INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PETROLATUM	008009038					
TOPICAL; OINTMENT, AUGMENTED		1				
TRANSDERMAL; OINTMENT		1				
VAGINAL; OINTMENT		1				
PETROLATUM, WHITE						
NASAL; OINTMENT		1				
OPHTHALMIC; OINTMENT		15		10/30/95	600	49.8% - 85.0%
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		37		04/01/94	UNK	1.0% - 53.9%
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		41		03/31/95	600	0.0867% - 95.785%
TOPICAL; OINTMENT, AUGMENTED		1				
PHARMA-SWEET 24052						
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
PHARMACEUTICAL GLAZE						
ORAL; CAPSULE		31		03/30/95	600	0.945MG - 34.48MG
ORAL; CAPSULE, COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		43		02/14/94	600	0.0126ML - 0.055ML
ORAL; TABLET		13		01/12/89	600	18.0MG
ORAL; TABLET, COATED		10		09/10/87	600	0.76MG - 3.4MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		05/15/90	600	
ORAL; TABLET, FILM COATED		4		06/23/82	600	0.182MG - 0.74MG
ORAL; TABLET, SUSTAINED ACTION		5		06/30/94	600	4.5MG - 6.8MG
PHARMACOAT 606						
ORAL; TABLET		1				
PHENOL	000108952					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		26		12/20/94	600	0.125% - 2.5%
IM - SC; INJECTION		3		04/14/95	600	0.068% - 0.5%
IM - SC; INJECTION, SUSTAINED ACTION		2		09/12/57	510	0.5%
INTRA-ARTICULAR; INJECTION		3		01/30/79	600	0.5%
INTRADERMAL; INJECTION		2		02/08/77	510	0.25%
INTRALESIONAL; INJECTION		3		01/30/79	600	0.5%
INTRAMUSCULAR; INJECTION		18		05/31/94	600	0.25% - 0.5%
INTRASYNOVIAL; INJECTION		1				
INTRAVENOUS; INJECTION		7		08/31/94	600	0.25% - 0.5%
IV(INFUSION); INJECTION		5		05/31/94	600	0.5%
SOFT TISSUE; INJECTION		2		01/30/79	600	0.5%
SUBCUTANEOUS; INJECTION		9		07/01/91	510	0.065% - 0.5%
SUBCUTANEOUS; SOLUTION, INJECTION		1				
PHENOL, LIQUEFIED						
IM - IV; INJECTION		2		01/30/92	600	0.5%
SUBCUTANEOUS; INJECTION		3		04/25/89	510	0.065%
SUBCUTANEOUS; SUSPENSION, INJECTION		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PHENYLETHYL ALCOHOL	000060128					
NASAL; SPRAY, METERED		3		10/19/94	UNK	0.25%
OPHTHALMIC; SOLUTION		1				
OPHTHALMIC; SUSPENSION		1				
OTIC; SOLUTION		1				
PHENYLMERCURIC ACETATE	000062384					
OPHTHALMIC; OINTMENT		3		12/03/86	600	0.0008%
PHENYLMERCURIC NITRATE	000055685					
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
OPHTHALMIC; SOLUTION		1				
PHOSPHATE BUFFER						
ORAL; SOLUTION		1				
PHOSPHOLIPID						
IV(INFUSION); INJECTION		3		12/30/93	510	1.2%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
PHOSPHORIC ACID	007664382					
IM - IV; INJECTION		1				
INTRA-ARTICULAR; INJECTION		1				
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		2		10/17/85	510	
INTRAVENOUS; SOLUTION		1				
IV(INFUSION); INJECTION		3		02/26/93	150	
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		5		08/31/95	600	
ORAL; SOLUTION		1				
SOFT TISSUE; INJECTION		1				
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		2		10/24/89	510	
TOPICAL; EMULSION, CREAM		16		04/30/92	600	0.002% - 10.0%
TOPICAL; LOTION		4		03/30/89	UNK	
TOPICAL; LOTION, AUGMENTED		1				
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		2		01/07/87	520	
VAGINAL; EMULSION, CREAM		6		06/09/86	600	0.07% - 0.8%
PINE NEEDLE OIL						
TOPICAL; LOTION		1				
PINEAPPLE						
ORAL; POWDER, FOR RECONSTITUTION		1				
PIPERAZINE	000110850					
ORAL; TABLET		4		09/23/93	600	
PIPERAZINE HEXAHYDRATE	000142632					
VAGINAL; EMULSION, CREAM		1				
PLASTIBASE-50W						
TOPICAL; OINTMENT		3		06/15/88	600	99.95%
PLUSWEET						
SUBLINGUAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLACRILIN	050602216					
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
POLACRILIN POTASSIUM	039393765					
ORAL; CAPSULE		9		08/22/88	600	1.56MG - 23.0MG
ORAL; TABLET		37		09/23/93	600	0.45MG - 45.8MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-21; TABLET		9		12/30/91	600	1.0MG - 3.0MG
