2% - 21.0%
TOPICAL; LOTION		9		05/31/89	UNK	3.0% - 10.0%
TOPICAL; OINTMENT		2		03/09/78	600	5.0%
TOPICAL; SOLUTION		4		01/28/92	600	3.43% - 65.7%
TOPICAL; SPONGE		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
VAGINAL; EMULSION, CREAM		4		09/19/85	600	5.0% - 7.0%
VAGINAL; SUPPOSITORY		3		01/27/87	520	227.9MG
GLYCERIN HYDROCHLORIDE						
ORAL; TABLET		1				
GLYCEROL ESTER OF HYDROGENATED ROSIN	008050291					
NASAL; OINTMENT		1				
GLYCERYL BEHENATE						
ORAL; TABLET		2		11/22/91	600	
GLYCERYL DISTEARATE	001323837					
ORAL; CAPSULE, SUSTAINED ACTION		4		01/06/81	UNK	6.6MG - 39.2MG
GLYCERYL LAURATE	001322889					
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
GLYCERYL OLEATE	000544763					
ORAL; CAPSULE		2		01/27/81	120	
ORAL; CAPSULE, HARD GELATIN		2		03/28/67	UNK	
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET, COATED		1				
ORAL-28; TABLET		2		11/17/95	510	0.15MG
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
GLYCERYL OLEATE/PROPYLENE GLYCOL						
TOPICAL; CREAM, AUGMENTED		1				
GLYCERYL PALMITATE	001330730					
RECTAL; SUPPOSITORY		2		04/30/73	120	
GLYCERYL RICINOLEATE	001323382					
TOPICAL; SUSPENSION, SHAMPOO		4		01/10/91	600	1.0% - 2.0%
GLYCERYL STEARATE	031566311					
OPHTHALMIC; SUSPENSION		2		05/11/88	600	0.5%
ORAL; CAPSULE, SUSTAINED ACTION		5		08/02/76	120	0.823MG - 27.0MG
ORAL; TABLET, SUSTAINED ACTION		5		01/04/95	600	52.86MG - 154.0MG
OTIC; SUSPENSION		2		09/29/87	600	0.05% - 0.5%
RECTAL; SUPPOSITORY		7		11/24/93	600	6.0MG - 36.85MG
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		44		09/13/95	UNK	0.3% - 20.0%
TOPICAL; LOTION		12		12/07/92	UNK	0.25% - 3.0%
TOPICAL; OINTMENT		4		10/10/85	600	5.0%
VAGINAL; EMULSION, CREAM		12		12/21/95	520	2.0% - 17.0%
GLYCERYL STEARATE SE						
TOPICAL; LOTION		1				
GLYCERYL STEARATE-STEARAMIDOETHYL DIETHYLAMINE						
TOPICAL; OINTMENT		1				
GLYCERYL STEARATE/PEG-100 STEARATE						
TOPICAL; EMULSION, CREAM		2		04/01/94	UNK	5.0%
TOPICAL; LOTION		2		11/26/85	600	1.42%
GLYCERYL STEARATE/PEG-40 STEARATE						
RECTAL; SUPPOSITORY		4		02/27/95	600	15.0MG - 35.0MG
GLYCINE	000056406					
INTRAMUSCULAR; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
ORAL; CAPSULE		1				
ORAL; POWDER, FOR RECONSTITUTION		2		12/23/91	520	2.1%
ORAL; SOLUTION		1				
ORAL; TABLET		3		06/01/84	600	8.0MG - 163.31MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		07/22/85	520	100.0MG - 200.0MG
RECTAL; SOLUTION		1				
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		2		11/17/93	510	
GLYCOL STEARATE	000111604					
TOPICAL; SUSPENSION, SHAMPOO		1				
GLYCYRRHIZA						
ORAL; POWDER, FOR RECONSTITUTION		1				
GLYCYRRHIZIN, AMMONIATED	001407030					
ORAL; GRANULE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
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
GUANIDINE HYDROCHLORIDE	000050011					
INTRAVENOUS; INJECTION		2		02/23/76	160	0.04%
GUAR GUM	009000300					
BUCCAL/SUBLINGUAL; TABLET		1				
ORAL; POWDER, FOR RECONSTITUTION		2		04/18/91	600	0.2%
ORAL; SUSPENSION		1				
ORAL; TABLET		8		12/01/86	600	1.296MG - 12.0MG
ORAL; TABLET, FILM COATED		4		01/06/78	110	6.0MG - 35.4MG
ORAL; TABLET, SUSTAINED ACTION		3		06/10/83	UNK	4.0MG - 5.04MG
TOPICAL; LOTION		1				
VAGINAL; TABLET		1				
GUM BASE, CHEWING						
BUCCAL; GUM, CHEWING		2		06/08/92	UNK	
ORAL; TABLET		1				
GUM ROSIN	008050100					
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		4		11/14/94	UNK	9.0MG
GUM, NATURAL						
ORAL; TABLET, SUSTAINED ACTION		1				
HERBACOL	006365839					
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		1				
HEXYLENE GLYCOL	000107415					
TOPICAL; EMULSION, CREAM		1				
TOPICAL; OINTMENT		2		04/30/87	UNK	12.0%
HIGH FRUCTOSE CORN SYRUP						
ORAL; SUSPENSION, SUSTAINED ACTION		1				
HISTIDINE	000071001					
INTRAVENOUS; SUSPENSION, INJECTION		1				
HYDROCARBON GEL, PLASTICIZED	008049658					
DENTAL; PASTE		1				
OPHTHALMIC; OINTMENT		1				
TOPICAL; OINTMENT		4		10/29/82	UNK	
HYDROCHLORIC ACID	007647010					
CAUDAL BLOCK; INJECTION		15		10/13/87	600	1.414% - 2.827%
DENTAL; SOLUTION		1				
EPIDURAL; INJECTION		31		10/30/92	UNK	
IM - IV - SC; INJECTION		50		12/29/93	600	
IM - IV; INJECTION		113		12/27/94	600	0.00022%
IM - IV; POWDER, FOR INJECTION SOLUTION		11		10/30/95	600	
IM - IV; SOLUTION, INJECTION		3		03/05/90	600	
IM - SC; INJECTION		10		01/23/85	510	
IM - SC; POWDER, FOR INJECTION SOLUTION		1				
INHALATION; AEROSOL, METERED		3		09/03/92	510	0.7% - 1.72%
INHALATION; SOLUTION		23		07/28/95	600	
INTERSTITIAL; INJECTION		1				
INTRA-ARTICULAR; INJECTION		12		11/05/81	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
HYDROCHLORIC ACID	007647010					
INTRABURSAL; INJECTION		2		09/30/64	UNK	
INTRACARDIAC; INJECTION		3		06/01/88	600	
INTRACAVITARY; INJECTION		1				