ORAL-28; TABLET		8		12/30/91	600	1.0MG - 3.0MG
POLISTIREX						
OPHTHALMIC; SUSPENSION		1				
POLOXAMER	106392125					
OPHTHALMIC; SOLUTION		1				
ORAL; GRANULE		1				
ORAL; POWDER, FOR RECONSTITUTION		2		09/15/80	600	0.5% - 0.51852%
ORAL; SOLUTION		3		06/01/94	600	10.0%
ORAL; SUSPENSION		2		12/18/87	120	0.02%
ORAL; TABLET		1		05/02/89	520	2.68MG
TOPICAL; SOLUTION		2		07/25/89	520	0.2%
TOPICAL; SPONGE		1				
POLOXAMER 188	106392125					
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
ORAL; CONCENTRATE		1				
ORAL; GRANULE		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
ORAL; TABLET		1				
POLOXAMER 331						
ORAL; POWDER, FOR RECONSTITUTION		3		02/13/87	600	0.5286%
ORAL; SUSPENSION		1				
POLOXAMER 407	009003116					
OPHTHALMIC; SOLUTION		4		12/29/95	600	0.1% - 0.16%
POLYBUTENE						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
POLYDEXTROSE	068424044					
ORAL; TABLET		2		01/27/94	600	
ORAL; TABLET, FILM COATED		1				
POLYDEXTROSE K						
ORAL; TABLET, FILM COATED		2		08/31/94	600	
POLYESTER						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYETHYLENE	009002884					
DENTAL; PASTE		2		10/01/86	600	4.0%
INTRAUTERINE; INTRAUTERINE DEVICE		1				
OPHTHALMIC; OINTMENT		3		12/12/80	UNK	
ORAL; CAPSULE		1				
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; OINTMENT		7		06/15/88	600	5.0% - 9.0%
TOPICAL; PASTE		1				
TOPICAL; TAPE		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE		2		12/22/94	510	27.0MG - 85.0MG
VAGINAL; SUPPOSITORY		1				
POLYETHYLENE GLYCOL	025322683					
IM - IV; INJECTION		1				
INTRAVENOUS; INJECTION		1				
OPHTHALMIC; SOLUTION		1				
ORAL; CAPSULE		14		07/31/92	600	1.1MG - 10.0MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; CONCENTRATE		2		07/20/88	600	50.0% - 60.0%
ORAL; SOLUTION		1				
ORAL; TABLET		202		10/04/95	510	0.045MG - 37.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		19		05/19/92	110	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		4		11/30/95	600	4.5MG - 4.63MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		49		12/08/95	UNK	0.013MG - 3.17MG
ORAL; TABLET, SUSTAINED ACTION		15		09/22/94	110	1.3MG - 3.46MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
TOPICAL; EMULSION		1				
TOPICAL; EMULSION, CREAM		6		02/06/89	600	
TOPICAL; GEL		1				
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		2		10/10/85	600	
TOPICAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		1				
POLYETHYLENE GLYCOL T-DODECYLTHIOETHER						
ORAL; TABLET		1				
POLYETHYLENE GLYCOL 1000						
ORAL; CONCENTRATE		1				
ORAL; SOLUTION		4		12/05/88	600	15.0%
RECTAL; SUPPOSITORY		3		10/05/81	600	1.6255GM
TOPICAL; EMULSION, CREAM		6		11/06/88	600	3.1% - 5.0%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYETHYLENE GLYCOL 1450						
ORAL; SUSPENSION		1				
TOPICAL; OINTMENT		1				
POLYETHYLENE GLYCOL 1500						
TOPICAL; OINTMENT		1				
POLYETHYLENE GLYCOL 1540						
DENTAL; GEL		1				
DENTAL; PASTE		1				
ORAL; CAPSULE		1				
ORAL; SOLUTION		1				
RECTAL; SUPPOSITORY		1				
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		1				
POLYETHYLENE GLYCOL 200						
ORAL; CAPSULE		2		01/27/81	120	
TOPICAL; OINTMENT		2		06/03/77	600	39.0%
POLYETHYLENE GLYCOL 20000						
ORAL; CAPSULE		1				
ORAL-28; TABLET		2		11/17/95	510	0.3MG
POLYETHYLENE GLYCOL 300	025322683					
IM - IV; INJECTION		2		07/03/80	600	50.0%
INTRAVENOUS; INJECTION		2		07/17/95	600	65.0%
IV(INFUSION); INJECTION		5		08/30/95	600	50.0% - 65.0%
OPHTHALMIC; OINTMENT		1				
OPHTHALMIC; SOLUTION		1				
ORAL; TABLET, FILM COATED		1				
TOPICAL; OINTMENT		3		06/15/77	UNK	57.0%
TOPICAL; SOLUTION		2		07/22/81	600	29.7%
POLYETHYLENE GLYCOL 3350						
INTRA-ARTICULAR; INJECTION		2		09/05/61	UNK	3.0%
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		3		10/29/92	510	2.03% - 3.0%
INTRASYNOVIAL; INJECTION		2		09/05/61	UNK	3.0%
NASAL; SPRAY, METERED		1				
ORAL; CAPSULE		1				
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; TABLET		6		10/22/87	520	0.6MG - 10.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		4		06/01/94	110	0.6MG - 1.25MG
RECTAL; SUPPOSITORY		3		08/31/92	600	
SOFT TISSUE; INJECTION		1				
TOPICAL; OINTMENT		6		12/31/87	520	22.0% - 39.0%
VAGINAL; SUPPOSITORY		1				
POLYETHYLENE GLYCOL 3500						
ORAL; TABLET, FILM COATED		2		10/03/90	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYETHYLENE GLYCOL 40 SORBITAN DIISOSTEARATE						
DENTAL; SOLUTION		3		12/28/95	600	0.08% - 2.265%
POLYETHYLENE GLYCOL 400	009004960					
IM - IV; INJECTION		6		05/27/94	600	18.0%
INTRAVENOUS; INJECTION		1				
NASAL; SPRAY, METERED		1				
ORAL; AEROSOL SPRAY		1				
ORAL; CAPSULE		7		07/30/93	600	0.25GM
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, SOFT GELATIN		4		02/19/92	600	520.41MG - 544.0MG
ORAL; SUSPENSION		1				
ORAL; SYRUP		2		01/04/85	600	
ORAL; TABLET		19		07/29/94	600	0.15MG - 1.5MG
ORAL; TABLET, FILM COATED		2		10/27/89	110	1.45GM
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-28; TABLET		1				
RECTAL; SUPPOSITORY		1				
TOPICAL; EMULSION, CREAM		5		09/04/92	UNK	1.0% - 53.0%
TOPICAL; LOTION		2		06/13/88	UNK	5.0%
TOPICAL; OINTMENT		8		12/31/87	520	58.9% - 65.0%
TOPICAL; SOLUTION		5		04/01/91	600	8.89% - 49.8875%
VAGINAL; SUPPOSITORY		2		01/27/87	520	
POLYETHYLENE GLYCOL 4000						
INTRA-ARTICULAR; INJECTION		5		03/26/79	600	2.8% - 3.0%
INTRALESIONAL; INJECTION		3		03/26/79	600	2.8% - 2.96%
INTRAMUSCULAR; INJECTION		5		03/26/79	600	2.8% - 3.0%
INTRASYNOVIAL; INJECTION		3		03/26/79	600	2.8% - 3.0%
ORAL; CAPSULE		2		09/20/90	600	23.0MG - 449.6MG
ORAL; TABLET		15		10/06/95	UNK	0.2MG - 15.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		2		08/31/94	600	
ORAL; TABLET, SUSTAINED ACTION, COATED		1				
RECTAL; SUPPOSITORY		2		10/05/81	600	875.0MG
SOFT TISSUE; INJECTION		3		03/26/79	600	2.8% - 2.96%
SUBLINGUAL; TABLET		2		04/16/81	600	2.5MG
TOPICAL; EMULSION, CREAM		2		01/29/93	600	25.0%
TOPICAL; OINTMENT		5		05/11/81	600	34.8% - 84.0%
TOPICAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		1				
VAGINAL; SUPPOSITORY		1				
POLYETHYLENE GLYCOL 600	006790096					
INTRAVENOUS; SOLUTION, INJECTION		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYETHYLENE GLYCOL 6000						
ORAL; CAPSULE		2		12/10/81	600	1.372MG - 1.79MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		3		01/04/95	600	5.82MG - 17.46MG
ORAL; TABLET		15		06/29/95	600	0.033MG - 375.0MG
ORAL; TABLET, COATED		2		09/06/73	600	5.2MG - 40.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		3		12/12/95	120	0.89MG - 30.0MG
ORAL; TABLET, SUSTAINED ACTION		3		03/30/95	110	0.5MG - 4.0MG
RECTAL; SUPPOSITORY		1				
TOPICAL; EMULSION, CREAM		1				
POLYETHYLENE GLYCOL 8000						
OPHTHALMIC; SOLUTION		2		07/11/94	UNK	2.0%
ORAL; CAPSULE		9		07/18/85	600	1.0MG - 4.9MG
ORAL; TABLET		26		12/21/93	600	0.096MG - 100.0MG
ORAL; TABLET, COATED		6		05/19/92	110	0.08MG - 0.67MG
ORAL; TABLET, SUSTAINED ACTION		4		06/01/94	110	0.06MG - 13.3MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
RECTAL; SUPPOSITORY		2		08/31/92	600	
TOPICAL; EMULSION, CREAM		4		10/17/94	600	5.0%
TOPICAL; POWDER		1				
TOPICAL; SOLUTION		1				
VAGINAL; TABLET		1				
POLYETHYLENE GLYCOL 900						
TOPICAL; OINTMENT		1				
POLYETHYLENE OXIDE	025322683					
ORAL; TABLET, SUSTAINED ACTION		3		06/01/94	110	82.49MG - 277.15MG
POLYETHYLENE TEREPHTHALATES	009003514					
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
POLYGLACTIN	026780507					
IMPLANTATION; PELLET, IMPLANT		2		12/18/95	150	
INTRAMUSCULAR; INJECTION		2		01/21/94	510	6.62%
SUBCUTANEOUS; PELLET, IMPLANT		2		12/18/95	150	
POLYGLYCERYL-10 TETRALINOLEATE						
ORAL; SUSPENSION		1				
POLYISOBUTYLENE	009003274					
TRANSDERMAL; FILM, CONTROLLED RELEASE		6		10/28/94	510	6.3MG - 119.0MG