INTRACAVITARY; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
INTRADERMAL; INJECTION		5		10/16/87	UNK	
INTRALESIONAL; INJECTION		8		10/16/87	UNK	
INTRAMUSCULAR; INJECTION		51		10/29/92	510	
INTRAMUSCULAR; SOLUTION, INJECTION		2		04/17/95	UNK	
INTRAOCULAR; SOLUTION		2		04/28/95	600	
INTRAPERITONEAL; INJECTION		1				
INTRAPERITONEAL; SOLUTION		5		08/19/92	160	
INTRAPLEURAL; INJECTION		1				
INTRASYNOVIAL; INJECTION		5		11/05/81	600	
INTRATHECAL; INJECTION		13		10/30/92	UNK	
INTRATHECAL; POWDER, FOR INJECTION SOLUTION		2		12/21/87	150	
INTRATHECAL; SOLUTION		1				
INTRATRACHEAL; POWDER, FOR RECONSTITUTION		1				
INTRATRACHEAL; SUSPENSION		1				
INTRATUMOR; INJECTION		1				
INTRATUMOR; POWDER, FOR INJECTION SOLUTION		1				
INTRAVASCULAR; INJECTION		4		01/22/92	160	
INTRAVASCULAR; SOLUTION		1				
INTRAVENOUS; INJECTION		165		08/18/95	110	1.414% - 10.0%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		14		04/19/95	600	
INTRAVENOUS; SOLUTION		7		12/26/85	160	0.9%
INTRAVENOUS; SUSPENSION, INJECTION		1				
IONTOPHORESIS; SOLUTION		1				
IRRIGATION; SOLUTION		7		11/27/91	600	
IV - SC; INJECTION		23		10/10/95	600	
IV - SC; POWDER, FOR INJECTION SOLUTION		2		08/31/90	600	
IV(INFUSION); INJECTION		176		08/18/95	110	1.414% - 2.827%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		7		12/21/90	600	
IV(INFUSION); SOLUTION		1				
NASAL; SOLUTION		3		12/26/90	510	
NASAL; SPRAY		1				
NASAL; SPRAY, METERED		11		10/20/95	UNK	
NERVE BLOCK; INJECTION		58		06/23/95	600	1.36% - 2.827%
OPHTHALMIC; GEL		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		61		12/29/95	600	
OPHTHALMIC; SUSPENSION		14		09/13/95	600	
ORAL; CONCENTRATE		6		06/30/92	600	0.217% - 0.47%
ORAL; SOLUTION		24		10/24/95	160	0.9%
ORAL; SUSPENSION		6		03/18/87	600	10.0%
ORAL; SYRUP		6		10/28/94	600	
OTIC; SOLUTION		6		12/29/95	600	0.56%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
HYDROCHLORIC ACID	007647010					
OTIC; SUSPENSION		1				
PERFUSION/CARDIAC; SOLUTION		1				
PERIDURAL; INJECTION		1				
RECTAL; SOLUTION		2		10/24/95	160	
RETROBULBAR; INJECTION		1				
SOFT TISSUE; INJECTION		6		06/19/80	600	
SPINAL; INJECTION		5		12/11/87	600	
SUBCONJUNCTIVAL; INJECTION		1				
SUBCUTANEOUS; INJECTION		19		03/31/94	510	
SUBCUTANEOUS; SOLUTION, INJECTION		1				
TOPICAL; EMULSION		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		2		10/08/85	600	
TOPICAL; GEL		2		02/07/89	600	
TOPICAL; GEL, JELLY		3		04/29/93	600	
TOPICAL; OINTMENT		1				
TOPICAL; SHAMPOO		2		08/31/90	UNK	
TOPICAL; SOLUTION		9		07/31/84	600	
URETERAL; SOLUTION		1				
HYDROCHLORIC ACID, DILUTED						
INTRA-ARTERIAL; INJECTION		1				
INTRAVASCULAR; INJECTION		1				
INTRAVENOUS; INJECTION		1				
IV(INFUSION); SOLUTION, INJECTION		1				
ORAL; CONCENTRATE		1				
TOPICAL; SOLUTION		1				
HYDROGEN PEROXIDE	007722841					
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		2		01/07/87	520	
HYDROXYETHYL CELLULOSE	009004620					
OPHTHALMIC; SOLUTION		3		06/29/76	600	0.25% - 0.35%
OPHTHALMIC; SUSPENSION		4		07/21/89	UNK	0.05% - 0.35%
ORAL; SYRUP		4		07/22/92	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, SUSTAINED ACTION		3		11/08/93	UNK	10.0MG - 20.0MG
OTIC; SOLUTION		1				
TOPICAL; SOLUTION		2		07/22/86	520	0.5% - 0.75%
TOPICAL; SPONGE		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
HYDROXYMETHYL CELLULOSE						
ORAL; TABLET		1				
TOPICAL; SOLUTION		1				
HYDROXYPROPYL CELLULOSE	009004642					
ORAL; CAPSULE		4		01/25/95	600	10.0MG - 36.0MG
ORAL; CAPSULE, COATED PELLETS		2		10/30/85	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		10/05/95	180	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
HYDROXYPROPYL CELLULOSE	009004642					
ORAL; CAPSULE, SUSTAINED ACTION		9		01/04/95	600	0.2MG - 5.49MG
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET		168		11/09/95	600	0.05MG - 46.0MG
ORAL; TABLET, COATED		14		06/20/88	600	0.5MG - 1.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		06/19/95	520	15.0MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		47		12/11/95	530	0.58GM
ORAL; TABLET, SUSTAINED ACTION		15		03/30/95	110	2.0MG - 37.5MG
SUBLINGUAL; TABLET		1				
TOPICAL; GEL		5		01/29/95	600	2.1% - 3.0%
TOPICAL; LOTION		2		03/30/89	UNK	0.15% - 0.54%
TOPICAL; LOTION, AUGMENTED		1				
TOPICAL; SOLUTION		2		07/03/85	600	0.05%
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
HYDROXYPROPYL METHYLCELLULOSE	009004653					
OPHTHALMIC; SOLUTION		6		10/31/95	600	0.1% - 0.5%
OPHTHALMIC; SUSPENSION		13		09/13/95	600	0.002% - 0.6%
ORAL; CAPSULE		15		09/11/92	530	1.58MG - 150.0MG
ORAL; CAPSULE, COATED PELLETS		2		10/30/85	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		10/05/95	180	
ORAL; CAPSULE, SUSTAINED ACTION		6		05/29/92	110	1.4MG - 119.7MG
ORAL; CONCENTRATE		1				