POLYISOBUTYLENE 1,200,000						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
POLYLACTIDE						
INTRAMUSCULAR; POWDER, FOR INJECTION SUSPENSION, LYOPHI		1				
POLYMERS						
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
POLYOLS						
DENTAL; GEL		1				
DENTAL; PASTE		1				
POLYOXYETHYLENE - POLYOXYPROPYLENE 1800						
OPHTHALMIC; SOLUTION		1				
TOPICAL; EMULSION, CREAM		1				
POLYOXYETHYLENE ALCOHOLS	009007630					
TOPICAL; EMULSION, CREAM		7		12/22/87	600	7.5% - 9.14%
TOPICAL; OINTMENT		1				
POLYOXYETHYLENE FATTY ACID ESTERS						
IM - IV - SC; INJECTION		1				
IM - SC; INJECTION		1				
POLYOXYETHYLENE PROPYLENE						
TOPICAL; LOTION		1				
POLYOXYETHYLENE SORBITAN MONOISOSTEARATE						
INTRAMUSCULAR; INJECTION		1				
POLYOXYL CASTOR OIL	008047163					
IV(INFUSION); INJECTION		1				
POLYOXYL DISTEARATE	009005087					
TOPICAL; OINTMENT		1				
POLYOXYL GLYCERYL STEARATE						
TOPICAL; EMULSION, CREAM		1				
POLYOXYL LANOLIN						
TOPICAL; SOLUTION		1				
POLYOXYL STEARATE						
OTIC; SUSPENSION		1				
RECTAL; SUPPOSITORY		1				
TOPICAL; EMULSION		1				
TOPICAL; EMULSION, CREAM		8		07/26/88	600	1.0% - 20.0%
TOPICAL; LOTION		1				
POLYOXYL 100 GLYCERYL STEARATE						
TOPICAL; EMULSION, CREAM		1				
VAGINAL; EMULSION, CREAM		1				
POLYOXYL 100 STEARATE						
TOPICAL; EMULSION, CREAM		2		02/16/94	600	
TOPICAL; LOTION		1				
VAGINAL; EMULSION, CREAM		1				
POLYOXYL 15 COCAMINE	008051523					
TOPICAL; SPONGE		1				
POLYOXYL 150 DISTEARATE						
TOPICAL; SOLUTION		1				
POLYOXYL 2 STEARATE						
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYOXYL 20 STEARATE						
ORAL; TABLET, SUSTAINED ACTION		1				
POLYOXYL 35 CASTOR OIL	061791126					
IV(INFUSION); INJECTION		1				
IV(INFUSION); SOLUTION, INJECTION		1				
OPHTHALMIC; SOLUTION		1				
POLYOXYL 40 CASTOR OIL						
IV(INFUSION); INJECTION		1				
POLYOXYL 40 HYDROGENATED CASTOR OIL	061788850					
ORAL; SOLUTION		1				
POLYOXYL 40 STEARATE	009004993					
DENTAL; SOLUTION		1				
OPHTHALMIC; OINTMENT		1				
OPHTHALMIC; SOLUTION		3		09/25/85	600	7.0%
OPHTHALMIC; SUSPENSION		2		05/11/88	600	0.5%
ORAL; CAPSULE		4		08/03/87	600	1.0MG - 2.64MG
ORAL; CONCENTRATE		1				
ORAL; GRANULE		1				
ORAL; TABLET		16		02/21/92	520	0.8MG - 8.48MG
OTIC; SUSPENSION		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		44		11/20/92	UNK	0.45% - 8.8%
TOPICAL; LOTION		7		04/24/85	UNK	0.502% - 3.5%
TOPICAL; OINTMENT		2		07/24/78	600	3.0%
POLYOXYL 50 STEARATE						
TOPICAL; OINTMENT		1				
POLYOXYL 60 CASTOR OIL						
IV(INFUSION); INJECTION		1				
POLYOXYL 75 LANOLIN						
TOPICAL; SOLUTION		1				
POLYOXYL 8 STEARATE	009004993					
TOPICAL; EMULSION, CREAM		3		10/08/85	600	1.0% - 8.0%
POLYOXYPROPYLENE 15 STEARYL ETHER						
TOPICAL; OINTMENT		1				
POLYOXYPROPYLENE 26 OLEATE						
TOPICAL; EMULSION, CREAM		2		06/13/88	600	1.5% - 4.0%
POLYPROPYLENE	009003070					
TRANSDERMAL; FILM, CONTROLLED RELEASE		2		10/10/84	110	4.5MG - 13.5MG
POLYPROPYLENE GLYCOL	009003150					
OPHTHALMIC; SOLUTION		1				
ORAL; TABLET		2		12/28/88	110	1.26MG
POLYSACCHARIDE						
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
POLYSILOXANE	009014135					
ORAL; CAPSULE		4		04/21/87	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYSORBATE 80	009005656					
IV(INFUSION); INJECTION		6		08/30/95	600	0.8% - 8.0%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
IV(INFUSION); SOLUTION, INJECTION		1				
NASAL; SPRAY, METERED		3		10/19/94	UNK	0.005%
OPHTHALMIC; SOLUTION		2		05/25/71	600	0.01% - 0.2%
OPHTHALMIC; SUSPENSION		18		12/30/94	UNK	0.002% - 0.1%
ORAL; CAPSULE		12		12/20/95	520	0.16MG - 418.37MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		2		09/11/95	110	0.071MG
ORAL; CONCENTRATE		1				
ORAL; DROPS		1				
ORAL; GRANULE		4		06/15/88	600	
ORAL; POWDER		1				
ORAL; POWDER, FOR RECONSTITUTION		5		12/20/95	520	0.028% - 2.625%
ORAL; SOLUTION		5		04/22/87	600	0.1512%
ORAL; SUSPENSION		22		06/16/95	UNK	0.02% - 0.375%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; TABLET		125		05/31/95	600	0.04MG - 12.0MG
ORAL; TABLET, COATED		2		12/20/82	600	
ORAL; TABLET, FILM COATED		29		09/14/95	110	0.0334MG - 14.8MG
ORAL; TABLET, SUSTAINED ACTION		5		01/31/94	600	17.5MG
OTIC; SOLUTION		1				
OTIC; SUSPENSION		3		11/06/85	600	0.01% - 5.0%
RECTAL; ENEMA		2		05/27/94	600	
RECTAL; POWDER, FOR RECONSTITUTION		1				
RECTAL; SOLUTION		2		12/19/77	160	
RECTAL; SUPPOSITORY		8		02/27/95	600	
SOFT TISSUE; INJECTION		3		05/24/82	600	0.075% - 0.2%
SUBCUTANEOUS; INJECTION		1				
SUBLINGUAL; TABLET		2		04/16/81	600	0.075MG
TOPICAL; EMULSION, CREAM		8		10/26/94	600	0.1% - 1.5%
TOPICAL; GEL		1				
TOPICAL; LOTION		2		06/05/78	600	0.01%
TOPICAL; OINTMENT		1				
VAGINAL; EMULSION, CREAM		2		02/21/91	520	
VAGINAL; SUPPOSITORY		3		01/27/87	520	28.0MG
POLYSORBATE 85	009005703					
IM - IV; INJECTION		1				
IV(INFUSION); INJECTION		1				
POLYVINYL ACETATE	009003207					
ORAL; TABLET		2		09/10/87	520	7.0MG
ORAL; TABLET, SUSTAINED ACTION		3		01/25/93	600	20.0MG - 46.0MG
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYSORBATE						
ORAL; TABLET		4		11/20/92	110	0.04MG - 0.9MG
ORAL; TABLET, SUSTAINED ACTION		3		05/14/85	UNK	
TOPICAL; EMULSION, CREAM		1				
POLYSORBATE 20	009005645					
INTRAVENOUS; INJECTION		1				
IV(INFUSION); INJECTION		2		08/08/85	510	0.014%
NASAL; SPRAY, METERED		1				
OPHTHALMIC; SUSPENSION		5		09/15/95	600	0.05%
ORAL; CAPSULE		1				
ORAL; SUSPENSION		2		09/15/95	180	0.1%
ORAL; TABLET		2		12/23/93	180	0.18MG - 1.8MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		1				
SUBCUTANEOUS; SOLUTION, INJECTION		1				
TOPICAL; LOTION		6		08/14/86	600	0.65% - 7.8%
TOPICAL; SOLUTION		2		04/17/92	600	
POLYSORBATE 40	009005667					
INTRAMUSCULAR; INJECTION		1				
ORAL; SOLUTION, ELIXIR		2		04/29/93	600	
ORAL; SUSPENSION		2		09/25/92	600	
ORAL; SYRUP		1				
TOPICAL; EMULSION, CREAM		7		04/29/94	600	2.5% - 2.8%
TOPICAL; LOTION		2		11/30/82	600	0.5263% - 3.0%
TOPICAL; OINTMENT		1				
POLYSORBATE 60	009005678					
ORAL; SUSPENSION		1				
ORAL; TABLET, COATED		1				
RECTAL; SUPPOSITORY		1				
TOPICAL; EMULSION, CREAM		59		06/13/95	600	0.2% - 6.1%
TOPICAL; LOTION		4		09/30/92	UNK	1.225% - 3.36%
TOPICAL; OINTMENT		2		10/09/85	600	
TOPICAL; SHAMPOO		4		09/18/84	600	9.0% - 15.0%
TOPICAL; SUPPOSITORY		1				
VAGINAL; EMULSION, CREAM		9		12/21/95	520	1.0% - 5.0%
VAGINAL; SUPPOSITORY		1				
POLYSORBATE 80	009005656					
INTRA-ARTICULAR; INJECTION		21		02/17/84	600	0.04% - 0.4%
INTRABURSAL; INJECTION		5		02/13/74	600	0.04% - 0.2%
INTRADERMAL; INJECTION		2		10/16/87	UNK	0.04%
INTRALESIONAL; INJECTION		7		10/16/87	UNK	0.04% - 0.4%
INTRAMUSCULAR; INJECTION		20		10/29/92	510	0.04% - 12.0%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		2		07/21/61	120	
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRASYNOVIAL; INJECTION		11		02/17/84	600	0.04% - 0.4%
INTRAVENOUS; INJECTION		4		07/17/95	600	8.0%
INTRAVENOUS; SOLUTION, INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POLYVINYL ACETATE PHTHALATE						
ORAL; TABLET, SUSTAINED ACTION		1				
POLYVINYL ALCOHOL	009002895					
OPHTHALMIC; SOLUTION		13		01/04/95	600	0.25% - 1.4%
OPHTHALMIC; SUSPENSION		8		09/29/89	UNK	1.4%
ORAL; TABLET		1				
ORAL; TABLET, FILM COATED		1				
POLYVINYLACETAL						
ORAL; TABLET		1				
POLYVINYLPYRIDINE	000100936					
ORAL; TABLET		1				
POLYVINYLPYRROLIDONE ETHYLCELLULOSE						
ORAL; TABLET		1				
POPPY SEED OIL	008002117					
INTRALYMPHATIC; OIL		1				
INTRAUTERINE; OIL		1				
POTASSIUM ACETATE	000127082					
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
RECTAL; ENEMA		1				
POTASSIUM CARBONATE	000584087					
ORAL; CAPSULE		11		08/26/88	600	2.552MG - 20.0MG
ORAL; SOLUTION		2		04/22/87	600	0.496% - 0.62%
ORAL; TABLET		15		05/11/90	600	3.5MG - 20.0MG