ORAL; GRANULE, ENTERIC COATED		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		3		04/29/93	600	0.45%
ORAL; TABLET		317		10/04/95	510	0.4MG - 48.0MG
ORAL; TABLET, COATED		15		05/24/88	600	1.54MG - 6.04MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		11/30/95	600	3.0MG - 18.7MG
ORAL; TABLET, FILM COATED		97		12/27/95	150	0.764MG - 170.0MG
ORAL; TABLET, SUSTAINED ACTION		35		03/30/95	110	2.81MG - 240.0MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
TOPICAL; GEL, JELLY		2		04/29/93	600	3.5%
TOPICAL; SOLUTION		1				
VAGINAL; TABLET		1				
HYDROXYPROPYL METHYLCELLULOSE PHTHALATE						
ORAL; CAPSULE, COATED PELLETS		3		06/30/92	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		10/05/95	180	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		03/29/82	600	29.2MG - 44.57MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
HYDROXYPROPYL METHYLCELLULOSE 2208	009004653					
ORAL; TABLET, SUSTAINED ACTION		6		04/28/95	600	30.0MG - 250.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
HYDROXYPROPYL METHYLCELLULOSE 2906	009004653					
OPHTHALMIC; SOLUTION		2		07/29/94	600	0.5%
ORAL; GRANULE, ENTERIC COATED		1				
ORAL; SYRUP		1				
HYDROXYPROPYL METHYLCELLULOSE 2910	009004653					
OPHTHALMIC; SOLUTION		3		09/23/93	UNK	0.5%
OPHTHALMIC; SUSPENSION		3		05/09/89	600	0.5%
ORAL; CAPLET		1				
ORAL; CAPSULE		9		04/08/94	530	1.0MG - 10.8MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		2		09/11/95	110	1.77MG - 10.88MG
ORAL; SUSPENSION		1				
ORAL; SYRUP		3		09/25/95	600	
ORAL; TABLET		68		10/06/95	UNK	0.2MG - 33.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		9		12/30/92	110	0.7MG - 9.8MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		12		10/24/94	600	3.8MG - 60.0MG
ORAL; TABLET, SUSTAINED ACTION		11		11/18/94	UNK	5.7MG - 144.0MG
ORAL-28; TABLET		1				
IMIDAZOLIDINYL UREA						
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
IMIDUREA	039236469					
TOPICAL; EMULSION, CREAM		2		06/17/94	UNK	0.2% - 0.3%
TOPICAL; SHAMPOO		1				
INK BLACK						
ORAL; CAPSULE		12		12/22/92	600	65.0MG - 102.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		4		11/30/95	600	
INK BLACK A-10450						
ORAL; TABLET		1				
ORAL; TABLET, FILM COATED		1				
INK BLACK A-10509						
ORAL; CAPSULE		3		06/11/87	600	
INK BLACK A-1057						
ORAL; TABLET, SUSTAINED ACTION		1		-- --		
INK BLACK IMPRINTING FGE-1386						
ORAL; CAPSULE		1				
INK BLUE BLACK A-9371						
ORAL; CAPSULE		1				
INK EDIBLE						
ORAL; CAPSULE		11		03/02/92	600	
ORAL; CAPSULE, SOFT GELATIN		2		12/30/86	150	
ORAL; CAPSULE, SUSTAINED ACTION		5		02/28/92	600	
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		4		05/14/85	UNK	
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
INK EDIBLE BLACK						
ORAL; CAPSULE		7		03/25/94	UNK	
ORAL; TABLET		4		02/16/88	600	
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		2		01/09/87	600	1.0MG
ORAL-28; TABLET		1				
INK EDIBLE GRAY						
ORAL; CAPSULE		1				
INK EDIBLE RED						
ORAL; CAPSULE		1				
INK EDIBLE RED A-8032						
ORAL; TABLET, SUSTAINED ACTION		2		07/13/87	UNK	
INK EDIBLE WHITE						
ORAL; CAPSULE		3		03/25/94	UNK	
ORAL; TABLET, FILM COATED		1				
INK FINE BLACK 2202C						
ORAL; TABLET		2		02/22/85	600	0.0007ML - 0.0011ML
INK FINE BLACK 2212						
ORAL; TABLET		3		09/25/84	600	
INK GREEN A-10454						
ORAL; TABLET, SUSTAINED ACTION		1				
INK LIGHT REDWOOD						
ORAL; CAPSULE		1				
INK PINK IMPRINTING SB-1003						
ORAL; CAPSULE		1				
INK RED A-8032						
ORAL; CAPSULE		1				
ORAL; TABLET, COATED		1				
INK RED S-1-9005						
ORAL; CAPSULE		1				
INK WHITE						
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; TABLET		1				
INK WHITE A-8154						
ORAL; CAPSULE		1				
ORAL; TABLET, FILM COATED		1				
INK WHITE 21-K						
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		1				
INVERT SUGAR	008013170					
ORAL; CAPSULE		1				
ORAL; SYRUP		4		06/07/85	600	
IODINE	007553562					
INTRA-ARTERIAL; INJECTION		2		08/29/74	160	28.2% - 37.0%
INTRA-ARTICULAR; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRADISCAL; INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
IODINE	007553562					
INTRAVASCULAR; INJECTION		3		08/29/78	160	20.2% - 40.0%
INTRAVENOUS; INJECTION		3		08/29/74	160	14.1% - 37.0%
PERIARTICULAR; INJECTION		1				
IOFETAMINE HYDROCHLORIDE						
INTRAVENOUS; INJECTION		1				
IRISH MOSS EXTRACT						
TOPICAL; LOTION		2		08/16/84	600	0.3%
IRON OXIDE						
ORAL; CAPSULE		18		03/03/95	110	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		5		05/24/82	600	
ORAL; DROPS		1				
ORAL; TABLET		30		07/12/95	110	0.04MG - 0.8MG
ORAL; TABLET, COATED		9		05/19/92	110	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		4		10/14/94	UNK	
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		11		06/01/95	120	0.25MG
ORAL; TABLET, SUSTAINED ACTION		6		12/24/92	120	0.3MG - 0.576MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
IRON OXIDE, BROWN						
ORAL; CAPSULE		2		04/30/92	600	
ORAL; TABLET		3		09/24/86	600	
ORAL; TABLET, FILM COATED		1				
IRON OXIDE, RED-BROWN						
ORAL; TABLET, SUSTAINED ACTION		1				
IRON OXIDE, YELLOW						
ORAL; CAPSULE		19		10/18/95	600	0.0608MG - 0.8MG
ORAL; CAPSULE, HARD GELATIN		3		05/03/95	530	
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		2		11/30/93	600	
ORAL; TABLET		26		11/30/95	510	0.02MG - 2.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		11		12/19/95	180	0.0015MG - 0.092MG
ORAL; TABLET, SUSTAINED ACTION		2		03/30/95	110	0.04MG
ORAL-21; TABLET		1				
ISOBUTANE	000075285					
TOPICAL; AEROSOL SPRAY		3		05/24/77	UNK	77.6%
ISOCETETH-20						
TOPICAL; SOLUTION		1				
ISOOCTYLACRYLATE						
TOPICAL; TAPE		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ISOPROPYL ALCOHOL	000067630					
ORAL; TABLET		3		09/10/87	520	
TOPICAL; AEROSOL SPRAY		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; LOTION		21		09/30/94	600	2.7% - 78.0%
TOPICAL; LOTION, AUGMENTED		1				
TOPICAL; OIL		1				
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		19		12/15/95	600	4.0% - 51.5%
TOPICAL; SPONGE		5		02/28/91	600	4.2MG
ISOPROPYL ISOSTEARATE						
TOPICAL; EMULSION, CREAM		1				
ISOPROPYL MYRISTATE	000110270					
OTIC; SUSPENSION		1				
TOPICAL; AEROSOL		1				
TOPICAL; AEROSOL SPRAY		2		06/15/73	600	21.96%
TOPICAL; EMULSION, CREAM		33		06/17/94	UNK	1.0% - 10.0%
TOPICAL; GEL		3		06/06/84	600	10.0%
TOPICAL; LOTION		3		12/07/92	UNK	2.0%
TOPICAL; OIL		1				
TOPICAL; OINTMENT		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
VAGINAL; EMULSION, CREAM		2		02/21/91	520	
ISOPROPYL PALMITATE	000142916					
TOPICAL; AEROSOL SPRAY		1				
TOPICAL; EMULSION, CREAM		48		09/20/95	UNK	0.325% - 9.9375%
TOPICAL; LOTION		5		06/13/88	UNK	0.6% - 5.0%
TOPICAL; OINTMENT		2		07/24/78	600	
ISOPROPYL STEARATE						
TOPICAL; EMULSION, CREAM		1				
ISOSTEARIC ACID						
TOPICAL; OINTMENT		1				
ISOSTEARYL ALCOHOL						
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		1				
ISOTONIC SODIUM CHLORIDE SOLUTION	008028771					
INTRAVENOUS; SOLUTION		1				
ORAL; SOLUTION		1				
JELENE	008049669					
TOPICAL; OINTMENT		1				
KAOLIN	001332587					
ORAL; CAPSULE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SYRUP		1				
ORAL; TABLET		5		05/31/94	530	7.95MG - 30.4MG
ORAL; TABLET, COATED		2		09/29/76	600	2.87MG - 8.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
KAOLIN	001332587					
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
KATHON CG						
TOPICAL; EMULSION, CREAM		1				
LAC RESIN						
ORAL; CAPSULE, SUSTAINED ACTION		1				
LACTATE	000050215					
TOPICAL; LOTION		1				
LACTIC ACID	008012213					
IM - IV - SC; INJECTION		4		01/06/76	600	0.012%
IM - IV; INJECTION		16		08/31/95	600	0.25% - 1.1578%
INTRACARDIAC; INJECTION		3		01/06/76	600	0.012%
INTRAMUSCULAR; INJECTION		9		02/25/93	600	
INTRAVENOUS; INJECTION		1				
IV(INFUSION); INJECTION		8		08/09/94	110	0.012% - 1.86%
IV(INFUSION); SOLUTION, INJECTION		1				
ORAL; CONCENTRATE		8		09/28/93	600	0.00022% - 0.3%
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
ORAL; TABLET		1				
TOPICAL; EMULSION, CREAM		25		09/20/95	UNK	0.015% - 2.0%
TOPICAL; LOTION		4		12/30/88	UNK	0.51% - 6.6%
TOPICAL; OINTMENT		3		10/01/86	600	
TOPICAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		2		01/27/87	520	0.81%
VAGINAL; SUPPOSITORY		1				
VAGINAL; TABLET		1				
LACTIC ACID, DL-	000598823					
IM - IV; INJECTION		1				
IMPLANTATION; PELLET, IMPLANT		2		12/18/95	150	
SUBCUTANEOUS; PELLET, IMPLANT		2		12/18/95	150	
VAGINAL; SUPPOSITORY		1				
LACTOBIONIC ACID	000096822					
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
LACTOSE	000063423					
BUCCAL; TABLET		1				
BUCCAL/SUBLINGUAL; TABLET		1				
IM - IV - SC; INJECTION		1				
IM - IV - SC; POWDER, FOR INJECTION SOLUTION		1				
IM - IV; INJECTION		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		2		11/20/64	UNK	
INHALATION; CAPSULE		1				
INHALATION; CAPSULE, HARD GELATIN		1				
INTRACAVITARY; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
INTRAMUSCULAR; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
LACTOSE	000063423					
INTRAVENOUS; INJECTION		5		04/13/89	600	
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		2		03/31/89	600	0.1% - 2.5%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); SOLUTION, INJECTION		1				
ORAL; CAPSULE		288		12/29/95	600	14.805%
ORAL; CAPSULE, COATED PELLETS		2		10/30/85	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		10/05/95	180	
ORAL; CAPSULE, HARD GELATIN		4		12/06/95	530	14.25MG - 100.0MG