POTASSIUM CHLORIDE	007447407					
CAUDAL BLOCK; INJECTION		1				
EPIDURAL; INJECTION		3		12/01/86	600	0.03%
INTRAOCULAR; SOLUTION		2		04/28/95	600	0.075%
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; SOLUTION, INJECTION		1				
NERVE BLOCK; INJECTION		6		12/01/86	600	0.012% - 0.03%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		2		09/22/93	UNK	
OPHTHALMIC; SOLUTION		9		01/04/95	600	0.042% - 0.23%
OPHTHALMIC; SUSPENSION		1				
ORAL; CAPSULE		13		08/26/88	600	0.57MG - 31.0MG
ORAL; TABLET		12		05/11/90	600	7.75MG - 31.0MG
POTASSIUM CITRATE	006100056					
ORAL; SOLUTION		1				
POTASSIUM HYDROXIDE	001310583					
INTRAVENOUS; INJECTION		1				
IV(INFUSION); INJECTION		3		03/10/88	600	
ORAL; CAPSULE		2		04/21/87	600	
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; PASTILLE		1				
ORAL; SOLUTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POTASSIUM METABISULFITE	004429429					
IM - IV; INJECTION		1				
INTRAVENOUS; INJECTION		1				
IV(INFUSION); INJECTION		5		11/18/85	510	
NERVE BLOCK; INJECTION		2		01/22/85	600	0.12%
OTIC; SOLUTION		4		12/29/95	600	0.1%
OTIC; SUSPENSION		1				
RECTAL; ENEMA		1				
POTASSIUM PHOSPHATE, DIBASIC	007758114					
INTRAVENOUS; SOLUTION		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		2		03/18/87	600	
ORAL; SYRUP		1				
ORAL; TABLET		1				
SUBCUTANEOUS; INJECTION		1				
POTASSIUM PHOSPHATE, MONOBASIC	007778770					
IM - IV; INJECTION		1				
INTRA-ARTICULAR; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
INTRAVENOUS; INJECTION		2		11/16/76	160	0.076%
NERVE BLOCK; INJECTION		1				
OPHTHALMIC; SOLUTION		6		03/04/94	600	0.03402% - 0.2%
OPHTHALMIC; SUSPENSION		1				
ORAL; SYRUP		3		10/31/95	600	0.095%
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OTIC; SOLUTION		1				
POTASSIUM POLACRILIN						
ORAL; TABLET		2		05/15/90	600	24.0MG
POTASSIUM SORBATE	000590001					
ORAL; CAPSULE		5		09/30/86	UNK	
ORAL; CONCENTRATE		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SOLUTION		3		12/03/86	600	
ORAL; SUSPENSION		1				
ORAL; SYRUP		3		11/10/83	600	
ORAL; TABLET		3		11/05/92	600	0.8MG
TOPICAL; EMULSION, CREAM		10		02/01/89	600	0.095% - 0.15%
TOPICAL; LOTION		2		06/05/78	600	0.1%
POVIDONE	009003398					
INTRAMUSCULAR; INJECTION		7		07/07/83	600	0.03% - 0.9%
IV(INFUSION); INJECTION		1				
OPHTHALMIC; SOLUTION		1				
OPHTHALMIC; SUSPENSION		1				
ORAL; CAPSULE		33		10/18/95	600	1.8MG - 44.1MG
ORAL; CAPSULE, COATED PELLETS		3		06/30/92	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		12/22/86	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POVIDONE	009003398					
ORAL; CAPSULE, SUSTAINED ACTION		26		02/16/94	600	0.15MG - 72.0MG
ORAL; DROPS		1				
ORAL; GRANULE		2		05/20/88	600	
ORAL; GRANULE FOR RECONSTITUTION, CR		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
ORAL; TABLET		441		12/28/95	600	0.012GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		05/14/82	520	5.0MG
ORAL; TABLET, COATED		31		06/23/95	600	3.0MG - 49.2MG
ORAL; TABLET, CONTROLLED RELEASE		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		6		08/28/95	600	1.0MG - 10.66MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		25		12/27/95	150	1.91MG - 40.0MG
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		30		11/18/94	UNK	3.0MG - 60.0MG
ORAL; TABLET, UNCOATED, TROCHE		1				
ORAL-21; TABLET		18		12/13/93	600	0.14MG - 4.0MG
ORAL-28; TABLET		24		11/17/95	510	0.14MG - 4.5MG
SUBLINGUAL; TABLET		2		04/16/81	600	6.0MG
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
TOPICAL; SOLUTION		1				
VAGINAL; TABLET		3		12/26/91	520	50.0MG
POVIDONE K25	0					
ORAL; TABLET		2		06/29/95	600	0.44MG - 52.0MG
ORAL; TABLET, SUSTAINED ACTION		1				
POVIDONE K29-32						
ORAL; TABLET		6		07/29/94	600	0.5MG - 1.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-28; TABLET		1				
POVIDONE K30						