ORAL; CAPSULE, SUSTAINED ACTION		6		02/21/92	110	21.6MG - 120.0MG
ORAL; CONCENTRATE		1				
ORAL; GRANULE		2		05/20/88	600	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		6		08/25/92	600	8.0%
ORAL; TABLET		1,160		12/29/95	600	0.031GM - 1.02GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		5		09/11/95	600	30.0MG - 117.7MG
ORAL; TABLET, COATED		63		12/30/92	110	2.8MG - 346.5MG
ORAL; TABLET, CONTROLLED RELEASE		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		5		11/30/95	600	40.0MG - 209.0MG
ORAL; TABLET, FILM COATED		68		12/27/95	150	1.96MG - 590.0MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	122.99MG - 153.2MG
ORAL; TABLET, SUSTAINED ACTION		24		11/23/94	UNK	10.0MG - 400.0MG
ORAL-21; TABLET		31		12/13/93	600	7.8MG - 89.007MG
ORAL-28; TABLET		35		11/17/95	510	21.0MG - 179.2MG
RECTAL; SOLUTION		2		08/25/92	600	8.0%
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		2		09/30/82	510	
SUBLINGUAL; TABLET		12		07/29/88	110	9.967MG - 327.5MG
TRANSDERMAL; OINTMENT		1				
VAGINAL; EMULSION, CREAM		1				
VAGINAL; SUPPOSITORY		2		01/27/87	520	
VAGINAL; TABLET		8		12/26/91	520	395.0MG - 696.0MG
LACTOSE MONOHYDRATE	010039266					
IM - IV; INJECTION		1				
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
ORAL; CAPSULE		7		09/29/95	600	18.0MG - 360.0MG
ORAL; TABLET		35		11/22/95	600	12.5MG - 370.0MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		2		05/24/95	600	92.04MG - 260.0MG
LACTOSE MONOHYDRATE, ALPHA	005989811					
ORAL; CAPSULE		1				
LACTOSE, ANHYDROUS	000063423					
ORAL; CAPSULE		1				
ORAL; TABLET		15		11/30/95	600	7.5MG - 415.8MG
ORAL; TABLET, COATED		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
LACTOSE, ANHYDROUS	000063423					
ORAL; TABLET, FILM COATED		4		12/11/95	530	26.85MG - 180.6MG
ORAL; TABLET, SUSTAINED ACTION		1				
LACTOSE, HYDROUS						
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
ORAL; CAPSULE		45		03/30/95	600	20.0MG - 498.65MG
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; TABLET		138		11/30/95	510	1.0MG - 535.6MG
ORAL; TABLET, COATED		2		02/25/92	600	32.0MG - 41.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		9		12/08/95	UNK	12.75MG - 556.0MG
ORAL; TABLET, SUSTAINED ACTION		6		01/04/95	600	58.0MG - 156.8MG
SUBLINGUAL; TABLET		3		02/19/88	600	1.7MG - 11.6MG
VAGINAL; EMULSION, CREAM		1				
LANOLIN	008020846					
TOPICAL; EMULSION, CREAM		2		04/10/79	600	2.0%
TOPICAL; LOTION		4		12/16/81	600	0.0014% - 1.0%
TRANSDERMAL; OINTMENT		1				
VAGINAL; EMULSION, CREAM		3		06/09/86	600	1.0% - 2.0%
LANOLIN ALCOHOLS	008013341					
OPHTHALMIC; OINTMENT		3		12/01/89	UNK	10.0%
TOPICAL; EMULSION, CREAM		16		10/29/93	UNK	2.0% - 3.0%
TOPICAL; GEL		1				
TOPICAL; OINTMENT		3		03/23/95	600	
LANOLIN ALCOHOLS, ACETYLATED	008028986					
TOPICAL; SPONGE		1				
LANOLIN CHOLESTEROLS						
TOPICAL; EMULSION		1				
LANOLIN NONIONIC DERIVATIVES						
OPHTHALMIC; OINTMENT		1				
LANOLIN OIL	008038435					
OPHTHALMIC; OINTMENT		2		07/25/94	600	2.0% - 3.0%
LANOLIN, ANHYDROUS	008006540					
OPHTHALMIC; OINTMENT		5		06/17/81	UNK	3.0% - 10.0%
TOPICAL; EMULSION, CREAM		1				
VAGINAL; EMULSION, CREAM		1				
LANOLIN, HYDROGENATED	008031445					
TOPICAL; OINTMENT		3		12/19/85	600	0.5% - 10.0%
LAURAMINE OXIDE						
TOPICAL; SOLUTION		1				
LAURDIMONIUM HYDROLYZED ANIMAL COLLAGEN						
TOPICAL; SHAMPOO		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
LAURETH SULFATE						
TOPICAL; SPONGE		1				
LAURETH 23						
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
LAURETH 4	009002920					
TOPICAL; LOTION		1				
TOPICAL; SOLUTION		1				
LAURIC DIETHANOLAMIDE	000120401					
TOPICAL; LOTION		1				
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		1				
TOPICAL; SUSPENSION, SHAMPOO		2		01/10/91	600	
LAURIC MYRISTIC DIETHANOLAMIDE						
TOPICAL; EMULSION		1				
LAURYL SULFATE						
ORAL; CAPSULE		1				
LECITHIN	008002435					
INHALATION; AEROSOL, METERED		1				
INTRAMUSCULAR; INJECTION		4		05/24/79	520	0.3% - 0.6%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
ORAL; CAPSULE		13		08/29/95	600	0.5MG - 15.0MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET		2		05/31/94	530	
ORAL; TABLET, FILM COATED		1				
RECTAL; SUPPOSITORY		2		05/12/87	600	1.2MG - 6.5MG
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				G
VAGINAL; EMULSION, CREAM		5		06/09/86	600	0.3% - 1.0%
LECITHIN, HYDROGENATED SOY						
INTRAVENOUS; SUSPENSION, INJECTION		1				
LECITHIN, SOY BEAN	008030760					
INHALATION; AEROSOL, METERED		1				
ORAL; CAPSULE, SOFT GELATIN		2		12/15/86	530	4.0MG - 20.0MG
VAGINAL; EMULSION, CREAM		1				
LEMON OIL	008008568					
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		1				
TOPICAL; GEL		1				
LEVOMENTHOL	002216515					
ORAL; SYRUP		1				
LIDOFENIN	059160291					
INTRAVENOUS; INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
LIME OIL	008008262					
ORAL; SOLUTION, ELIXIR		1				
LIMONENE, DL-	000138863					
TOPICAL; LOTION		1				
LINEAR ALCOHOL ETHYLENE OXIDE ADDUCT						
TOPICAL; LOTION		1				
LUBRITAB						
ORAL; TABLET		2		12/22/82	600	3.45MG - 10.0MG
LYSINE	000056871					
IM - IV; INJECTION		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
MAGNESIUM ALUMINUM SILICATE	001327431					
ORAL; DROPS		1				
ORAL; GRANULE		3		12/18/80	600	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		26		03/30/94	600	0.15% - 2.0%
ORAL; SYRUP		1				
ORAL; TABLET		2		11/03/70	UNK	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		07/05/88	520	8.0MG
RECTAL; SUSPENSION		2		11/17/86	600	
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		3		04/24/85	UNK	1.5%
TOPICAL; SHAMPOO		2		08/27/90	UNK	0.5%
VAGINAL; OINTMENT		1				
MAGNESIUM CARBONATE	000546930					
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; TABLET		8		02/21/95	600	5.37MG - 250.0MG
ORAL; TABLET, COATED		2		09/30/59	120	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
MAGNESIUM CHLORIDE	007791186					
INTRAMUSCULAR; INJECTION		1				
INTRAOCULAR; SOLUTION		2		04/28/95	600	0.03%
INTRAPERITONEAL; SOLUTION		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		2		09/22/93	UNK	
MAGNESIUM HYDROXIDE	001309428					
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		10/09/91	530	12.44MG - 450.0MG
MAGNESIUM NITRATE	010377603					
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
MAGNESIUM OXIDE	001309484					
ORAL; CAPSULE		2		04/25/88	600	10.0MG
ORAL; TABLET		15		02/09/94	510	5.0MG - 20.0MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
MAGNESIUM SILICATE	001343880					
ORAL; CAPSULE		1				
ORAL; TABLET		1				
ORAL; TABLET, FILM COATED		1				
MAGNESIUM STEARATE	000557040					
BUCCAL; TABLET		1				
BUCCAL/SUBLINGUAL; TABLET		1				
ORAL; CAPSULE		464		12/29/95	600	0.003GM
ORAL; CAPSULE, COATED PELLETS		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		09/12/89	120	
ORAL; CAPSULE, HARD GELATIN		6		12/06/95	530	1.05MG - 7.0MG
ORAL; CAPSULE, SUSTAINED ACTION		16		09/11/95	110	0.25MG - 100.0MG
ORAL; CONCENTRATE		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		1		[illegible]	[illegible]	
ORAL; TABLET		1,818		12/29/95	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		17		09/11/95	600	1.5MG - 50.0MG
ORAL; TABLET, COATED		84		04/23/95	600	0.18MG - 40.0MG
ORAL; TABLET, CONTROLLED RELEASE		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		14		11/30/95	600	0.8MG - 12.0MG
ORAL; TABLET, DISPERSIBLE		3		04/28/95	600	
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		116		12/27/95	150	0.48MG - 10.9MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	0.7MG - 1.2MG
ORAL; TABLET, SUSTAINED ACTION		84		04/28/95	600	0.3MG - 70.0MG
ORAL; TABLET, UNCOATED, TROCHE		1				
ORAL-21; TABLET		30		12/13/93	600	0.25MG - 1.0MG
ORAL-28; TABLET		37		11/17/95	510	0.05MG - 75.0MG
SUBLINGUAL; TABLET		15		07/29/88	110	0.08MG - 1.5MG
TOPICAL; EMULSION, CREAM		3		10/29/93	UNK	
VAGINAL; TABLET		5		12/26/91	520	4.0MG - 17.0MG
MAGNESIUM SULFATE	010034998					
ORAL; TABLET		7		05/15/86	600	0.5MG - 2.9MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		1				
MAGNESIUM SULFATE, ANHYDROUS	007487889					
ORAL; CAPSULE		1				
MAGNESIUM TRISILICATE	001343904					
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
MALEIC ACID	000110167					
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		1				
ORAL; CAPSULE		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
MALEIC ACID	000110167					
ORAL; SYRUP		1				
ORAL; TABLET		2		06/28/78	510	2.0MG
MALIC ACID	006915157					
ORAL; SOLUTION		1				
MALIC ACID, DL-						
ORAL; SOLUTION		2		04/27/88	510	
ORAL; TABLET		1				
MALTODEXTRIN						
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; TABLET		4		11/05/92	600	
MALTOL	000118718					
ORAL; CONCENTRATE		1				
ORAL; SOLUTION		4		10/31/93	600	
ORAL; SYRUP		1				
MALTOLDEXTRIN	009050366					
ORAL; GRANULE, FOR RECONSTITUTION		1				
MALTOSE	000069794					
ORAL; SOLUTION		1				
MANNITOL	000069658					
IM - IV; INJECTION		2		04/22/70	510	
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
IM - SC; POWDER, FOR INJECTION SOLUTION		1				
INTRAMUSCULAR; INJECTION		3		01/21/94	510	
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		2		10/17/85	510	
INTRAVENOUS; INJECTION		12		12/24/91	510	10.0%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		12		11/29/95	600	100.0MG
IV(INFUSION); INJECTION		5		02/16/89	600	2.0% - 10.0%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		11		09/20/95	180	37.0MG - 1500.0MG
OPHTHALMIC; POWDER, FOR RECONSTITUTION		3		09/22/93	UNK	
OPHTHALMIC; SOLUTION		4		11/04/93	UNK	4.05% - 4.6%