ORAL; CAPSULE		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET		8		05/31/94	530	4.5MG - 18.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, FILM COATED		1				.0MG
POVIDONE K90						
ORAL; CAPSULE		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; TABLET		4		04/28/95	600	0.88MG - 18.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
POVIDONE K90						
ORAL; TABLET, FILM COATED		1				
PROMALGEN TYPE G						
TOPICAL; LOTION		1				
PROMULGEN D						
TOPICAL; LOTION		1				
VAGINAL; EMULSION, CREAM		1				
PROMULGEN G	009009614					
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		2		11/26/85	600	2.16%
TOPICAL; SHAMPOO		1				
PROPANE	000074986					
TOPICAL; AEROSOL SPRAY		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
PROPENYL GUAETHOL	000094860					
ORAL; CAPSULE		1				
PROPYL GALLATE	000121799					
INTRAMUSCULAR; INJECTION		1				
ORAL; CONCENTRATE		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; GEL		2		12/30/94	600	0.01% - 0.05%
TOPICAL; OINTMENT		2		11/13/81	UNK	
PROPYLENE CARBONATE	000108327					
TOPICAL; OINTMENT		1				
PROPYLENE GLYCOL	000057556					
IM - IV; INJECTION		41		05/27/94	600	0.04% - 50.0%
IM - IV; SOLUTION, INJECTION		1				
INHALATION; SOLUTION		1				
INTRAMUSCULAR; INJECTION		3		02/12/86	600	2.07% - 40.0%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		2		07/21/61	120	
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRAVENOUS; INJECTION		5		12/29/89	110	4.5% - 37.5%
IV(INFUSION); INJECTION		16		09/11/92	600	0.09% - 40.0%
IV(INFUSION); SOLUTION, INJECTION		2		12/31/86	110	25.0% - 30.0%
NASAL; SPRAY, METERED		2		03/08/95	UNK	
OPHTHALMIC; SOLUTION		5		10/11/88	600	0.12% - 1.0%
OPHTHALMIC; SUSPENSION		5		11/10/93	UNK	0.12% - 5.0%
ORAL; CAPLET		1				
ORAL; CAPSULE		16		03/16/95	600	52.0MG
ORAL; CAPSULE, SOFT GELATIN		3		07/14/95	530	0.135MG - 148.31MG
ORAL; CAPSULE, SUSTAINED ACTION		7		04/25/95	UNK	0.15MG - 0.39MG
ORAL; CONCENTRATE		17		11/30/94	600	0.07% - 90.0%
ORAL; SOLUTION		31		11/17/95	530	2.0% - 50.0%
ORAL; SOLUTION, ELIXIR		9		10/27/92	600	25.8917%
ORAL; SUSPENSION		17		12/18/87	120	0.69% - 8.0%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; SYRUP		44		07/17/95	600	5.0% - 7.0%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PROPYLENE GLYCOL	000057556					
ORAL; TABLET		93		08/24/95	600	0.00006ML - 0.00012ML
ORAL; TABLET, COATED		4		12/30/92	110	0.4MG - 1.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		5		06/19/95	520	3.46MG - 6.95MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		25		05/25/95	600	0.33MG - 2.1MG
ORAL; TABLET, SUSTAINED ACTION		7		04/17/86	110	0.71MG - 5.3MG
OTIC; SOLUTION		11		12/29/95	600	2.0% - 94.9%
OTIC; SOLUTION, DROPS		1				
OTIC; SUSPENSION		4		09/29/87	600	2.0% - 10.0%
RECTAL; EMULSION, AEROSOL FOAM		1				
RECTAL; SUSPENSION		2		11/17/86	600	
TOPICAL; AEROSOL		1				
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, AEROSOL FOAM		3		12/19/79	600	5.3/6%
TOPICAL; EMULSION, CREAM		124		09/13/95	UNK	0.2% - 67.43%
TOPICAL; GEL		9		12/30/94	600	3.0% - 98.09%
TOPICAL; LOTION		24		09/30/92	UNK	2.0% - 50.0%
TOPICAL; LOTION, AUGMENTED		1				
TOPICAL; OINTMENT		26		03/31/95	600	0.012% - 38.0%
TOPICAL; OINTMENT, AUGMENTED		2		08/31/95	600	10.0%
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		39		11/30/95	600	3.0% - 99.99%
TOPICAL; SPONGE		2		02/28/91	600	
TOPICAL; SUSPENSION, SHAMPOO		1				
TOPICAL; SWAB		3		07/30/95	600	25.0%
VAGINAL; EMULSION, CREAM		12		12/21/95	520	3.0% - 14.0%
VAGINAL; GEL		1				
VAGINAL; SUPPOSITORY		1				
PROPYLENE GLYCOL ALGINATE	009005372					
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER		2		12/05/88	510	0.002%
PROPYLENE GLYCOL DIACETATE	000623847					
OTIC; SOLUTION		7		10/31/94	600	3.0%
PROPYLENE GLYCOL MONOLAURATE	001322878					
TOPICAL; EMULSION, CREAM		1				
PROPYLENE GLYCOL MONOSTEARATE	001323393					
TOPICAL; EMULSION, CREAM		25		05/31/91	600	0.3% - 8.0%