OPHTHALMIC; SUSPENSION		2		12/30/94	UNK	4.5%
ORAL; CAPSULE		4		03/30/89	600	65.6MG - 283.1MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; POWDER, FOR RECONSTITUTION		5		08/06/84	520	4.0% - 4.188%
ORAL; TABLET		30		12/05/94	120	5.0MG - 302.8MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		9		09/11/95	600	75.0MG - 600.0MG
ORAL; TABLET, COATED		2		02/11/82	120	25.8MG - 177.7MG
ORAL; TABLET, SUSTAINED ACTION		4		11/18/94	UNK	110.0MG - 392.2MG
SUBCUTANEOUS; INJECTION		1				
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		4		05/25/95	510	
SUBLINGUAL; TABLET		6		02/26/88	600	10.0MG - 143.4475MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
MANNOSE, D-	000530267					
ORAL; TABLET		1				
MEBROFENIN	078266065					
INTRAVENOUS; INJECTION		1				
MEDICAL ANTIFOAM EMULSION C						
ORAL; SUSPENSION		6		03/18/87	600	0.08%
ORAL; TABLET		1				
MEDICAL ANTIFORM A-F EMULSION						
TOPICAL; EMULSION, CREAM		1				
MEDRONATE DISODIUM	025681894					
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
MEDRONIC ACID	001984152					
INTRAVENOUS; INJECTION		3		02/17/81	160	
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
MEGLUMINE	006284408					
INTRA-ARTERIAL; INJECTION		3		07/26/85	160	
INTRA-ARTICULAR; INJECTION		1				
INTRACARDIAC; INJECTION		2		07/26/85	160	
INTRADISCAL; INJECTION		1				
INTRAUTERINE; INJECTION		1				
INTRAUTERINE; SOLUTION		1				
INTRAVENOUS; INJECTION		4		06/02/88	160	0.039%
IV(INFUSION); INJECTION		1				
PERIARTICULAR; INJECTION		1				
URETERAL; SOLUTION		1				
MENTHOL	000089781					
DENTAL; SOLUTION		1				
INHALATION; AEROSOL, METERED		2		12/28/84	UNK	0.02%
ORAL; AEROSOL		1				
ORAL; CAPSULE		1				
ORAL; CONCENTRATE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		2		12/22/94	600	
ORAL; SUSPENSION		3		01/26/84	600	0.003%
ORAL; SYRUP		11		02/21/85	600	0.01%
ORAL; TABLET		1				
TOPICAL; LOTION		3		04/29/82	600	0.05001%
METAPHOSPHORIC ACID	013478983					
IV(INFUSION); INJECTION		1				
METHACRYLIC ACID COPOLYMER						
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		05/10/95	180	
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; GRANULE, ENTERIC COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		4		06/19/95	520	19.5MG - 48.65MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
METHANESULFONIC ACID	000075752					
IM - IV; INJECTION		1				
METHIONINE	000063683					
ORAL; TABLET		1				
METHYL ACRYLATE - METHYL METHACRYLATE						
ORAL; TABLET, SUSTAINED ACTION		1				
METHYL BORONIC ACID						
INTRAVENOUS; INJECTION		1				
METHYL GLUCETH-120 DIOLEATE						
TOPICAL; SHAMPOO		1				
METHYL HYDROXYETHYL CELLULOSE						
ORAL; TABLET		2		06/02/87	600	12.0MG - 24.0MG
METHYL LAURATE						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
METHYL SALICYLATE	000119368					
TOPICAL; GEL		1				
METHYL STEARATE	000112618					
TOPICAL; EMULSION, CREAM		1				
VAGINAL; EMULSION, CREAM		1				
METHYLATED SPIRITS						
ORAL; CAPSULE		1				
ORAL; TABLET, COATED		1				
METHYLCELLULOSE	009004675					
INTRAMUSCULAR; INJECTION		2		07/07/83	600	0.03%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		3		08/24/72	600	0.1641% - 0.5%
ORAL; CAPSULE		4		12/04/85	600	1.0MG - 6.04MG
ORAL; POWDER, FOR RECONSTITUTION		4		04/18/91	600	0.08% - 0.1%
ORAL; SUSPENSION		3		12/16/93	180	
ORAL; TABLET		74		10/30/92	600	0.756MG - 55.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		7		01/26/84	510	0.29MG - 30.0MG
ORAL; TABLET, FILM COATED		15		05/09/88	600	2.3MG - 21.0MG
ORAL; TABLET, SUSTAINED ACTION		2		08/21/92	UNK	3.0MG
ORAL-28; TABLET		2		11/17/95	510	12.0MG - 15.0MG
SUBLINGUAL; TABLET		2		07/07/80	600	
TOPICAL; EMULSION		1				
TOPICAL; LOTION		2		04/24/85	UNK	0.2% - 1.5%
VAGINAL; EMULSION, CREAM		1				
METHYLCELLULOSE 400						
OPHTHALMIC; SOLUTION		1				
METHYLCHLOROISOTHIAZOLINONE	026172554					
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
METHYLENE BLUE	000061734					
INTRAVENOUS; INJECTION		1				
METHYLISOTHIAZOLINONE	002682204					
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
METHYLPARABEN	000099763					
CAUDAL BLOCK; INJECTION		4		03/03/87	600	0.1%
EPIDURAL; INJECTION		9		03/03/87	600	0.1%
IM - IV - SC; INJECTION		26		04/11/89	600	0.15% - 0.2%
IM - IV; INJECTION		37		11/27/91	600	0.065% - 0.2%
IM - SC; INJECTION		2		06/10/87	510	0.1% - 0.15%
INHALATION; SOLUTION		5		06/30/81	600	0.0249934% - 0.07%
INTRA-ARTICULAR; INJECTION		3		06/19/80	600	0.15%
INTRABURSAL; INJECTION		1				
INTRADERMAL; INJECTION		1				
INTRALESIONAL; INJECTION		3		06/19/80	600	0.15%
INTRAMUSCULAR; INJECTION		31		02/25/93	600	0.09% - 0.18%
INTRASYNOVIAL; INJECTION		2		03/01/77	UNK	0.15%
INTRAVENOUS; INJECTION		23		12/20/91	UNK	0.05% - 0.18%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IRRIGATION; SOLUTION		1				