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		4		04/30/87	UNK	2.0%
TOPICAL; OINTMENT, AUGMENTED		2		08/31/95	600	2.0%
VAGINAL; EMULSION, CREAM		1				
PROPYLPARABEN	000094133					
IM - IV - SC; INJECTION		25		04/11/89	600	0.02% - 0.2%
IM - IV; INJECTION		33		11/27/91	600	0.01% - 0.2%
IM - SC; INJECTION		1				
INHALATION; SOLUTION		5		06/30/81	600	0.007% - 0.0150066%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT / ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PROPYLPARABEN	000094133					
INTRA-ARTICULAR; INJECTION		3		06/19/80	600	0.02%
INTRABURSAL; INJECTION		1				
INTRALESIONAL; INJECTION		3		06/19/80	600	0.02%
INTRAMUSCULAR; INJECTION		27		02/25/93	600	0.01% - 0.02%
INTRASYNOVIAL; INJECTION		2		03/01/77	UNK	0.02%
INTRAVENOUS; INJECTION		16		12/20/91	UNK	0.005% - 0.03%
IV - SC; INJECTION		2		12/05/85	180	0.015% - 0.02%
IV(INFUSION); INJECTION		19		03/25/94	160	0.0005% - 0.056%
NASAL; SOLUTION		2		05/18/70	510	0.017%
NERVE BLOCK; INJECTION		2		03/06/72	600	0.005% - 0.015%
OPHTHALMIC; OINTMENT		8		08/31/95	600	0.01%
OPHTHALMIC; SOLUTION		9		10/18/88	600	0.01% - 0.015%
OPHTHALMIC; SUSPENSION		2		12/28/82	600	0.01%
ORAL; CAPSULE		79		12/20/95	520	0.0246MG - 0.188MG
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	0.05MG - 0.081MG
ORAL; CAPSULE, SUSTAINED ACTION		8		04/25/95	UNK	
ORAL; CONCENTRATE		23		09/28/93	600	0.004% - 0.03%
ORAL; POWDER, FOR RECONSTITUTION		5		12/31/91	520	0.01% - 0.08%
ORAL; SOLUTION		42		11/17/95	530	0.01% - 20.0%
ORAL; SOLUTION, ELIXIR		10		10/27/92	600	0.02%
ORAL; SUSPENSION		41		09/15/95	180	0.01% - 0.05%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; SYRUP		39		07/17/95	600	0.0085% - 0.02%
ORAL; TABLET		19		03/30/94	600	0.0045MG - 0.14MG
ORAL; TABLET, COATED		11		04/08/81	600	0.002MG
ORAL; TABLET, FILM COATED		2		12/28/87	520	0.02MG - 0.04MG
ORAL; TABLET, SUSTAINED ACTION		3		05/22/87	UNK	0.12MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
OTIC; SUSPENSION		1				
RECTAL; EMULSION, AEROSOL FOAM		1				
RECTAL; SOLUTION		1				
RECTAL; SUSPENSION		2		11/17/86	600	
SOFT TISSUE; INJECTION		2		06/19/80	600	0.02%
SUBCUTANEOUS; INJECTION		2		12/19/91	UNK	0.015%
TOPICAL; EMULSION, AEROSOL FOAM		2		12/19/79	600	0.011%
TOPICAL; EMULSION, CREAM		50		10/31/94	600	0.001% - 0.15%
TOPICAL; GEL		1				
TOPICAL; GEL, JELLY		2		04/29/93	600	0.003%
TOPICAL; LOTION		12		12/07/92	UNK	0.02% - 0.2%
TOPICAL; OINTMENT		10		09/30/94	600	0.01% - 0.2%
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PROPYLPARABEN	000094133					
URETERAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		11		12/21/95	520	0.02% - 0.1%
VAGINAL; GEL		1				
VAGINAL; SUPPOSITORY		1				
PROPYLPARABEN SODIUM	035285699					
ORAL; CAPSULE, SOFT GELATIN		2		12/30/86	150	0.12MG - 0.28MG
ORAL; POWDER, FOR RECONSTITUTION		1				
PROSWEET						
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
PROSWEET 604						
ORAL; SYRUP		1				
PROTAMINE SULFATE	009009658					
IM - SC; INJECTION		1				
INTRADERMAL; INJECTION		2		02/08/77	510	0.043% - 0.12%
SUBCUTANEOUS; INJECTION		4		10/28/82	510	0.035% - 0.036%
SUBCUTANEOUS; SUSPENSION, INJECTION		1				
PROTEIN HYDROLYSATE	009015547					
TOPICAL; LOTION		1				
RA-2397						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
RA-3011						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
ROSIN	008050097					
ORAL; CAPSULE		1				
ORAL; TABLET		1				
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		1				
SACCHARIN	000081072					
INHALATION; AEROSOL, METERED		1				
ORAL; AEROSOL SPRAY		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		3		01/05/78	520	
ORAL; SYRUP		2		01/19/83	UNK	0.02%
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
SUBLINGUAL; TABLET		1				
TOPICAL; OINTMENT		4		06/03/77	600	0.4% - 0.5%
TOPICAL; SOLUTION		1				
SACCHARIN CALCIUM	006381915					
ORAL; SOLUTION		1				
ORAL; SYRUP		1				