IV - SC; INJECTION		2		12/05/85	180	0.15% - 0.18%
IV(INFUSION); INJECTION		26		03/25/94	160	0.005% - 0.18%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
NASAL; SOLUTION		2		05/18/78	510	0.033%
NERVE BLOCK; INJECTION		32		06/21/88	600	0.05% - 0.101%
OPHTHALMIC; OINTMENT		8		08/31/95	600	0.05%
OPHTHALMIC; SOLUTION		9		10/18/88	600	0.015% - 0.05%
OPHTHALMIC; SUSPENSION		2		12/28/82	600	0.05%
ORAL; CAPSULE		79		12/20/95	520	0.128MG - 1.0MG
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		2		01/29/93	600	
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SOFT GELATIN		4		11/22/95	150	0.22MG - 0.323MG
ORAL; CAPSULE, SUSTAINED ACTION		8		04/25/95	UNK	
ORAL; CONCENTRATE		25		09/28/93	600	0.005% - 0.2%
ORAL; GRANULE		2		05/20/88	600	
ORAL; POWDER, FOR RECONSTITUTION		5		12/31/91	520	0.02% - 0.1%
ORAL; SOLUTION		46		11/17/95	530	0.015% - 0.4%
ORAL; SOLUTION, ELIXIR		14		10/27/92	600	0.05% - 0.1%
ORAL; SUSPENSION		47		09/15/95	180	0.05% - 0.2%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; SYRUP		69		07/11/95	600	0.05% - 0.18%
ORAL; TABLET		26		03/30/94	600	0.005MG - 0.186MG
ORAL; TABLET, COATED		14		02/25/92	600	0.01UGM
ORAL; TABLET, FILM COATED		3		12/28/87	520	0.06MG - 0.23MG
ORAL; TABLET, SUSTAINED ACTION		3		05/22/87	UNK	0.17MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
METHYLPARABEN	000099763					
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
OTIC; SUSPENSION		1				
PERIDURAL; INJECTION		1				
RECTAL; EMULSION, AEROSOL FOAM		1				
RECTAL; ENEMA		2		05/27/94	600	0.18%
RECTAL; SOLUTION		1				
RECTAL; SUSPENSION		2		11/17/86	600	
SOFT TISSUE; INJECTION		2		06/19/80	600	0.15%
SUBCUTANEOUS; INJECTION		9		12/19/91	UNK	0.1% - 0.15%
TOPICAL; EMULSION, AEROSOL FOAM		2		12/19/79	600	0.108%
TOPICAL; EMULSION, CREAM		65		10/31/94	600	0.018% - 0.3%
TOPICAL; GEL		2		11/22/88	UNK	0.08% - 0.3%
TOPICAL; GEL, JELLY		2		04/29/93	600	0.007%
TOPICAL; LOTION		15		12/07/92	UNK	0.08% - 0.3%
TOPICAL; OINTMENT		12		09/30/94	600	0.02% - 0.4%
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		4		04/03/85	600	0.1%
TOPICAL; SUSPENSION, SHAMPOO		1				
URETERAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		12		12/21/95	520	0.1% - 0.18%
VAGINAL; GEL		1				
VAGINAL; SUPPOSITORY		1				
METHYLPARABEN SODIUM	005026620					
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		2		11/17/86	600	
RECTAL; SUSPENSION		1				
MICROCRYSTALLINE WAX	008063089					
TOPICAL; GEL		1				
TOPICAL; OINTMENT		2		12/14/90	UNK	25.0%
MINERAL OIL	008012951					
DENTAL; PASTE		2		10/01/86	600	45.795% - 95.0%
OPHTHALMIC; OINTMENT		37		10/30/95	600	3.0% - 59.5%
OPHTHALMIC; SUSPENSION		2		05/11/88	600	
ORAL; CAPSULE		7		01/06/75	600	3.24MG - 5.0MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; DROPS		1				
ORAL; TABLET		13		09/26/89	120	0.0002ML - 85.0ML
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OTIC; OINTMENT		1				
OTIC; SUSPENSION		1				
TOPICAL; AEROSOL SPRAY		1				
TOPICAL; EMULSION, CREAM		64		06/13/95	600	3.0% - 40.0%
TOPICAL; LOTION		9		12/30/88	UNK	1.0% - 16.0%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT / ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
MINERAL OIL						
TOPICAL; OIL	008012951	1				
TOPICAL; OINTMENT		44		03/23/95	600	0.1% - 95.0%
TOPICAL; PASTE		1				
TOPICAL; SUPPOSITORY		2		04/26/93	520	
TRANSDERMAL; FILM, CONTROLLED RELEASE		3		10/28/94	510	2.31MG - 12.4MG
VAGINAL; EMULSION, CREAM		8		12/21/95	520	2.0% - 7.0%
VAGINAL; SUPPOSITORY		1				
MINERAL OIL, LIGHT						
OPHTHALMIC; OINTMENT		1				
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; PASTILLE		1				
ORAL; TABLET		9		04/28/89	520	0.625MG - 7.5MG
ORAL; TABLET, COATED		2		02/11/82	120	0.00015ML
ORAL; TABLET, FILM COATED		2		08/10/82	520	1.07MG - 2.494MG
ORAL; TABLET, SUSTAINED ACTION		1				
RECTAL; SUSPENSION		1				
TOPICAL; EMULSION, CREAM		8		09/28/92	600	3.0% - 20.0%
TOPICAL; LOTION		5		01/24/92	600	7.0% - 16.0%
TOPICAL; OINTMENT		15		03/23/95	600	4.4% - 23.0%
TRANSDERMAL; FILM, CONTROLLED RELEASE		2		09/10/86	510	19.0MG - 57.0MG
MONOGLYCERIDE CITRATE						
TOPICAL; EMULSION, CREAM		1				
MONOGLYCERIDES						
TOPICAL; LOTION		1				
MULTISTEROL EXTRACT						
TOPICAL; EMULSION, CREAM		2		04/10/79	600	
MYRISTIC ACID						
ORAL; CAPSULE, SUSTAINED ACTION	000544638	2		01/28/92	600	
MYRISTYL ALCOHOL						
ORAL; TABLET, SUSTAINED ACTION	000112721	5		01/04/95	600	0.58MG - 2.0MG
TOPICAL; EMULSION, CREAM		2		11/20/92	UNK	3.0%
TOPICAL; LOTION		1				
MYRISTYL-GAMMA-PICOLINIUM CHLORIDE						
INTRA-ARTICULAR; INJECTION	002748881	1				
INTRALESIONAL; INJECTION		3		03/26/79	600	0.019%
INTRAMUSCULAR; INJECTION		3		03/26/79	600	0.019%
INTRASYNOVIAL; INJECTION		4		03/26/79	600	0.019% - 0.169%
SOFT TISSUE; INJECTION		1				
		3		03/26/79	600	0.019%